US010709364B2

(12) **United States Patent**
Kamath et al.

(10) **Patent No.: US 10,709,364 B2**
(45) **Date of Patent: Jul. 14, 2020**

(54) **SYSTEM AND METHODS FOR PROCESSING ANALYTE SENSOR DATA FOR SENSOR CALIBRATION**

(71) Applicant: **DexCom, Inc.**, San Diego, CA (US)

(72) Inventors: **Apurv Ullas Kamath**, San Diego, CA (US); **James H. Brauker**, Addison, MI (US); **Paul V. Goode, Jr.**, Round Rock, TX (US); **Aarthi Mahalingam**, San Diego, CA (US); **Jack Pryor**, Ladera Ranch, CA (US)

(73) Assignee: **DexCom, Inc.**, San Diego, CA (US)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.

(21) Appl. No.: **16/690,977**

(22) Filed: **Nov. 21, 2019**

(65) **Prior Publication Data**

US 2020/0085354 A1 Mar. 19, 2020

Related U.S. Application Data

(63) Continuation of application No. 16/457,628, filed on Jun. 28, 2019, which is a continuation of application
(Continued)

(51) **Int. Cl.**
A61B 5/1473 (2006.01)
A61B 5/1486 (2006.01)
(Continued)

(52) **U.S. Cl.**
CPC ***A61B 5/14546*** (2013.01); ***A61B 5/0031*** (2013.01); ***A61B 5/1451*** (2013.01);
(Continued)

(58) **Field of Classification Search**
CPC . A61B 5/14546; A61B 5/0031; A61B 5/1451; A61B 5/14532; A61B 5/1473;
(Continued)

(56) **References Cited**

U.S. PATENT DOCUMENTS

1,498,738 A 6/1924 Lahousse et al.
1,564,641 A 12/1925 Robert St et al.
(Continued)

FOREIGN PATENT DOCUMENTS

CA 2127172 C 7/1998
CN 101026994 A 8/2007
(Continued)

OTHER PUBLICATIONS

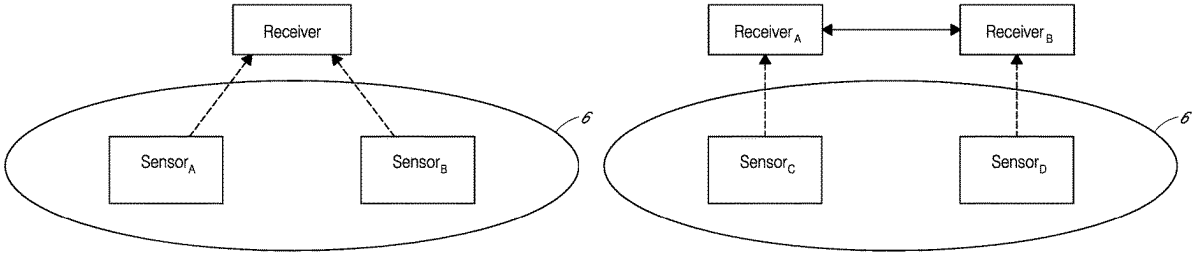
US 7,530,950 B2, 05/2009, Brister et al. (withdrawn)
(Continued)

Primary Examiner — Navin Natnithithadha
(74) *Attorney, Agent, or Firm* — Knobbe Martens Olson & Bear, LLP

(57) **ABSTRACT**

Systems and methods for processing sensor analyte data are disclosed, including initiating calibration, updating calibration, evaluating clinical acceptability of reference and sensor analyte data, and evaluating the quality of sensor calibration. The sensor can be calibrated using a calibration set of one or more matched sensor and reference analyte data pairs. Reference data resulting from benchtop testing an analyte sensor prior to its insertion can be used to provide initial calibration of the sensor data. Reference data from a short term continuous analyte sensor implanted in a user can be used to initially calibrate or update sensor data from a long term continuous analyte sensor.

10 Claims, 48 Drawing Sheets



Related U.S. Application Data

No. 15/787,595, filed on Oct. 18, 2017, which is a continuation of application No. 15/065,623, filed on Mar. 9, 2016, now Pat. No. 9,918,668, which is a continuation of application No. 13/607,162, filed on Sep. 7, 2012, now Pat. No. 9,314,196, which is a continuation of application No. 12/683,755, filed on Jan. 7, 2010, now Pat. No. 8,611,978, which is a continuation of application No. 11/373,628, filed on Mar. 9, 2006, now Pat. No. 7,920,906.

(60) Provisional application No. 60/660,743, filed on Mar. 10, 2005.

(51) **Int. Cl.**

A61B 5/1468	(2006.01)
A61B 5/1495	(2006.01)
A61B 5/145	(2006.01)
A61B 5/00	(2006.01)

(52) **U.S. Cl.**

CPC ***A61B 5/1473*** (2013.01); ***A61B 5/1495*** (2013.01); ***A61B 5/14532*** (2013.01); ***A61B 5/14865*** (2013.01); *A61B 5/1468* (2013.01); *A61B 5/1486* (2013.01); *A61B 5/14503* (2013.01); *A61B 5/14735* (2013.01); *A61B 2560/0223* (2013.01); *Y02A 90/26* (2018.01)

(58) **Field of Classification Search**

CPC A61B 5/14865; A61B 5/1495; A61B 5/14503; A61B 5/1468; A61B 5/14735; A61B 5/1486; A61B 2560/0223; Y02A 90/26

USPC 600/309, 345–366

See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

1,726,766	A	9/1929	Rector et al.
2,057,029	A	10/1936	Johnstone
2,402,306	A	6/1946	Henry et al.
2,497,894	A	2/1950	Clare et al.
2,719,797	A	10/1955	Rosenblatt et al.
2,882,696	A	4/1959	Herrmann et al.
2,910,256	A	10/1959	Richard et al.
3,022,639	A	2/1962	Brown et al.
3,210,578	A	10/1965	Sherer
3,218,819	A	11/1965	Crotser et al.
3,219,533	A	11/1965	Mullins
3,381,371	A	5/1968	Russell et al.
3,506,032	A	4/1970	Eveleigh et al.
3,539,455	A	11/1970	Clark, Jr. et al.
3,556,950	A	1/1971	Dahms et al.
3,562,352	A	2/1971	Nyilas
3,581,062	A	5/1971	Aston
3,607,329	A	9/1971	Manjikian
3,610,226	A	10/1971	Albisser et al.
3,652,475	A	3/1972	Wada et al.
3,728,678	A	4/1973	Tong
3,775,182	A	11/1973	Patton et al.
3,780,727	A	12/1973	King
3,791,871	A	2/1974	Rowley
3,826,244	A	7/1974	Salcman et al.
3,837,339	A	9/1974	Aisenberg et al.
3,838,033	A	9/1974	Mindt et al.
3,838,682	A	10/1974	Clark et al.
3,872,455	A	3/1975	Fuller et al.
3,874,850	A	4/1975	Sorensen et al.
3,882,011	A	5/1975	Hines et al.
3,898,984	A	8/1975	Mandel et al.
3,910,256	A	10/1975	Clark et al.
3,926,760	A	12/1975	Allen et al.
3,929,971	A	12/1975	Roy et al.
3,930,462	A	1/1976	Day et al.
3,933,593	A	1/1976	Sternberg et al.
3,943,918	A	3/1976	Lewis et al.
3,949,388	A	4/1976	Fuller et al.
3,957,613	A	5/1976	Macur
3,957,651	A	5/1976	Kesting et al.
3,964,974	A	6/1976	Banauch et al.
3,966,580	A	6/1976	Janata et al.
3,978,684	A	9/1976	Taylor et al.
3,979,274	A	9/1976	Newman
3,982,530	A	9/1976	Storch et al.
4,003,621	A	1/1977	Lamp et al.
4,008,717	A	2/1977	Kowarski et al.
4,016,866	A	4/1977	Lawton et al.
4,024,312	A	5/1977	Korpman et al.
4,036,749	A	7/1977	Anderson
4,037,563	A	7/1977	Pflueger et al.
4,040,908	A	8/1977	Clark, Jr. et al.
4,052,754	A	10/1977	Homsy et al.
4,055,175	A	10/1977	Clemens et al.
4,067,322	A	1/1978	Johnson et al.
4,068,660	A	1/1978	Beck et al.
4,073,713	A	2/1978	Newman
4,076,656	A	2/1978	White et al.
4,101,395	A	7/1978	Motani et al.
4,109,505	A	8/1978	Clark et al.
4,110,997	A	9/1978	Klotz et al.
4,116,920	A	9/1978	Honma et al.
4,119,406	A	10/1978	Clemens et al.
4,129,128	A	12/1978	McFarlane
4,151,845	A	5/1979	Clemens
4,172,770	A	10/1979	Semersky et al.
4,176,659	A	12/1979	Rolfe
4,187,390	A	2/1980	Gore
4,197,840	A	4/1980	Beck et al.
4,197,852	A	4/1980	Schindler et al.
4,206,755	A	6/1980	Klein
4,215,703	A	8/1980	Willson
4,225,410	A	9/1980	Pace et al.
4,230,118	A	10/1980	Holman et al.
4,240,438	A	12/1980	Shults et al.
4,240,889	A	12/1980	Yoda et al.
4,245,634	A	1/1981	Albisser et al.
4,248,246	A	2/1981	Ikeda
4,253,469	A	3/1981	Aslan et al.
4,255,500	A	3/1981	Hooke et al.
4,259,540	A	3/1981	Sabia et al.
4,260,725	A	4/1981	Keogh et al.
4,260,726	A	4/1981	Deubzer et al.
4,265,249	A	5/1981	Schindler et al.
4,286,039	A	8/1981	Landa et al.
4,319,578	A	3/1982	Enger et al.
4,324,256	A	4/1982	Vesterager et al.
4,324,257	A	4/1982	Albarda et al.
4,327,725	A	5/1982	Cortese et al.
4,344,438	A	8/1982	Schultz
4,349,728	A	9/1982	Phillips et al.
4,353,368	A	10/1982	Slovak et al.
4,353,888	A	10/1982	Sefton et al.
4,366,040	A	12/1982	Marsoner et al.
4,369,785	A	1/1983	Rehkopf et al.
4,374,013	A	2/1983	Enfors
4,378,016	A	3/1983	Loeb et al.
4,388,166	A	6/1983	Suzuki et al.
4,402,694	A	9/1983	Ash et al.
4,403,847	A	9/1983	Chrestensen
4,403,984	A	9/1983	Ash et al.
4,415,666	A	11/1983	D'Orazio et al.
4,418,148	A	11/1983	Oberhardt
4,419,535	A	12/1983	O'Hara
4,425,920	A	1/1984	Bourland et al.
4,431,004	A	2/1984	Bessman et al.
4,431,507	A	2/1984	Nankai et al.
4,432,366	A	2/1984	Margules
4,436,094	A	3/1984	Cerami
4,442,841	A	4/1984	Uehara et al.
4,453,537	A	6/1984	Spitzer
4,454,295	A	6/1984	Wittmann et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

4,457,339 A 7/1984 Juan et al.
4,469,110 A 9/1984 Slama
4,477,314 A 10/1984 Richter et al.
4,478,222 A 10/1984 Koning et al.
4,478,976 A 10/1984 Goertz et al.
4,484,987 A 11/1984 Gough
4,486,290 A 12/1984 Cahalan et al.
4,492,575 A 1/1985 Mabille
4,493,714 A 1/1985 Ueda et al.
4,494,950 A 1/1985 Fischell
4,506,680 A 3/1985 Stokes
4,509,531 A 4/1985 Ward
4,519,973 A 5/1985 Cahalan et al.
RE31,916 E 6/1985 Oswin et al.
4,526,569 A 7/1985 Bernardi
4,527,240 A 7/1985 Kvitash
4,527,999 A 7/1985 Lee
4,534,355 A 8/1985 Potter
4,534,825 A 8/1985 Koning et al.
4,535,786 A 8/1985 Kater
4,538,616 A 9/1985 Rogoff
4,545,382 A 10/1985 Higgins et al.
4,554,927 A 11/1985 Fussell
4,561,963 A 12/1985 Owen et al.
4,565,665 A 1/1986 Fogt
4,565,666 A 1/1986 Cahalan et al.
4,568,444 A 2/1986 Nakamura et al.
4,571,292 A 2/1986 Liu et al.
4,573,968 A 3/1986 Parker
4,577,642 A 3/1986 Stokes
4,578,215 A 3/1986 Bradley
4,579,120 A 4/1986 MacGregor
4,583,976 A 4/1986 Ferguson
4,592,824 A 6/1986 Smith et al.
4,600,495 A 7/1986 Fogt
4,602,922 A 7/1986 Cabasso et al.
4,603,152 A 7/1986 Laurin et al.
4,614,514 A 9/1986 Carr et al.
4,619,793 A 10/1986 Lee
4,625,730 A 12/1986 Fountain et al.
4,626,104 A 12/1986 Pointon et al.
RE32,361 E 2/1987 Duggan
4,647,643 A 3/1987 Zdrahala et al.
4,650,547 A 3/1987 Gough
4,655,880 A 4/1987 Liu
4,663,824 A 5/1987 Kenmochi
4,671,288 A 6/1987 Gough
4,672,970 A 6/1987 Uchida et al.
4,675,346 A 6/1987 Lin et al.
4,680,268 A 7/1987 Clark, Jr.
4,684,538 A 8/1987 Klemarczyk
4,684,558 A 8/1987 Keusch et al.
4,685,463 A 8/1987 Williams
4,685,903 A 8/1987 Cable et al.
4,686,137 A 8/1987 Ward, Jr. et al.
4,689,149 A 8/1987 Kanno et al.
4,689,309 A 8/1987 Jones
4,694,861 A 9/1987 Goodale et al.
4,703,756 A 11/1987 Gough et al.
4,703,989 A 11/1987 Price et al.
4,705,503 A 11/1987 Dorman et al.
4,711,245 A 12/1987 Higgins et al.
4,711,251 A 12/1987 Stokes
4,721,677 A 1/1988 Clark, Jr.
4,726,381 A 2/1988 Jones
4,731,726 A 3/1988 Allen, III
4,734,092 A 3/1988 Millerd
4,736,748 A 4/1988 Nakamura et al.
4,739,380 A 4/1988 Lauks et al.
4,747,822 A 5/1988 Peabody
4,749,985 A 6/1988 Corsberg
4,750,496 A 6/1988 Reinhart et al.
4,752,935 A 6/1988 Beck
4,753,652 A 6/1988 Langer et al.
4,755,168 A 7/1988 Romanelli et al.
4,757,022 A 7/1988 Shults et al.
4,759,828 A 7/1988 Young et al.
4,761,748 A 8/1988 Le Rat et al.
4,763,648 A 8/1988 Wyatt
4,763,658 A 8/1988 Jones
4,776,343 A 10/1988 Hubbard et al.
4,777,953 A 10/1988 Ash et al.
4,779,618 A 10/1988 Mund et al.
4,781,680 A 11/1988 Redmond et al.
4,781,798 A 11/1988 Gough
4,784,157 A 11/1988 Halls et al.
4,786,394 A 11/1988 Enzer et al.
4,787,398 A 11/1988 Garcia et al.
4,789,467 A 12/1988 Lindsay et al.
4,791,932 A 12/1988 Margules
4,793,555 A 12/1988 Lee et al.
4,795,435 A 1/1989 Steer
4,795,542 A 1/1989 Ross et al.
4,803,243 A 2/1989 Fujimoto et al.
4,805,624 A 2/1989 Yao et al.
4,805,625 A 2/1989 Wyler
4,807,632 A 2/1989 Liess et al.
4,808,089 A 2/1989 Buchholtz et al.
4,808,292 A 2/1989 Kessler et al.
4,809,704 A 3/1989 Sogawa et al.
4,810,243 A 3/1989 Howson
4,810,470 A 3/1989 Burkhardt et al.
4,813,424 A 3/1989 Wilkins
4,815,471 A 3/1989 Stobie
4,820,281 A 4/1989 Lawler, Jr.
4,822,336 A 4/1989 Ditraglia
4,826,706 A 5/1989 Hilker
4,828,544 A 5/1989 Lane et al.
4,830,013 A 5/1989 Maxwell
4,831,070 A 5/1989 McInally et al.
4,832,005 A 5/1989 Takamiya et al.
4,832,034 A 5/1989 Pizziconi et al.
4,834,101 A 5/1989 Collison et al.
4,838,281 A 6/1989 Rogers et al.
4,841,974 A 6/1989 Gumbrecht et al.
4,849,458 A 7/1989 Reed et al.
4,852,573 A 8/1989 Kennedy
4,852,604 A 8/1989 Wales et al.
4,854,322 A 8/1989 Ash et al.
4,858,615 A 8/1989 Meinema
4,861,454 A 8/1989 Ushizawa et al.
4,867,741 A 9/1989 Portnoy
4,871,440 A 10/1989 Nagata et al.
4,874,363 A 10/1989 Abell
4,883,057 A 11/1989 Broderick
4,883,467 A 11/1989 Franetzki et al.
4,886,070 A 12/1989 Demarest
4,886,562 A 12/1989 Pinson
4,889,528 A 12/1989 Nadai et al.
4,889,744 A 12/1989 Quaid
4,890,620 A 1/1990 Gough
4,890,621 A 1/1990 Hakky
4,894,339 A 1/1990 Hanazato et al.
4,900,305 A 2/1990 Smith et al.
4,902,294 A 2/1990 Gosserez
4,907,857 A 3/1990 Giuliani et al.
4,909,786 A 3/1990 Gijselhart et al.
4,909,908 A 3/1990 Ross et al.
4,919,114 A 4/1990 Miyazaki
4,919,141 A 4/1990 Zier et al.
4,919,649 A 4/1990 Timothy et al.
4,921,477 A 5/1990 Davis
4,921,480 A 5/1990 Sealfon
4,925,268 A 5/1990 Iyer et al.
4,925,444 A 5/1990 Orkin et al.
4,927,407 A 5/1990 Dorman
4,927,516 A 5/1990 Yamaguchi et al.
4,928,694 A 5/1990 Maxwell
4,934,369 A 6/1990 Maxwell
4,934,375 A 6/1990 Cole et al.
4,935,345 A 6/1990 Guilbeau et al.
4,940,065 A 7/1990 Tanagho et al.
4,944,299 A 7/1990 Silvian
4,946,439 A 8/1990 Eggers

(56) **References Cited**

U.S. PATENT DOCUMENTS

4,951,657 A 8/1990 Pfister et al.
4,951,669 A 8/1990 Maxwell et al.
4,952,618 A 8/1990 Olsen
4,953,552 A 9/1990 Demarzo
4,955,861 A 9/1990 Enegren et al.
4,957,483 A 9/1990 Gonser et al.
4,958,148 A 9/1990 Olson
4,960,594 A 10/1990 Honeycutt
4,961,434 A 10/1990 Stypulkowski
4,963,595 A 10/1990 Ward et al.
4,966,579 A 10/1990 Polaschegg
4,967,940 A 11/1990 Blette et al.
4,970,145 A 11/1990 Bennetto et al.
4,974,592 A 12/1990 Branco
4,974,929 A 12/1990 Curry
4,975,175 A 12/1990 Karube et al.
4,975,636 A 12/1990 Desautels
4,976,687 A 12/1990 Martin
4,979,509 A 12/1990 Hakky
4,984,929 A 1/1991 Roeck et al.
4,986,271 A 1/1991 Wilkins
4,986,671 A 1/1991 Sun et al.
4,988,341 A 1/1991 Columbus et al.
4,988,758 A 1/1991 Fukuda et al.
4,989,607 A 2/1991 Keusch et al.
4,990,231 A 2/1991 Stewart et al.
4,992,794 A 2/1991 Brouwers
4,994,026 A 2/1991 Fecondini
4,994,167 A 2/1991 Shults et al.
4,995,402 A 2/1991 Smith et al.
4,997,627 A 3/1991 Bergkuist et al.
5,000,180 A 3/1991 Kuypers et al.
5,000,194 A 3/1991 Van Den Honert et al.
5,002,054 A 3/1991 Ash et al.
5,002,055 A 3/1991 Merki et al.
5,002,572 A 3/1991 Picha
5,002,590 A 3/1991 Friesen et al.
5,006,050 A 4/1991 Cooke et al.
5,006,111 A 4/1991 Inokuchi et al.
5,007,929 A 4/1991 Quaid
5,009,251 A 4/1991 Pike et al.
5,019,096 A 5/1991 Fox, Jr. et al.
5,019,974 A 5/1991 Beckers
5,026,348 A 6/1991 Venegas
5,030,199 A 7/1991 Barwick et al.
5,030,310 A 7/1991 Wogoman
5,030,333 A 7/1991 Clark, Jr.
5,034,112 A 7/1991 Murase et al.
5,035,711 A 7/1991 Aoki et al.
5,037,497 A 8/1991 Stypulkowski
5,041,092 A 8/1991 Barwick
5,045,057 A 9/1991 Van Driessche et al.
5,045,601 A 9/1991 Capelli et al.
5,046,496 A 9/1991 Betts et al.
5,048,525 A 9/1991 Maxwell
5,050,612 A 9/1991 Matsumura
5,055,171 A 10/1991 Peck
5,055,198 A 10/1991 Shettigar
5,059,654 A 10/1991 Hou et al.
5,063,081 A 11/1991 Cozzette et al.
5,067,491 A 11/1991 Taylor, II et al.
5,068,536 A 11/1991 Rosenthal
5,077,476 A 12/1991 Rosenthal
5,082,550 A 1/1992 Rishpon et al.
5,088,981 A 2/1992 Howson et al.
5,089,112 A 2/1992 Skotheim et al.
5,089,421 A 2/1992 Dieffenbach
5,096,669 A 3/1992 Lauks et al.
5,097,834 A 3/1992 Skrabal
5,098,377 A 3/1992 Borsanyi et al.
5,101,814 A 4/1992 Palti
5,106,365 A 4/1992 Hernandez
5,108,819 A 4/1992 Heller et al.
5,109,850 A 5/1992 Blanco et al.
5,112,301 A 5/1992 Fenton, Jr. et al.
5,116,313 A 5/1992 McGregor
5,120,420 A 6/1992 Nankai et al.
5,120,813 A 6/1992 Ward, Jr.
5,122,925 A 6/1992 Inpyn
5,127,405 A 7/1992 Alcala et al.
5,130,009 A 7/1992 Marsoner et al.
5,130,231 A 7/1992 Kennedy et al.
5,135,297 A 8/1992 Valint, Jr.
5,137,028 A 8/1992 Nishimura
5,140,985 A 8/1992 Schroeder et al.
5,145,565 A 9/1992 Kater et al.
5,147,725 A 9/1992 Pinchuk
5,152,746 A 10/1992 Atkinson et al.
5,160,418 A 11/1992 Mullen
5,161,532 A 11/1992 Joseph
5,162,407 A 11/1992 Turner
5,165,406 A 11/1992 Wong
5,165,407 A 11/1992 Wilson et al.
5,171,689 A 12/1992 Kawaguri et al.
5,174,123 A 12/1992 Erickson
5,174,291 A 12/1992 Schoonen et al.
5,176,632 A 1/1993 Bernardi
5,176,658 A 1/1993 Ranford
5,176,662 A 1/1993 Bartholomew et al.
5,178,142 A 1/1993 Harjunmaa et al.
5,178,957 A 1/1993 Kolpe et al.
5,182,004 A 1/1993 Kohno
5,183,549 A 2/1993 Joseph et al.
5,188,591 A 2/1993 Dorsey, III
5,190,038 A 3/1993 Polson et al.
5,190,041 A 3/1993 Palti
5,195,963 A 3/1993 Yafuso et al.
5,198,771 A 3/1993 Fidler et al.
5,200,051 A 4/1993 Cozzette et al.
5,202,261 A 4/1993 Musho et al.
5,207,218 A 5/1993 Carpentier et al.
5,208,147 A 5/1993 Kagenow et al.
5,208,313 A 5/1993 Krishnan
5,212,050 A 5/1993 Mier et al.
5,220,917 A 6/1993 Cammilli et al.
5,220,920 A 6/1993 Gharib
5,222,980 A 6/1993 Gealow
5,224,929 A 7/1993 Remiszewski
5,225,063 A 7/1993 Gumbrecht et al.
5,232,434 A 8/1993 Inagaki et al.
5,235,003 A 8/1993 Ward et al.
5,243,696 A 9/1993 Carr et al.
5,243,982 A 9/1993 Moestl et al.
5,243,983 A 9/1993 Tarr et al.
5,246,867 A 9/1993 Lakowicz et al.
5,249,576 A 10/1993 Goldberger et al.
5,250,439 A 10/1993 Musho et al.
5,254,102 A 10/1993 Ogawa
5,261,892 A 11/1993 Bertaud et al.
5,262,035 A 11/1993 Gregg et al.
5,262,305 A 11/1993 Heller et al.
5,264,104 A 11/1993 Gregg et al.
5,264,105 A 11/1993 Gregg et al.
5,265,594 A 11/1993 Olsson et al.
5,265,999 A 11/1993 Wenschhof et al.
5,266,179 A 11/1993 Nankai et al.
5,269,891 A 12/1993 Colin
5,271,736 A 12/1993 Picha
5,271,815 A 12/1993 Wong
5,279,294 A 1/1994 Anderson et al.
5,281,319 A 1/1994 Kaneko et al.
5,282,844 A 2/1994 Stokes et al.
5,282,848 A 2/1994 Schmitt
5,284,140 A 2/1994 Allen et al.
5,284,570 A 2/1994 Savage et al.
5,285,513 A 2/1994 Kaufman et al.
5,285,792 A 2/1994 Sjoquist et al.
5,286,364 A 2/1994 Yacynych et al.
5,287,753 A 2/1994 Routh et al.
5,293,877 A 3/1994 O'Hara et al.
5,296,144 A 3/1994 Sternina et al.
5,298,022 A 3/1994 Bernardi
5,298,144 A 3/1994 Spokane
5,299,571 A 4/1994 Mastrototaro

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,302,093 A 4/1994 Owens et al.
5,302,440 A 4/1994 Davis
5,304,468 A 4/1994 Phillips et al.
5,307,263 A 4/1994 Brown
5,310,469 A 5/1994 Cunningham et al.
5,311,908 A 5/1994 Barone et al.
5,312,361 A 5/1994 Zadini et al.
5,314,471 A 5/1994 Brauker et al.
5,316,008 A 5/1994 Suga et al.
5,316,452 A 5/1994 Bogen et al.
5,318,511 A 6/1994 Riquier et al.
5,320,725 A 6/1994 Gregg et al.
5,322,063 A 6/1994 Allen et al.
5,324,322 A 6/1994 Grill, Jr. et al.
5,326,356 A 7/1994 Della Valle et al.
5,326,449 A 7/1994 Cunningham
5,328,451 A 7/1994 Davis et al.
5,330,521 A 7/1994 Cohen
5,330,634 A 7/1994 Wong et al.
5,331,555 A 7/1994 Hashimoto et al.
5,335,658 A 8/1994 Bedingham
5,336,102 A 8/1994 Cairns et al.
5,337,747 A 8/1994 Neftel
5,340,352 A 8/1994 Nakanishi et al.
5,340,722 A 8/1994 Wolfbeis et al.
5,342,409 A 8/1994 Mullett
5,342,789 A 8/1994 Chick et al.
5,343,869 A 9/1994 Pross et al.
5,344,451 A 9/1994 Dayton
5,344,454 A 9/1994 Clarke et al.
5,345,932 A 9/1994 Yafuso et al.
5,348,788 A 9/1994 White
5,352,348 A 10/1994 Young et al.
5,352,351 A 10/1994 White et al.
5,354,272 A 10/1994 Swendson et al.
5,354,449 A 10/1994 Band et al.
5,356,375 A 10/1994 Higley
5,356,378 A 10/1994 Doan
5,356,786 A 10/1994 Heller et al.
5,358,409 A 10/1994 Obara
5,360,404 A 11/1994 Novacek et al.
5,362,761 A 11/1994 Uragami et al.
5,368,028 A 11/1994 Palti
5,368,224 A 11/1994 Richardson et al.
5,368,562 A 11/1994 Blomquist et al.
5,372,133 A 12/1994 Hogen Esch et al.
5,372,135 A 12/1994 Mendelson et al.
5,372,427 A 12/1994 Padovani et al.
5,372,709 A 12/1994 Hood
5,372,719 A 12/1994 Afeyan et al.
5,376,070 A 12/1994 Purvis et al.
5,378,229 A 1/1995 Layer et al.
5,379,238 A 1/1995 Stark
5,380,268 A 1/1995 Wheeler
5,380,422 A 1/1995 Negishi et al.
5,380,491 A 1/1995 Carver, Jr. et al.
5,380,536 A 1/1995 Hubbell et al.
5,380,665 A 1/1995 Cusack et al.
5,382,514 A 1/1995 Passaniti et al.
5,384,028 A 1/1995 Ito
5,387,327 A 2/1995 Khan
5,390,671 A 2/1995 Lord et al.
5,391,250 A 2/1995 Cheney, II et al.
5,393,401 A 2/1995 Knoll
5,397,848 A 3/1995 Yang et al.
5,400,267 A 3/1995 Denen et al.
5,404,877 A 4/1995 Nolan et al.
5,405,510 A 4/1995 Betts et al.
5,408,999 A 4/1995 Singh et al.
5,411,052 A 5/1995 Murray
5,411,647 A 5/1995 Johnson et al.
5,411,866 A 5/1995 Luong et al.
5,417,115 A 5/1995 Burns
5,417,206 A 5/1995 Kaneyoshi
5,417,395 A 5/1995 Fowler et al.
5,421,328 A 6/1995 Bedingham
5,421,923 A 6/1995 Clarke et al.
5,423,738 A 6/1995 Robinson et al.
5,423,749 A 6/1995 Merte et al.
5,425,717 A 6/1995 Mohiuddin
5,426,032 A 6/1995 Phillips et al.
5,428,123 A 6/1995 Ward et al.
5,429,129 A 7/1995 Lovejoy et al.
5,429,485 A 7/1995 Dodge
5,429,602 A 7/1995 Hauser
5,429,735 A 7/1995 Johnson et al.
5,431,160 A 7/1995 Wilkins
5,431,174 A 7/1995 Knute
5,431,921 A 7/1995 Thombre
5,434,412 A 7/1995 Sodickson et al.
5,437,635 A 8/1995 Fields et al.
5,437,999 A 8/1995 Diebold et al.
5,438,984 A 8/1995 Schoendorfer
5,445,610 A 8/1995 Evert
5,448,992 A 9/1995 Kupershmidt
5,451,260 A 9/1995 Versteeg et al.
5,453,199 A 9/1995 Afeyan et al.
5,453,278 A 9/1995 Chan et al.
5,458,631 A 10/1995 Xavier
5,462,051 A 10/1995 Oka et al.
5,462,064 A 10/1995 D'Angelo et al.
5,462,645 A 10/1995 Albery et al.
5,466,356 A 11/1995 Schneider et al.
5,466,575 A 11/1995 Cozzette et al.
5,469,846 A 11/1995 Khan
5,472,317 A 12/1995 Field et al.
5,474,552 A 12/1995 Palti
5,476,094 A 12/1995 Allen et al.
5,476,776 A 12/1995 Wilkins
5,480,711 A 1/1996 Ruefer
5,482,008 A 1/1996 Stafford et al.
5,482,446 A 1/1996 Williamson et al.
5,482,473 A 1/1996 Lord et al.
5,484,404 A 1/1996 Schulman et al.
5,486,776 A 1/1996 Chiang
5,489,414 A 2/1996 Schreiber et al.
5,490,323 A 2/1996 Thacker et al.
5,491,474 A 2/1996 Suni et al.
5,494,562 A 2/1996 Maley et al.
5,496,453 A 3/1996 Uenoyama et al.
5,497,772 A 3/1996 Schulman et al.
5,502,396 A 3/1996 Desarzens et al.
5,505,828 A 4/1996 Wong et al.
5,507,288 A 4/1996 Boecker et al.
5,508,203 A 4/1996 Fuller et al.
5,508,509 A 4/1996 Yafuso et al.
5,509,410 A 4/1996 Hill et al.
5,509,888 A 4/1996 Miller
5,512,046 A 4/1996 Pusinelli et al.
5,512,248 A 4/1996 Van
5,513,636 A 5/1996 Palti
5,514,253 A 5/1996 Davis et al.
5,514,718 A 5/1996 Lewis et al.
5,515,851 A 5/1996 Goldstein
5,516,832 A 5/1996 Kennan et al.
5,518,601 A 5/1996 Foos et al.
5,527,288 A 6/1996 Gross et al.
5,529,066 A 6/1996 Palti
5,529,676 A 6/1996 Maley et al.
5,531,679 A 7/1996 Schulman et al.
5,531,878 A 7/1996 Vadgama et al.
5,540,828 A 7/1996 Yacynych
5,545,200 A 8/1996 West et al.
5,545,220 A 8/1996 Andrews et al.
5,545,223 A 8/1996 Neuenfeldt et al.
5,549,547 A 8/1996 Cohen et al.
5,549,548 A 8/1996 Larsson
5,549,569 A 8/1996 Lynn et al.
5,549,651 A 8/1996 Lynn
5,549,675 A 8/1996 Neuenfeldt et al.
5,551,850 A 9/1996 Williamson et al.
5,552,112 A 9/1996 Schiffmann et al.
5,553,616 A 9/1996 Ham et al.
5,554,339 A 9/1996 Cozzette et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,558,957 A 9/1996 Datta et al.
5,562,614 A 10/1996 O'Donnell
5,562,615 A 10/1996 Nassif
5,564,439 A 10/1996 Picha
5,568,806 A 10/1996 Cheney, II et al.
5,569,186 A 10/1996 Lord et al.
5,569,188 A 10/1996 Mackool
5,569,219 A 10/1996 Hakki et al.
5,569,462 A 10/1996 Martinson et al.
5,571,395 A 11/1996 Park et al.
5,575,293 A 11/1996 Miller et al.
5,575,930 A 11/1996 Tietje-Girault et al.
5,577,499 A 11/1996 Teves
5,582,184 A 12/1996 Erickson et al.
5,582,497 A 12/1996 Noguchi
5,582,593 A 12/1996 Hultman
5,582,697 A 12/1996 Ikeda et al.
5,584,813 A 12/1996 Livingston et al.
5,584,876 A 12/1996 Bruchman et al.
5,586,553 A 12/1996 Halili et al.
5,587,273 A 12/1996 Yan et al.
5,588,560 A 12/1996 Benedict et al.
5,589,133 A 12/1996 Suzuki
5,589,563 A 12/1996 Ward et al.
5,590,651 A 1/1997 Shaffer et al.
5,593,440 A 1/1997 Brauker et al.
5,593,852 A 1/1997 Heller et al.
5,601,435 A 2/1997 Quy
5,607,565 A 3/1997 Azarnia et al.
5,609,572 A 3/1997 Lang
5,609,575 A 3/1997 Larson et al.
5,611,900 A 3/1997 Worden et al.
5,624,409 A 4/1997 Seale
5,624,537 A 4/1997 Turner et al.
5,626,561 A 5/1997 Butler et al.
5,626,563 A 5/1997 Dodge et al.
5,628,310 A 5/1997 Rao et al.
5,628,619 A 5/1997 Wilson
5,628,890 A 5/1997 Carter et al.
5,630,978 A 5/1997 Domb
5,637,083 A 6/1997 Bertrand et al.
5,637,135 A 6/1997 Ottenstein et al.
5,640,470 A 6/1997 Iyer et al.
5,640,954 A 6/1997 Pfeiffer et al.
5,643,195 A 7/1997 Drevet et al.
5,651,767 A 7/1997 Schulman et al.
5,653,239 A 8/1997 Pompei et al.
5,653,756 A 8/1997 Clarke et al.
5,653,863 A 8/1997 Genshaw et al.
5,658,247 A 8/1997 Henley
5,658,250 A 8/1997 Blomquist et al.
5,658,330 A 8/1997 Carlisle et al.
5,658,802 A 8/1997 Hayes et al.
5,660,163 A 8/1997 Schulman et al.
5,660,177 A 8/1997 Faupel et al.
5,662,616 A 9/1997 Bousquet
5,665,061 A 9/1997 Antwiler
5,665,065 A 9/1997 Colman et al.
5,665,215 A 9/1997 Bussmann et al.
5,665,222 A 9/1997 Heller et al.
5,667,504 A 9/1997 Baumann et al.
5,673,694 A 10/1997 Rivers
5,676,651 A 10/1997 Larson, Jr. et al.
5,676,820 A 10/1997 Wang et al.
5,681,572 A 10/1997 Seare, Jr.
5,682,884 A 11/1997 Hill et al.
5,683,562 A 11/1997 Schaffar et al.
5,686,829 A 11/1997 Girault
5,688,239 A 11/1997 Walker
5,688,244 A 11/1997 Lang
5,690,119 A 11/1997 Rytky et al.
5,695,623 A 12/1997 Michel et al.
5,696,314 A 12/1997 McCaffrey et al.
5,697,366 A 12/1997 Kimball et al.
5,697,899 A 12/1997 Hillman et al.
5,697,976 A 12/1997 Chesterfield et al.
5,703,359 A 12/1997 Wampler, III
5,704,354 A 1/1998 Preidel et al.
5,706,807 A 1/1998 Picha
5,707,502 A 1/1998 McCaffrey et al.
5,711,001 A 1/1998 Bussan et al.
5,711,302 A 1/1998 Lampropoulos et al.
5,711,685 A 1/1998 Wood
5,711,861 A 1/1998 Ward et al.
5,713,842 A 2/1998 Kay
5,713,888 A 2/1998 Neuenfeldt et al.
5,714,123 A 2/1998 Sohrab
5,714,391 A 2/1998 Omura et al.
5,720,293 A 2/1998 Quinn et al.
5,730,654 A 3/1998 Brown
5,733,336 A 3/1998 Neuenfeldt et al.
5,735,273 A 4/1998 Kurnik et al.
5,735,285 A 4/1998 Albert et al.
5,741,319 A 4/1998 Woloszko et al.
5,741,330 A 4/1998 Brauker et al.
5,741,634 A 4/1998 Nozoe et al.
5,743,262 A 4/1998 Lepper, Jr. et al.
5,746,898 A 5/1998 Preidel
5,749,832 A 5/1998 Vadgama et al.
5,749,907 A 5/1998 Mann
5,755,692 A 5/1998 Manicom
5,756,632 A 5/1998 Ward et al.
5,758,643 A 6/1998 Wong et al.
5,763,760 A 6/1998 Gumbrecht et al.
5,763,787 A 6/1998 Gravel et al.
5,766,151 A 6/1998 Valley et al.
5,770,028 A 6/1998 Maley et al.
5,771,890 A 6/1998 Tamada
5,772,586 A 6/1998 Heinonen et al.
5,773,270 A 6/1998 D'Orazio et al.
5,773,286 A 6/1998 Dionne et al.
5,776,324 A 7/1998 Usala
5,777,060 A 7/1998 Van Antwerp
5,779,665 A 7/1998 Mastrototaro et al.
5,781,455 A 7/1998 Hyodo
5,782,880 A 7/1998 Lahtinen et al.
5,782,912 A 7/1998 Brauker et al.
5,783,054 A 7/1998 Raguse et al.
5,786,439 A 7/1998 Van Antwerp et al.
5,787,900 A 8/1998 Butler et al.
5,791,344 A 8/1998 Schulman et al.
5,791,880 A 8/1998 Wilson
5,795,453 A 8/1998 Gilmartin
5,795,774 A 8/1998 Matsumoto et al.
5,798,065 A 8/1998 Picha
5,798,085 A 8/1998 Seaton et al.
5,800,383 A 9/1998 Chandler et al.
5,800,420 A 9/1998 Gross et al.
5,800,529 A 9/1998 Brauker et al.
5,804,048 A 9/1998 Wong et al.
5,806,517 A 9/1998 Gerhardt et al.
5,807,274 A 9/1998 Henning et al.
5,807,312 A 9/1998 Dzwonkiewicz
5,807,375 A 9/1998 Gross et al.
5,807,406 A 9/1998 Brauker et al.
5,810,736 A 9/1998 Pail
5,810,770 A 9/1998 Chin et al.
5,811,487 A 9/1998 Schulz, Jr. et al.
5,814,599 A 9/1998 Mitragotri et al.
5,820,570 A 10/1998 Erickson et al.
5,820,589 A 10/1998 Torgerson et al.
5,820,622 A 10/1998 Gross et al.
5,822,715 A 10/1998 Worthington et al.
5,823,802 A 10/1998 Bartley
5,824,651 A 10/1998 Nanci et al.
5,827,183 A 10/1998 Kurnik et al.
5,833,603 A 11/1998 Kovacs et al.
5,836,886 A 11/1998 Itoigawa et al.
5,836,887 A 11/1998 Oka et al.
5,836,989 A 11/1998 Shelton
5,837,454 A 11/1998 Cozzette et al.
5,837,728 A 11/1998 Purcell
5,840,026 A 11/1998 Uber, III et al.
5,840,148 A 11/1998 Campbell et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,840,240 A 11/1998 Stenoien et al.
5,843,069 A 12/1998 Butler et al.
5,848,991 A 12/1998 Gross et al.
5,851,197 A 12/1998 Marano et al.
5,857,983 A 1/1999 Douglas et al.
5,858,296 A 1/1999 Domb
5,858,365 A 1/1999 Faller
5,858,747 A 1/1999 Schinstine et al.
5,861,019 A 1/1999 Sun et al.
5,863,400 A 1/1999 Drummond et al.
5,863,460 A 1/1999 Slovacek et al.
5,871,499 A 2/1999 Hahn et al.
5,871,514 A 2/1999 Wiklund et al.
5,872,198 A 2/1999 Mosbach et al.
5,872,499 A 2/1999 Poulsen
5,873,862 A 2/1999 Lopez
5,874,500 A 2/1999 Rhee et al.
5,879,294 A 3/1999 Anderson et al.
5,879,373 A 3/1999 Roper et al.
5,879,713 A 3/1999 Roth et al.
5,879,828 A 3/1999 Debe et al.
5,882,354 A 3/1999 Brauker et al.
5,882,494 A 3/1999 Van Antwerp
5,895,235 A 4/1999 Droz
5,897,525 A 4/1999 Dey et al.
5,897,578 A 4/1999 Wiklund et al.
5,897,955 A 4/1999 Drumheller
5,899,855 A 5/1999 Brown
5,904,666 A 5/1999 Dedecker et al.
5,904,708 A 5/1999 Goedeke
5,910,554 A 6/1999 Kempe et al.
5,911,219 A 6/1999 Aylsworth et al.
5,913,998 A 6/1999 Butler et al.
5,914,026 A 6/1999 Blubaugh, Jr. et al.
5,914,182 A 6/1999 Drumheller
5,916,445 A 6/1999 Hjerten et al.
5,917,346 A 6/1999 Gord
5,919,215 A 7/1999 Wiklund et al.
5,921,951 A 7/1999 Morris
5,925,021 A 7/1999 Castellano et al.
5,928,130 A 7/1999 Schmidt
5,928,155 A 7/1999 Eggers et al.
5,928,182 A 7/1999 Kraus et al.
5,928,189 A 7/1999 Phillips et al.
5,928,195 A 7/1999 Malamud et al.
5,931,814 A 8/1999 Alex et al.
5,932,175 A 8/1999 Knute et al.
5,933,136 A 8/1999 Brown
5,935,785 A 8/1999 Reber et al.
5,938,636 A 8/1999 Kramer et al.
5,938,679 A 8/1999 Freeman et al.
5,942,979 A 8/1999 Luppino
5,944,661 A 8/1999 Swette et al.
5,945,498 A 8/1999 Hoepken et al.
5,947,911 A 9/1999 Wong et al.
5,951,521 A 9/1999 Mastrototaro et al.
5,954,643 A 9/1999 Van Antwerp et al.
5,954,685 A 9/1999 Tierney
5,954,954 A 9/1999 Houck et al.
5,957,854 A 9/1999 Besson et al.
5,957,903 A 9/1999 Mirzaee et al.
5,959,050 A 9/1999 Mosbach et al.
5,961,451 A 10/1999 Reber et al.
5,963,132 A 10/1999 Yoakum
5,964,261 A 10/1999 Neuenfeldt et al.
5,964,745 A 10/1999 Lyles et al.
5,964,804 A 10/1999 Brauker et al.
5,964,993 A 10/1999 Blubaugh, Jr. et al.
5,965,125 A 10/1999 Mineau-Hanschke
5,965,380 A 10/1999 Heller et al.
5,967,986 A 10/1999 Cimochowski et al.
5,971,922 A 10/1999 Arita et al.
5,972,199 A 10/1999 Heller et al.
5,976,085 A 11/1999 Kimball et al.
5,977,241 A 11/1999 Koloski et al.
5,985,129 A 11/1999 Gough et al.
5,985,693 A 11/1999 Leedy
5,987,352 A 11/1999 Klein et al.
5,989,409 A 11/1999 Kurnik et al.
5,995,208 A 11/1999 Sarge et al.
5,995,860 A 11/1999 Sun et al.
5,997,501 A 12/1999 Gross et al.
5,998,247 A 12/1999 Wu
5,999,848 A 12/1999 Gord et al.
5,999,849 A 12/1999 Gord et al.
6,001,067 A 12/1999 Shults et al.
6,001,471 A 12/1999 Bries et al.
6,007,845 A 12/1999 Domb et al.
6,011,984 A 1/2000 Van Antwerp et al.
6,013,113 A 1/2000 Mika
6,014,577 A 1/2000 Henning et al.
6,015,392 A 1/2000 Douglas et al.
6,015,572 A 1/2000 Lin et al.
6,016,448 A 1/2000 Busacker et al.
6,017,435 A 1/2000 Hassard et al.
6,018,033 A 1/2000 Chen et al.
6,023,629 A 2/2000 Tamada
6,024,699 A 2/2000 Surwit et al.
6,024,720 A 2/2000 Chandler et al.
6,027,445 A 2/2000 Von Bahr
6,027,479 A 2/2000 Alei et al.
6,032,059 A 2/2000 Henning et al.
6,032,667 A 3/2000 Heinonen
6,036,924 A 3/2000 Simons et al.
6,043,328 A 3/2000 Domschke et al.
6,045,671 A 4/2000 Wu et al.
6,048,691 A 4/2000 Maracas
6,049,727 A 4/2000 Crothall
6,051,372 A 4/2000 Bayerl et al.
6,051,389 A 4/2000 Ahl et al.
6,057,377 A 5/2000 Sasaki et al.
6,059,946 A 5/2000 Yukawa et al.
6,060,640 A 5/2000 Pauley et al.
6,063,637 A 5/2000 Arnold et al.
6,065,154 A 5/2000 Hulings et al.
6,066,083 A 5/2000 Slater et al.
6,066,088 A 5/2000 Davis
6,066,448 A 5/2000 Wohlstadter et al.
6,068,668 A 5/2000 Mastroianni
6,071,391 A 6/2000 Gotoh et al.
6,071,406 A 6/2000 Tsou
6,074,775 A 6/2000 Gartstein et al.
6,077,299 A 6/2000 Adelberg et al.
6,080,583 A 6/2000 Von Bahr
6,081,735 A 6/2000 Diab et al.
6,081,736 A 6/2000 Colvin et al.
6,082,289 A 7/2000 Cavallaro
6,083,523 A 7/2000 Dionne et al.
6,083,710 A 7/2000 Heller et al.
6,088,608 A 7/2000 Schulman et al.
6,090,087 A 7/2000 Tsukada et al.
6,091,975 A 7/2000 Daddona et al.
6,091,976 A 7/2000 Pfeiffer et al.
6,093,156 A 7/2000 Cunningham et al.
6,093,172 A 7/2000 Funderburk et al.
6,099,511 A 8/2000 Devos et al.
6,101,404 A 8/2000 Yoon et al.
6,103,033 A 8/2000 Say et al.
6,103,533 A 8/2000 Hassard et al.
6,106,486 A 8/2000 Tenerz et al.
6,107,083 A 8/2000 Collins et al.
6,115,622 A 9/2000 Minoz
6,115,634 A 9/2000 Donders et al.
6,117,290 A 9/2000 Say et al.
6,119,028 A 9/2000 Schulman et al.
6,120,676 A 9/2000 Heller et al.
6,121,009 A 9/2000 Heller et al.
6,121,611 A 9/2000 Lindsay et al.
6,122,351 A 9/2000 Schlueter, Jr. et al.
6,122,536 A 9/2000 Sun et al.
6,123,827 A 9/2000 Wong et al.
6,127,154 A 10/2000 Mosbach et al.
6,128,519 A 10/2000 Say
6,134,461 A 10/2000 Say et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

6,135,978 A 10/2000 Houben et al.
6,139,718 A 10/2000 Kurnik et al.
6,141,573 A 10/2000 Kurnik et al.
6,142,939 A 11/2000 Eppstein et al.
6,144,869 A 11/2000 Berner et al.
6,144,871 A 11/2000 Saito et al.
RE36,991 E 12/2000 Yamamoto et al.
6,156,051 A 12/2000 Schraga
6,157,860 A 12/2000 Hauser et al.
6,159,147 A 12/2000 Lichter et al.
6,159,186 A 12/2000 Wickham et al.
6,159,497 A 12/2000 Laprade et al.
6,162,201 A 12/2000 Cohen et al.
6,162,611 A 12/2000 Heller et al.
6,164,921 A 12/2000 Moubayed et al.
6,165,154 A 12/2000 Gray et al.
6,165,156 A 12/2000 Cesarczyk et al.
6,167,614 B1 1/2001 Tuttle et al.
6,168,568 B1 1/2001 Gavriely
6,169,155 B1 1/2001 Alvarez et al.
6,171,276 B1 1/2001 Lippe et al.
6,175,752 B1 1/2001 Say et al.
6,175,767 B1 1/2001 Doyle, Sr.
6,180,221 B1 1/2001 Crotzer et al.
6,180,416 B1 1/2001 Kurnik et al.
6,183,437 B1 2/2001 Walker
6,187,062 B1 2/2001 Oweis et al.
6,189,536 B1 2/2001 Martinez et al.
6,191,860 B1 2/2001 Klinger et al.
6,192,891 B1 2/2001 Gravel et al.
6,198,969 B1 3/2001 Kuzma
6,200,265 B1 3/2001 Walsh et al.
6,200,772 B1 3/2001 Vadgama et al.
6,201,979 B1 3/2001 Kurnik et al.
6,201,980 B1 3/2001 Darrow et al.
6,201,993 B1 3/2001 Kruse et al.
6,206,856 B1 3/2001 Mahurkar
6,208,894 B1 3/2001 Schulman et al.
6,212,416 B1 4/2001 Ward et al.
6,212,424 B1 4/2001 Robinson
6,213,739 B1 4/2001 Phallen et al.
6,214,185 B1 4/2001 Offenbacher et al.
6,214,901 B1 4/2001 Chudzik et al.
6,219,574 B1 4/2001 Cormier et al.
6,223,080 B1 4/2001 Thompson
6,223,083 B1 4/2001 Rosar
6,230,059 B1 5/2001 Duffin
6,231,879 B1 5/2001 Li et al.
6,232,783 B1 5/2001 Merrill
6,233,080 B1 5/2001 Brenner et al.
6,233,471 B1 5/2001 Berner et al.
6,241,663 B1 6/2001 Wu et al.
6,241,863 B1 6/2001 Monbouquette
6,248,067 B1 6/2001 Causey, III et al.
6,248,077 B1 6/2001 Elson et al.
6,248,093 B1 6/2001 Moberg
6,251,280 B1 6/2001 Dai et al.
6,254,586 B1 7/2001 Mann et al.
6,255,592 B1 7/2001 Pennington et al.
6,256,522 B1 7/2001 Schultz
6,259,937 B1 7/2001 Schulman et al.
6,263,222 B1 7/2001 Diab et al.
6,264,825 B1 7/2001 Blackburn et al.
6,266,551 B1 7/2001 Osadchy et al.
6,268,161 B1 7/2001 Han et al.
6,270,478 B1 8/2001 Mernoee
6,271,332 B1 8/2001 Lohmann et al.
6,272,364 B1 8/2001 Kurnik
6,272,382 B1 8/2001 Faltys et al.
6,272,480 B1 8/2001 Tresp et al.
6,274,285 B1 8/2001 Gries et al.
6,274,686 B1 8/2001 Mosbach et al.
6,275,717 B1 8/2001 Gross et al.
6,280,408 B1 8/2001 Sipin
6,283,761 B1 9/2001 Joao
6,284,126 B1 9/2001 Kurnik et al.
6,284,478 B1 9/2001 Heller et al.
6,285,897 B1 9/2001 Kilcoyne et al.
6,293,925 B1 9/2001 Safabash et al.
6,294,281 B1 9/2001 Heller
6,295,506 B1 9/2001 Heinonen et al.
6,296,615 B1 10/2001 Brockway et al.
6,298,254 B2 10/2001 Tamada
6,298,255 B1 10/2001 Cordero et al.
6,299,578 B1 10/2001 Kurnik et al.
6,299,583 B1 10/2001 Eggers et al.
6,300,002 B1 10/2001 Webb et al.
6,300,884 B1 10/2001 Wilson
6,302,855 B1 10/2001 Lav et al.
6,306,104 B1 10/2001 Cunningham et al.
6,306,424 B1 10/2001 Vyakarnam et al.
6,306,594 B1 10/2001 Cozzette et al.
6,308,089 B1 10/2001 Von Der Ruhr et al.
6,309,351 B1 10/2001 Kurnik et al.
6,309,384 B1 10/2001 Harrington et al.
6,309,526 B1 10/2001 Fujiwara et al.
6,309,884 B1 10/2001 Cooper et al.
6,310,110 B1 10/2001 Markowitz et al.
6,315,738 B1 11/2001 Nishikawa et al.
6,319,566 B1 11/2001 Polanyi et al.
6,325,978 B1 12/2001 Labuda et al.
6,325,979 B1 12/2001 Hahn et al.
6,326,160 B1 12/2001 Dunn et al.
6,329,161 B1 12/2001 Heller et al.
6,329,488 B1 12/2001 Terry et al.
6,329,929 B1 12/2001 Weijand et al.
6,330,464 B1 12/2001 Colvin, Jr. et al.
6,336,269 B1 1/2002 Eldridge et al.
6,340,588 B1 1/2002 Nova et al.
6,341,232 B1 1/2002 Conn et al.
6,343,225 B1 1/2002 Clark, Jr.
6,344,021 B1 2/2002 Juster et al.
6,346,114 B1 2/2002 Schraga
6,348,640 B1 2/2002 Navot et al.
6,355,000 B1 3/2002 Ogura
6,356,776 B1 3/2002 Berner et al.
6,358,225 B1 3/2002 Butterfield
6,359,444 B1 3/2002 Grimes
6,360,888 B1 3/2002 McIvor et al.
6,365,670 B1 4/2002 Fry
6,366,794 B1 4/2002 Moussy et al.
6,368,141 B1 4/2002 Van Antwerp et al.
6,368,274 B1 4/2002 Van Antwerp et al.
6,368,658 B1 4/2002 Schwarz et al.
6,370,410 B2 4/2002 Kurnik et al.
6,370,941 B2 4/2002 Nakamura et al.
6,371,963 B1 4/2002 Nishtala et al.
6,372,244 B1 4/2002 Antanavich et al.
6,377,828 B1 4/2002 Chaiken et al.
6,379,201 B1 4/2002 Biggs et al.
6,379,301 B1 4/2002 Worthington et al.
6,379,317 B1 4/2002 Kintzig et al.
6,379,883 B2 4/2002 Davis et al.
6,387,709 B1 5/2002 Mason et al.
6,391,019 B1 5/2002 Ito
6,393,318 B1 5/2002 Conn et al.
6,398,562 B1 6/2002 Butler et al.
6,400,974 B1 6/2002 Lesho
6,402,703 B1 6/2002 Kensey et al.
6,403,944 B1 6/2002 MacKenzie et al.
6,405,066 B1 6/2002 Essenpreis et al.
6,406,066 B1 6/2002 Uegane
6,406,426 B1 6/2002 Reuss et al.
6,407,195 B2 6/2002 Sherman et al.
6,409,674 B1 6/2002 Brockway et al.
6,413,393 B1 7/2002 Van Antwerp et al.
6,416,651 B1 7/2002 Millar
6,418,332 B1 7/2002 Mastrototaro et al.
6,418,346 B1 7/2002 Nelson et al.
6,424,847 B1 7/2002 Mastrototaro et al.
6,427,088 B1 7/2002 Bowman, IV et al.
6,430,437 B1 8/2002 Marro
6,432,050 B1 8/2002 Porat et al.
6,438,414 B1 8/2002 Conn et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

6,440,068 B1 8/2002 Brown et al.
6,442,413 B1 8/2002 Silver
6,447,448 B1 9/2002 Ishikawa et al.
6,447,542 B1 9/2002 Weadock
6,454,710 B1 9/2002 Ballerstadt et al.
6,459,917 B1 10/2002 Gowda et al.
6,461,496 B1 10/2002 Feldman et al.
6,464,849 B1 10/2002 Say et al.
6,465,066 B1 10/2002 Rule et al.
6,466,810 B1 10/2002 Ward et al.
6,467,480 B1 10/2002 Meier et al.
6,468,287 B1 10/2002 Baugh
6,471,689 B1 10/2002 Joseph et al.
6,471,993 B1 10/2002 Shastri et al.
6,474,360 B1 11/2002 Ito
6,475,372 B1 11/2002 Ohara et al.
6,475,750 B1 11/2002 Han et al.
6,477,392 B1 11/2002 Honigs et al.
6,477,395 B2 11/2002 Schulman et al.
6,478,736 B1 11/2002 Mault
6,481,440 B2 11/2002 Gielen et al.
6,484,045 B1 11/2002 Holker et al.
6,484,046 B1 11/2002 Say et al.
6,484,132 B1 11/2002 Hively et al.
6,485,449 B2 11/2002 Ito
6,487,429 B2 11/2002 Hockersmith et al.
6,488,652 B1 12/2002 Weijand et al.
6,494,830 B1 12/2002 Wessel
6,494,879 B2 12/2002 Lennox et al.
6,497,655 B1 12/2002 Linberg et al.
6,497,729 B1 12/2002 Moussy et al.
6,498,043 B1 12/2002 Schulman et al.
6,498,941 B1 12/2002 Jackson
6,501,976 B1 12/2002 Sohrab
6,510,239 B1 1/2003 Wieres et al.
6,510,329 B2 1/2003 Heckel
6,512,939 B1 1/2003 Colvin et al.
6,514,689 B2 2/2003 Han et al.
6,514,718 B2 2/2003 Heller et al.
6,517,508 B1 2/2003 Utterberg et al.
6,520,326 B2 2/2003 McIvor et al.
6,520,477 B2 2/2003 Trimmer
6,520,937 B2 2/2003 Hart et al.
6,520,938 B1 2/2003 Funderburk et al.
6,520,997 B1 2/2003 Pekkarinen et al.
6,522,903 B1 2/2003 Berman et al.
6,522,927 B1 2/2003 Bishay et al.
6,524,861 B1 2/2003 Anderson
6,526,298 B1 2/2003 Khalil et al.
6,527,729 B1 3/2003 Turcott
6,528,584 B2 3/2003 Kennedy et al.
6,529,755 B2 3/2003 Kurnik et al.
6,534,711 B1 3/2003 Pollack
6,536,433 B1 3/2003 Cewers
6,537,318 B1 3/2003 Ita et al.
6,541,107 B1 4/2003 Zhong et al.
6,541,266 B2 4/2003 Modzelewski et al.
6,542,765 B1 4/2003 Guy et al.
6,544,212 B2 4/2003 Galley et al.
6,545,085 B2 4/2003 Kilgour et al.
6,546,268 B1 4/2003 Ishikawa et al.
6,546,269 B1 4/2003 Kurnik
6,547,839 B2 4/2003 Zhang et al.
6,549,796 B2 4/2003 Sohrab
6,551,494 B1 4/2003 Heller et al.
6,551,496 B1 4/2003 Moles et al.
6,553,241 B2 4/2003 Mannheimer et al.
6,553,244 B2 4/2003 Lesho et al.
6,554,805 B2 4/2003 Hiejima
6,554,822 B1 4/2003 Holschneider et al.
6,558,320 B1 5/2003 Causey, III et al.
6,558,321 B1 5/2003 Burd et al.
6,558,347 B1 5/2003 Jhuboo et al.
6,558,351 B1 5/2003 Steil et al.
6,560,471 B1 5/2003 Heller et al.
6,561,978 B1 5/2003 Conn et al.
6,562,001 B2 5/2003 Lebel et al.
6,564,105 B2 5/2003 Starkweather et al.
6,565,509 B1 5/2003 Say et al.
6,565,535 B2 5/2003 Zaias et al.
6,565,807 B1 5/2003 Patterson et al.
6,569,309 B2 5/2003 Otsuka et al.
6,569,521 B1 5/2003 Sheridan et al.
6,571,128 B2 5/2003 Lebel et al.
6,572,545 B2 6/2003 Knobbe et al.
6,572,579 B1 6/2003 Raghavan et al.
6,572,745 B2 6/2003 Rappin et al.
6,574,490 B2 6/2003 Abbink et al.
6,575,905 B2 6/2003 Knobbe et al.
6,576,101 B1 6/2003 Heller et al.
6,577,899 B2 6/2003 Lebel et al.
6,579,257 B1 6/2003 Elgas et al.
6,579,498 B1 6/2003 Eglise
6,579,690 B1 6/2003 Bonnecaze et al.
6,584,335 B1 6/2003 Haar et al.
6,585,644 B2 7/2003 Lebel et al.
6,585,675 B1 7/2003 O'Mahony et al.
6,585,763 B1 7/2003 Keilman et al.
6,587,704 B1 7/2003 Fine et al.
6,587,705 B1 7/2003 Kim et al.
6,589,229 B1 7/2003 Connelly et al.
6,591,123 B2 7/2003 Fein et al.
6,591,125 B1 7/2003 Buse et al.
6,592,745 B1 7/2003 Feldman et al.
6,594,514 B2 7/2003 Berner et al.
6,595,756 B2 7/2003 Gray et al.
6,595,919 B2 7/2003 Berner et al.
6,602,221 B1 8/2003 Saravia et al.
6,603,995 B1 8/2003 Carter
6,605,072 B2 8/2003 Struys et al.
6,605,200 B1 8/2003 Mao et al.
6,605,201 B1 8/2003 Mao et al.
6,607,509 B2 8/2003 Bobroff et al.
6,607,658 B1 8/2003 Heller et al.
6,609,071 B2 8/2003 Shapiro et al.
6,610,012 B2 8/2003 Mault
6,612,984 B1 9/2003 Kerr, II
6,613,379 B2 9/2003 Ward et al.
6,615,061 B1 9/2003 Khalil et al.
6,615,078 B1 9/2003 Burson et al.
6,618,603 B2 9/2003 Varalli et al.
6,618,934 B1 9/2003 Feldman et al.
6,628,975 B1 9/2003 Fein et al.
6,633,772 B2 10/2003 Ford et al.
6,635,014 B2 10/2003 Starkweather et al.
6,641,533 B2 11/2003 Causey, III et al.
6,642,015 B2 11/2003 Vachon et al.
6,645,181 B1 11/2003 Lavi et al.
6,645,219 B2 11/2003 Roe
6,645,359 B1 11/2003 Bhullar et al.
6,648,821 B2 11/2003 Lebel et al.
6,653,091 B1 11/2003 Dunn et al.
6,654,625 B1 11/2003 Say et al.
6,656,157 B1 12/2003 Duchonet al.
6,659,948 B2 12/2003 Lebel et al.
6,662,439 B1 12/2003 Bhullar
6,663,615 B1 12/2003 Madou et al.
6,666,821 B2 12/2003 Keimel
6,668,196 B1 12/2003 Villegas et al.
6,673,022 B1 1/2004 Bobo et al.
6,673,596 B1 1/2004 Sayler et al.
6,679,865 B2 1/2004 Shekalim
6,683,535 B1 1/2004 Utke
6,684,904 B2 2/2004 Ito
6,685,668 B1 2/2004 Cho et al.
6,687,522 B2 2/2004 Tamada
6,687,546 B2 2/2004 Lebel et al.
6,689,056 B1 2/2004 Kilcoyne et al.
6,689,089 B1 2/2004 Tiedtke et al.
6,689,265 B2 2/2004 Heller et al.
6,692,456 B1 2/2004 Eppstein et al.
6,694,191 B2 2/2004 Starkweather et al.
6,695,860 B1 2/2004 Ward et al.
6,699,188 B2 3/2004 Wessel

(56) **References Cited**

U.S. PATENT DOCUMENTS

6,699,218 B2 3/2004 Flaherty et al.
6,699,383 B2 3/2004 Lemire et al.
6,702,249 B2 3/2004 Ito
6,702,857 B2 3/2004 Brauker et al.
6,702,972 B1 3/2004 Markle
6,705,833 B2 3/2004 Tam et al.
6,711,424 B1 3/2004 Fine et al.
6,712,796 B2 3/2004 Fentis et al.
6,713,773 B1 3/2004 Lyons et al.
6,721,586 B2 4/2004 Kiser et al.
6,721,587 B2 4/2004 Gough
6,723,077 B2 4/2004 Pickup et al.
6,723,086 B2 4/2004 Bassuk et al.
6,730,072 B2 5/2004 Shawgo et al.
6,730,200 B1 5/2004 Stewart et al.
6,731,976 B2 5/2004 Penn et al.
6,733,446 B2 5/2004 Lebel et al.
6,733,655 B1 5/2004 Davies et al.
6,736,777 B2 5/2004 Kim et al.
6,736,783 B2 5/2004 Blake et al.
6,737,158 B1 5/2004 Thompson
6,740,059 B2 5/2004 Flaherty
6,740,075 B2 5/2004 Lebel et al.
6,741,877 B1 5/2004 Shults et al.
6,742,635 B2 6/2004 Hirshberg
6,743,635 B2 6/2004 Neel et al.
6,746,582 B2 6/2004 Heller et al.
6,749,587 B2 6/2004 Flaherty
6,758,810 B2 7/2004 Lebel et al.
6,767,440 B1 7/2004 Bhullar et al.
6,767,632 B2 7/2004 Axelgaard et al.
6,770,030 B1 8/2004 Schaupp et al.
6,770,067 B2 8/2004 Lorenzen et al.
6,773,565 B2 8/2004 Kunimoto et al.
6,780,297 B2 8/2004 Matsumoto et al.
6,782,343 B2 8/2004 Hasper
6,784,274 B2 8/2004 Van Antwerp et al.
6,789,634 B1 9/2004 Denton
6,790,178 B1 9/2004 Mault et al.
6,793,632 B2 9/2004 Sohrab
6,793,802 B2 9/2004 Lee et al.
6,801,041 B2 10/2004 Karinka et al.
6,802,957 B2 10/2004 Jung et al.
6,804,002 B2 10/2004 Fine et al.
6,804,544 B2 10/2004 Van Antwerp et al.
6,804,558 B2 10/2004 Haller et al.
6,805,693 B2 10/2004 Gray et al.
6,809,507 B2 10/2004 Morgan et al.
6,809,653 B1 10/2004 Mann et al.
6,810,290 B2 10/2004 Lebel et al.
6,810,736 B2 11/2004 Ikezawa et al.
6,811,533 B2 11/2004 Lebel et al.
6,811,534 B2 11/2004 Bowman, IV et al.
6,811,548 B2 11/2004 Jeffrey
6,813,519 B2 11/2004 Lebel et al.
6,814,845 B2 11/2004 Wilson et al.
6,815,186 B2 11/2004 Clark, Jr.
6,830,551 B1 12/2004 Uchigaki et al.
6,835,553 B2 12/2004 Han et al.
6,843,899 B2 1/2005 Ufer
6,849,052 B2 2/2005 Uchigaki et al.
6,850,790 B2 2/2005 Berner et al.
6,858,020 B2 2/2005 Rusnak
6,862,465 B2 3/2005 Shults et al.
6,863,800 B2 3/2005 Karinka et al.
6,867,262 B1 3/2005 Angel et al.
6,869,413 B2 3/2005 Langley et al.
6,873,268 B2 3/2005 Lebel et al.
6,875,386 B1 4/2005 Ward et al.
6,878,112 B2 4/2005 Linberg et al.
6,881,551 B2 4/2005 Heller et al.
6,882,940 B2 4/2005 Potts et al.
6,885,883 B2 4/2005 Parris et al.
6,887,228 B2 5/2005 McKay
6,891,317 B2 5/2005 Pei et al.
6,892,085 B2 5/2005 McIvor et al.
6,893,552 B1 5/2005 Wang et al.
6,895,263 B2 5/2005 Shin et al.
6,895,265 B2 5/2005 Silver
6,902,544 B2 6/2005 Ludin et al.
6,908,681 B2 6/2005 Terry et al.
6,919,566 B1 7/2005 Cadell
6,923,763 B1 8/2005 Kovatchev et al.
6,925,393 B1 8/2005 Kalatz et al.
6,926,691 B2 8/2005 Miethke
6,931,327 B2 8/2005 Goode, Jr. et al.
6,932,584 B2 8/2005 Gray et al.
6,932,894 B2 8/2005 Mao et al.
6,936,006 B2 8/2005 Sabra
6,942,518 B2 9/2005 Liamos et al.
6,945,965 B2 9/2005 Whiting
6,948,492 B2 9/2005 Wermeling et al.
6,950,708 B2 9/2005 Bowman, IV et al.
6,952,604 B2 10/2005 Denuzzio et al.
6,954,662 B2 10/2005 Freger et al.
6,958,705 B2 10/2005 Lebel et al.
6,960,192 B1 11/2005 Flaherty et al.
6,965,791 B1 11/2005 Hitchcock et al.
6,966,325 B2 11/2005 Erickson
6,968,294 B2 11/2005 Gutta et al.
6,971,274 B2 12/2005 Olin
6,972,080 B1 12/2005 Tomioka et al.
6,973,706 B2 12/2005 Say et al.
6,974,437 B2 12/2005 Lebel et al.
6,975,893 B2 12/2005 Say et al.
6,979,315 B2 12/2005 Rogers et al.
6,983,867 B1 1/2006 Fugere
6,989,891 B2 1/2006 Braig et al.
6,990,366 B2 1/2006 Say et al.
6,991,643 B2 1/2006 Saadat
6,997,907 B2 2/2006 Safabash et al.
6,997,921 B2 2/2006 Gray et al.
6,998,247 B2 2/2006 Monfre et al.
7,003,336 B2 2/2006 Holker et al.
7,003,340 B2 2/2006 Say et al.
7,003,341 B2 2/2006 Say et al.
7,008,979 B2 3/2006 Schottman et al.
7,011,630 B2 3/2006 Desai et al.
7,016,713 B2 3/2006 Gardner et al.
7,022,072 B2 4/2006 Fox et al.
7,022,219 B2 4/2006 Mansouri et al.
7,024,245 B2 4/2006 Lebel et al.
7,025,425 B2 4/2006 Kovatchev et al.
7,025,727 B2 4/2006 Brockway et al.
7,025,743 B2 4/2006 Mann et al.
7,027,848 B2 4/2006 Robinson et al.
7,029,444 B2 4/2006 Shin et al.
7,033,322 B2 4/2006 Silver
7,039,446 B2 5/2006 Ruchti et al.
7,041,068 B2 5/2006 Freeman et al.
7,041,468 B2 5/2006 Drucker et al.
7,048,687 B1 5/2006 Reuss et al.
7,048,727 B1 5/2006 Moss
7,052,483 B2 5/2006 Wojcik
7,056,302 B2 6/2006 Douglas
7,058,437 B2 6/2006 Buse et al.
7,058,453 B2 6/2006 Nelson et al.
7,060,031 B2 6/2006 Webb et al.
7,060,059 B2 6/2006 Keith et al.
7,061,593 B2 6/2006 Braig et al.
7,063,086 B2 6/2006 Shahbazpour et al.
7,066,884 B2 6/2006 Custer et al.
7,070,577 B1 7/2006 Haller et al.
7,070,580 B2 7/2006 Nielsen
7,073,246 B2 7/2006 Bhullar et al.
7,074,307 B2 7/2006 Simpson et al.
7,078,582 B2 7/2006 Stebbings et al.
7,081,195 B2 7/2006 Simpson et al.
7,097,775 B2 8/2006 Greenberg et al.
7,098,803 B2 8/2006 Mann et al.
7,100,628 B1 9/2006 Izenson et al.
7,108,778 B2 9/2006 Simpson et al.
7,110,803 B2 9/2006 Shults et al.
7,113,821 B1 9/2006 Sun et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

7,114,502 B2 10/2006 Schulman et al.
7,115,884 B1 10/2006 Walt et al.
7,118,667 B2 10/2006 Lee
7,120,483 B2 10/2006 Russell et al.
7,125,382 B2 10/2006 Zhou et al.
7,131,967 B2 11/2006 Gray et al.
7,134,999 B2 11/2006 Brauker et al.
7,136,689 B2 11/2006 Shults et al.
7,141,034 B2 11/2006 Eppstein et al.
7,144,496 B2 12/2006 Meserol et al.
7,146,202 B2 12/2006 Ward et al.
7,150,741 B2 12/2006 Erickson et al.
7,153,265 B2 12/2006 Vachon
7,162,290 B1 1/2007 Levin
7,163,511 B2 1/2007 Conn et al.
7,166,074 B2 1/2007 Reghabi et al.
7,168,597 B1 1/2007 Jones et al.
7,169,289 B2 1/2007 Schuelein et al.
7,171,274 B2 1/2007 Starkweather et al.
7,172,075 B1 2/2007 Ji
7,183,102 B2 2/2007 Monfre et al.
7,184,810 B2 2/2007 Caduff et al.
7,190,988 B2 3/2007 Say et al.
7,192,450 B2 3/2007 Brauker et al.
7,198,606 B2 4/2007 Boecker et al.
7,207,968 B1 4/2007 Harcinske
7,207,974 B2 4/2007 Safabash et al.
7,211,074 B2 5/2007 Sansoucy
7,221,970 B2 5/2007 Parker
7,223,253 B2 5/2007 Hogendijk
7,225,535 B2 6/2007 Feldman et al.
7,226,978 B2 6/2007 Tapsak et al.
7,228,162 B2 6/2007 Ward et al.
7,229,288 B2 6/2007 Stuart et al.
7,238,165 B2 7/2007 Vincent et al.
7,241,266 B2 7/2007 Zhou et al.
7,241,586 B2 7/2007 Gulati et al.
7,247,138 B2 7/2007 Reghabi et al.
7,248,906 B2 7/2007 Dirac et al.
7,254,450 B2 8/2007 Christopherson et al.
7,255,690 B2 8/2007 Gray et al.
7,258,681 B2 8/2007 Houde
7,261,690 B2 8/2007 Teller et al.
7,266,400 B2 9/2007 Fine et al.
7,267,665 B2 9/2007 Steil et al.
7,276,029 B2 10/2007 Goode, Jr. et al.
7,276,146 B2 10/2007 Wilsey
7,276,147 B2 10/2007 Wilsey
7,278,983 B2 10/2007 Ireland et al.
7,287,318 B2 10/2007 Bhullar et al.
7,288,085 B2 10/2007 Olsen
7,291,497 B2 11/2007 Holmes et al.
7,295,867 B2 11/2007 Berner et al.
7,299,082 B2 11/2007 Feldman et al.
7,310,544 B2 12/2007 Brister et al.
7,311,690 B2 12/2007 Burnett
7,313,425 B2 12/2007 Finarov et al.
7,314,452 B2 1/2008 Madonia
7,315,767 B2 1/2008 Caduff et al.
7,316,662 B2 1/2008 Delnevo et al.
7,317,939 B2 1/2008 Fine et al.
7,318,814 B2 1/2008 Levine et al.
7,324,012 B2 1/2008 Mann et al.
7,327,273 B2 2/2008 Hung et al.
7,329,234 B2 2/2008 Sansoucy
7,329,239 B2 2/2008 Safabash et al.
7,334,594 B2 2/2008 Ludin
7,335,179 B2 2/2008 Burnett
7,335,195 B2 2/2008 Mehier
7,335,286 B2 2/2008 Abel et al.
7,335,294 B2 2/2008 Heller et al.
7,338,464 B2 3/2008 Blischak et al.
7,344,499 B1 3/2008 Prausnitz et al.
7,354,420 B2 4/2008 Steil et al.
7,359,723 B2 4/2008 Jones
7,361,155 B2 4/2008 Sage, Jr. et al.
7,364,562 B2 4/2008 Braig et al.
7,364,592 B2 4/2008 Carr-Brendel et al.
7,366,556 B2 4/2008 Brister et al.
7,366,566 B2 4/2008 Henry et al.
7,367,942 B2 5/2008 Grage et al.
7,379,765 B2 5/2008 Petisce et al.
7,381,184 B2 6/2008 Funderburk et al.
7,384,397 B2 6/2008 Zhang et al.
7,386,937 B2 6/2008 Bhullar et al.
7,392,080 B2 6/2008 Eppstein et al.
7,395,158 B2 7/2008 Monfre et al.
7,396,353 B2 7/2008 Lorenzen et al.
7,399,277 B2 7/2008 Saidara et al.
7,402,153 B2 7/2008 Steil et al.
7,404,819 B1 7/2008 Darios et al.
7,405,055 B2 7/2008 Dunn et al.
7,417,164 B2 8/2008 Suri
7,424,318 B2 9/2008 Brister et al.
7,426,408 B2 9/2008 Denuzzio et al.
7,433,727 B2 10/2008 Ward et al.
7,460,898 B2 12/2008 Brister et al.
7,467,003 B2 12/2008 Brister et al.
7,471,972 B2 12/2008 Rhodes et al.
7,494,465 B2 2/2009 Brister et al.
7,497,827 B2 3/2009 Brister et al.
7,519,408 B2 4/2009 Rasdal et al.
7,519,478 B2 4/2009 Bartkowiak et al.
7,523,004 B2 4/2009 Bartkowiak et al.
7,525,298 B2 4/2009 Morgan et al.
7,565,197 B2 7/2009 Haubrich et al.
7,574,266 B2 8/2009 Dudding et al.
7,582,059 B2 9/2009 Funderburk et al.
7,583,990 B2 9/2009 Goode, Jr. et al.
7,587,287 B2 9/2009 Connolly et al.
7,591,801 B2 9/2009 Brauker et al.
7,599,726 B2 10/2009 Goode, Jr. et al.
7,602,310 B2 10/2009 Mann et al.
7,604,593 B2 10/2009 Parris et al.
7,613,491 B2 11/2009 Boock et al.
7,615,007 B2 11/2009 Shults et al.
7,618,368 B2 11/2009 Brown
7,618,369 B2 11/2009 Hayter et al.
7,624,028 B1 11/2009 Brown
7,632,228 B2 12/2009 Brauker et al.
7,636,602 B2 12/2009 Baru Fassio et al.
7,637,868 B2 12/2009 Saint et al.
7,640,032 B2 12/2009 Jones
7,640,048 B2 12/2009 Brister et al.
7,647,237 B2 1/2010 Malave et al.
7,651,596 B2 1/2010 Petisce et al.
7,654,956 B2 2/2010 Brister et al.
7,657,297 B2 2/2010 Simpson et al.
7,687,586 B2 3/2010 Ward et al.
7,695,434 B2 4/2010 Malecha
7,697,967 B2 4/2010 Stafford
7,699,775 B2 4/2010 Desai et al.
7,711,402 B2 5/2010 Shults et al.
7,711,493 B2 5/2010 Bartkowiak et al.
7,713,574 B2 5/2010 Brister et al.
7,715,893 B2 5/2010 Kamath et al.
7,725,148 B2 5/2010 Shah et al.
7,731,659 B2 6/2010 Malecha
7,736,322 B2 6/2010 Roe et al.
7,761,126 B2 7/2010 Gardner et al.
7,766,830 B2 8/2010 Fox et al.
7,771,352 B2 8/2010 Shults et al.
7,774,145 B2 8/2010 Brauker et al.
7,792,562 B2 9/2010 Shults et al.
7,811,231 B2 10/2010 Jin et al.
7,826,981 B2 11/2010 Goode, Jr. et al.
7,857,760 B2 12/2010 Brister et al.
7,862,622 B2 1/2011 Dunlap et al.
7,874,985 B2 1/2011 Kovatchev et al.
7,879,010 B2 2/2011 Hunn et al.
7,881,763 B2 2/2011 Brauker et al.
7,885,697 B2 2/2011 Brister et al.
7,885,698 B2 2/2011 Feldman
7,890,295 B2 2/2011 Shin et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

7,896,809 B2 3/2011 Simpson et al.
7,899,511 B2 3/2011 Shults et al.
7,901,354 B2 3/2011 Shults et al.
7,905,833 B2 3/2011 Brister et al.
7,917,186 B2 3/2011 Kamath et al.
7,310,544 C1 4/2011 Brister et al.
7,920,906 B2 4/2011 Goode, Jr. et al.
7,925,321 B2 4/2011 Goode, Jr. et al.
7,927,274 B2 4/2011 Rasdal et al.
7,935,499 B2 5/2011 Dunn et al.
7,946,984 B2 5/2011 Brister et al.
7,946,985 B2 5/2011 Mastrototaro et al.
7,949,381 B2 5/2011 Brister et al.
7,640,048 C1 7/2011 Dobbles et al.
7,986,986 B2 7/2011 Goode et al.
7,998,071 B2 8/2011 Goode, Jr. et al.
8,000,901 B2 8/2011 Brauker et al.
8,005,524 B2 8/2011 Brauker et al.
8,005,525 B2 8/2011 Goode, Jr. et al.
8,010,174 B2 8/2011 Goode, Jr. et al.
8,019,421 B2 9/2011 Darvish et al.
7,497,827 C1 10/2011 Brister et al.
7,905,833 C1 11/2011 Brister et al.
8,060,173 B2 11/2011 Goode, Jr. et al.
RE43,039 E 12/2011 Brister et al.
8,073,520 B2 12/2011 Kamath et al.
8,118,877 B2 2/2012 Brauker et al.
7,774,145 C1 3/2012 Brauker et al.
8,133,178 B2 3/2012 Brauker et al.
8,152,789 B2 4/2012 Starkweather et al.
8,160,669 B2 4/2012 Brauker et al.
8,160,671 B2 4/2012 Kamath et al.
8,167,801 B2 5/2012 Goode, Jr. et al.
8,170,803 B2 5/2012 Kamath et al.
7,713,574 C1 6/2012 Brister et al.
8,216,139 B2 7/2012 Brauker et al.
8,229,534 B2 7/2012 Brister et al.
8,231,531 B2 7/2012 Brister et al.
8,233,958 B2 7/2012 Brauker et al.
8,251,906 B2 8/2012 Brauker et al.
8,257,259 B2 9/2012 Brauker et al.
8,265,725 B2 9/2012 Brauker et al.
8,275,438 B2 9/2012 Simpson et al.
8,277,713 B2 10/2012 Petisce et al.
8,280,475 B2 10/2012 Brister et al.
8,282,549 B2 10/2012 Brauker et al.
8,290,559 B2 10/2012 Shariati et al.
8,290,560 B2 10/2012 Kamath et al.
8,290,561 B2 10/2012 Brauker et al.
8,298,142 B2 10/2012 Simpson et al.
7,310,544 C2 11/2012 Brister et al.
8,311,749 B2 11/2012 Brauker et al.
8,313,434 B2 11/2012 Brister et al.
8,321,149 B2 11/2012 Brauker et al.
8,353,881 B2 1/2013 Jennewine
8,355,753 B2 1/2013 Bochenko et al.
8,364,229 B2 1/2013 Simpson et al.
8,366,614 B2 2/2013 Say et al.
8,374,667 B2 2/2013 Brauker et al.
8,452,368 B2 5/2013 Brister et al.
8,457,708 B2 6/2013 Brister et al.
8,463,350 B2 6/2013 Kamath et al.
8,469,886 B2 6/2013 Brauker et al.
8,474,397 B2 7/2013 Brister et al.
8,475,373 B2 7/2013 Brister et al.
8,483,791 B2 7/2013 Brister et al.
8,506,482 B2 8/2013 Feldman
8,515,516 B2 8/2013 Kamath et al.
8,515,519 B2 8/2013 Brister et al.
8,548,551 B2 10/2013 Kamath et al.
8,560,037 B2 10/2013 Goode, Jr. et al.
8,565,848 B2 10/2013 Brister et al.
8,565,849 B2 10/2013 Kamath et al.
8,571,625 B2 10/2013 Kamath et al.
8,579,816 B2 11/2013 Kamath et al.
8,611,978 B2 12/2013 Kamath et al.
8,615,282 B2 12/2013 Brister et al.
7,713,574 C2 3/2014 Brister et al.
8,663,109 B2 3/2014 Brister et al.
8,690,775 B2 4/2014 Brister et al.
8,731,630 B2 5/2014 Kamath et al.
8,750,955 B2 6/2014 Brister et al.
8,788,007 B2 7/2014 Brauker et al.
8,792,953 B2 7/2014 Brister et al.
8,792,954 B2 7/2014 Brister et al.
8,792,955 B2 7/2014 Brister et al.
8,825,127 B2 9/2014 Kamath et al.
8,858,434 B2 10/2014 Kamath et al.
8,886,272 B2 11/2014 Brister et al.
8,915,849 B2 12/2014 Brauker et al.
8,968,198 B2 3/2015 Brauker et al.
8,986,209 B2 3/2015 Brauker et al.
8,989,833 B2 3/2015 Brauker et al.
9,044,199 B2 6/2015 Brister et al.
9,055,901 B2 6/2015 Brister et al.
9,060,742 B2 6/2015 Brister et al.
9,078,608 B2 7/2015 Kamath et al.
9,078,626 B2 7/2015 Brister et al.
9,155,496 B2 10/2015 Shults et al.
9,220,449 B2 12/2015 Pryor et al.
9,247,900 B2 2/2016 Brister et al.
9,314,196 B2 4/2016 Pryor et al.
9,414,777 B2 8/2016 Brister et al.
9,603,557 B2 3/2017 Brister et al.
9,610,031 B2 4/2017 Brister et al.
9,668,677 B2 6/2017 Brister et al.
9,669,156 B2 6/2017 Jennewine
9,724,028 B2 8/2017 Brauker et al.
9,775,543 B2 10/2017 Brister et al.
9,801,572 B2 10/2017 Brister et al.
9,814,414 B2 11/2017 Brister et al.
9,833,176 B2 12/2017 Brister et al.
9,918,668 B2 3/2018 Pryor et al.
10,022,078 B2 7/2018 Brauker et al.
10,314,525 B2 6/2019 Simpson et al.
2001/0008187 A1 7/2001 Hanssen et al.
2001/0016682 A1 8/2001 Berner et al.
2001/0020546 A1 9/2001 Eldridge et al.
2001/0021817 A1 9/2001 Brugger et al.
2001/0027327 A1 10/2001 Schraga
2001/0039053 A1 11/2001 Liseo et al.
2001/0039387 A1 11/2001 Rutynowski et al.
2001/0041830 A1 11/2001 Varalli et al.
2001/0044413 A1 11/2001 Pierce et al.
2001/0044588 A1 11/2001 Mault
2001/0051766 A1 12/2001 Gazdzinski
2001/0051768 A1 12/2001 Schulman et al.
2001/0053933 A1 12/2001 Phaneuf et al.
2002/0009810 A1 1/2002 O'Connor et al.
2002/0016535 A1 2/2002 Martin et al.
2002/0019022 A1 2/2002 Dunn et al.
2002/0019330 A1 2/2002 Murray et al.
2002/0022883 A1 2/2002 Burg
2002/0023852 A1 2/2002 McIvor et al.
2002/0026110 A1 2/2002 Parris et al.
2002/0026111 A1 2/2002 Ackerman
2002/0032531 A1 3/2002 Mansky et al.
2002/0038081 A1 3/2002 Fein et al.
2002/0042090 A1 4/2002 Heller et al.
2002/0042561 A1 4/2002 Schulman et al.
2002/0043471 A1 4/2002 Ikeda et al.
2002/0043651 A1 4/2002 Darrow et al.
2002/0045808 A1 4/2002 Ford et al.
2002/0050250 A1 5/2002 Peterson et al.
2002/0055673 A1 5/2002 Van Antwerp et al.
2002/0065453 A1 5/2002 Lesho et al.
2002/0068860 A1 6/2002 Clark, Jr.
2002/0071776 A1 6/2002 Bandis et al.
2002/0077599 A1 6/2002 Wojcik
2002/0084196 A1 7/2002 Liamos et al.
2002/0099282 A1 7/2002 Knobbe et al.
2002/0099997 A1 7/2002 Piret
2002/0100474 A1 8/2002 Kellner et al.
2002/0100725 A1 8/2002 Lee et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2002/0103499 A1 8/2002 Perez et al.
2002/0106709 A1 8/2002 Potts et al.
2002/0111547 A1 8/2002 Knobbe et al.
2002/0119711 A1 8/2002 Van Antwerp et al.
2002/0123048 A1 9/2002 Gau
2002/0128419 A1 9/2002 Terry et al.
2002/0128546 A1 9/2002 Silver
2002/0128594 A1 9/2002 Das et al.
2002/0132279 A1 9/2002 Hockersmith
2002/0133063 A1 9/2002 Hockersmith et al.
2002/0133224 A1 9/2002 Bajgar et al.
2002/0137991 A1 9/2002 Scarantino et al.
2002/0151796 A1 10/2002 Koulik
2002/0151816 A1 10/2002 Rich et al.
2002/0155615 A1 10/2002 Novikov et al.
2002/0160722 A1 10/2002 Terranova et al.
2002/0161288 A1 10/2002 Shin et al.
2002/0164836 A1 11/2002 Ho
2002/0169369 A1 11/2002 Ward et al.
2002/0169405 A1 11/2002 Roberts
2002/0177763 A1 11/2002 Burns et al.
2002/0177764 A1 11/2002 Sohrab
2002/0182241 A1 12/2002 Borenstein et al.
2002/0185384 A1 12/2002 Leong et al.
2002/0186185 A1 12/2002 Ide et al.
2002/0188185 A1 12/2002 Sohrab
2002/0188216 A1 12/2002 Kayyali et al.
2002/0188252 A1 12/2002 Bardy
2002/0193679 A1 12/2002 Malave et al.
2002/0193885 A1 12/2002 Legeay et al.
2002/0198513 A1 12/2002 Lebel et al.
2003/0003524 A1 1/2003 Taniike et al.
2003/0004432 A1 1/2003 Assenheimer
2003/0004457 A1 1/2003 Andersson
2003/0006669 A1 1/2003 Pei et al.
2003/0009093 A1 1/2003 Silver
2003/0021729 A1 1/2003 Moller et al.
2003/0023171 A1 1/2003 Sato et al.
2003/0023317 A1 1/2003 Brauker et al.
2003/0023461 A1 1/2003 Quintanilla et al.
2003/0024811 A1 2/2003 Davies et al.
2003/0028089 A1 2/2003 Galley et al.
2003/0032867 A1 2/2003 Crothall et al.
2003/0032874 A1 2/2003 Rhodes et al.
2003/0036773 A1 2/2003 Whitehurst et al.
2003/0042137 A1 3/2003 Mao et al.
2003/0049166 A1 3/2003 Pendo et al.
2003/0050537 A1 3/2003 Wessel
2003/0050546 A1 3/2003 Desai et al.
2003/0054428 A1 3/2003 Monfre et al.
2003/0055464 A1 3/2003 Darvish et al.
2003/0059631 A1 3/2003 Al-Lamee
2003/0060765 A1 3/2003 Campbell et al.
2003/0065254 A1 4/2003 Schulman et al.
2003/0065308 A1 4/2003 Lebel et al.
2003/0070548 A1 4/2003 Clausen
2003/0072741 A1 4/2003 Berglund et al.
2003/0076082 A1 4/2003 Morgan et al.
2003/0078481 A1 4/2003 McIvor et al.
2003/0078560 A1 4/2003 Miller et al.
2003/0088166 A1 5/2003 Say et al.
2003/0091433 A1 5/2003 Tam et al.
2003/0096424 A1 5/2003 Mao et al.
2003/0097082 A1 5/2003 Purdy et al.
2003/0099682 A1 5/2003 Moussy et al.
2003/0100040 A1 5/2003 Bonnecaze et al.
2003/0100821 A1 5/2003 Heller et al.
2003/0104119 A1 6/2003 Wilson et al.
2003/0114735 A1 6/2003 Silver et al.
2003/0117296 A1 6/2003 Seely
2003/0119208 A1 6/2003 Yoon et al.
2003/0120152 A1 6/2003 Omiya
2003/0125612 A1 7/2003 Fox et al.
2003/0125613 A1 7/2003 Enegren et al.
2003/0130616 A1 7/2003 Steil et al.
2003/0132227 A1 7/2003 Geisler et al.
2003/0134100 A1 7/2003 Mao et al.
2003/0134347 A1 7/2003 Heller et al.
2003/0138674 A1 7/2003 Zeikus et al.
2003/0143746 A1 7/2003 Sage, Jr.
2003/0146110 A1 8/2003 Karinka et al.
2003/0153821 A1 8/2003 Berner et al.
2003/0157409 A1 8/2003 Huang
2003/0158520 A1 8/2003 Safabash et al.
2003/0168338 A1 9/2003 Gao et al.
2003/0176183 A1 9/2003 Drucker et al.
2003/0176933 A1 9/2003 Lebel et al.
2003/0181794 A1 9/2003 Rini et al.
2003/0186457 A1 10/2003 Iwaki et al.
2003/0187338 A1 10/2003 Say et al.
2003/0188427 A1 10/2003 Say et al.
2003/0190383 A1 10/2003 Kim
2003/0191377 A1 10/2003 Robinson et al.
2003/0199744 A1 10/2003 Buse et al.
2003/0199745 A1 10/2003 Burson et al.
2003/0199790 A1 10/2003 Boecker et al.
2003/0199878 A1 10/2003 Pohjonen et al.
2003/0200040 A1 10/2003 Trygg et al.
2003/0203498 A1 10/2003 Neel et al.
2003/0203991 A1 10/2003 Schottman et al.
2003/0208113 A1 11/2003 Mault et al.
2003/0211625 A1 11/2003 Cohan et al.
2003/0212317 A1 11/2003 Kovatchev et al.
2003/0212346 A1 11/2003 Yuzhakov et al.
2003/0212347 A1 11/2003 Sohrab
2003/0212379 A1 11/2003 Bylund et al.
2003/0217966 A1 11/2003 Tapsak et al.
2003/0225324 A1 12/2003 Anderson et al.
2003/0225360 A1 12/2003 Eppstein et al.
2003/0225361 A1 12/2003 Sabra
2003/0225437 A1 12/2003 Ferguson
2003/0228681 A1 12/2003 Ritts et al.
2003/0231550 A1 12/2003 MacFarlane
2003/0235817 A1 12/2003 Bartkowiak et al.
2004/0002682 A1 1/2004 Kovelman et al.
2004/0006263 A1 1/2004 Anderson et al.
2004/0008761 A1 1/2004 Kelliher et al.
2004/0010207 A1 1/2004 Flaherty et al.
2004/0011671 A1 1/2004 Shults et al.
2004/0015063 A1 1/2004 Denuzzio et al.
2004/0015134 A1 1/2004 Lavi et al.
2004/0018486 A1 1/2004 Dunn et al.
2004/0023253 A1 2/2004 Kunwar et al.
2004/0023317 A1 2/2004 Motamedi et al.
2004/0024327 A1 2/2004 Brodnick
2004/0024553 A1 2/2004 Monfre et al.
2004/0030285 A1 2/2004 Lavi et al.
2004/0030294 A1 2/2004 Mahurkar
2004/0039298 A1 2/2004 Abreu
2004/0039342 A1 2/2004 Eppstein et al.
2004/0039343 A1 2/2004 Eppstein et al.
2004/0039406 A1 2/2004 Jessen
2004/0040840 A1 3/2004 Mao et al.
2004/0045879 A1 3/2004 Shults et al.
2004/0052689 A1 3/2004 Yao
2004/0054263 A1 3/2004 Moerman et al.
2004/0054352 A1 3/2004 Adams et al.
2004/0064068 A1 4/2004 Denuzzio et al.
2004/0064156 A1 4/2004 Shah et al.
2004/0068230 A1 4/2004 Estes et al.
2004/0074785 A1 4/2004 Holker et al.
2004/0078219 A1 4/2004 Kaylor et al.
2004/0087671 A1 5/2004 Tamada et al.
2004/0088023 A1 5/2004 Imran et al.
2004/0106741 A1 6/2004 Kriesel et al.
2004/0106857 A1 6/2004 Gough
2004/0106858 A1 6/2004 Say et al.
2004/0106859 A1 6/2004 Say et al.
2004/0106860 A1 6/2004 Say et al.
2004/0111017 A1 6/2004 Say et al.
2004/0120848 A1 6/2004 Teodorczyk
2004/0122353 A1 6/2004 Shahmirian et al.
2004/0127777 A1 7/2004 Ruchti et al.
2004/0127818 A1 7/2004 Roe et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2004/0133131 A1 7/2004 Kuhn et al.
2004/0133164 A1 7/2004 Funderburk et al.
2004/0135684 A1 7/2004 Steinthal et al.
2004/0138543 A1 7/2004 Russell et al.
2004/0138588 A1 7/2004 Saikley et al.
2004/0143173 A1 7/2004 Reghabi et al.
2004/0146909 A1 7/2004 Duong et al.
2004/0147872 A1 7/2004 Thompson
2004/0152187 A1 8/2004 Haight et al.
2004/0152622 A1 8/2004 Keith et al.
2004/0158138 A1 8/2004 Kilcoyne et al.
2004/0158207 A1 8/2004 Hunn et al.
2004/0167382 A1 8/2004 Gardner et al.
2004/0167801 A1 8/2004 Say et al.
2004/0171921 A1 9/2004 Say et al.
2004/0173472 A1 9/2004 Jung et al.
2004/0176672 A1 9/2004 Silver et al.
2004/0180391 A1 9/2004 Gratzl et al.
2004/0186362 A1 9/2004 Brauker et al.
2004/0186365 A1 9/2004 Jin et al.
2004/0193025 A1 9/2004 Steil et al.
2004/0193090 A1 9/2004 Lebel et al.
2004/0197846 A1 10/2004 Hockersmith et al.
2004/0199059 A1 10/2004 Brauker et al.
2004/0204687 A1 10/2004 Mogensen et al.
2004/0204744 A1 10/2004 Penner et al.
2004/0206625 A1 10/2004 Bhullar et al.
2004/0206916 A1 10/2004 Colvin et al.
2004/0219664 A1 11/2004 Heller et al.
2004/0220517 A1 11/2004 Starkweather et al.
2004/0225199 A1 11/2004 Evanyk et al.
2004/0225338 A1 11/2004 Lebel et al.
2004/0228902 A1 11/2004 Benz
2004/0234575 A1 11/2004 Horres et al.
2004/0236200 A1 11/2004 Say et al.
2004/0236251 A1 11/2004 Roe et al.
2004/0242982 A1 12/2004 Sakata et al.
2004/0244151 A1 12/2004 Sakata et al.
2004/0249421 A1 12/2004 Harel et al.
2004/0253365 A1 12/2004 Warren et al.
2004/0254433 A1 12/2004 Bandis et al.
2004/0254434 A1 12/2004 Goodnow et al.
2004/0260164 A1 12/2004 Kilcoyne et al.
2004/0260234 A1 12/2004 Srinivasan et al.
2004/0265940 A1 12/2004 Slater et al.
2004/0267300 A1 12/2004 Mace
2005/0000829 A1 1/2005 Morita et al.
2005/0003399 A1 1/2005 Blackburn et al.
2005/0003470 A1 1/2005 Nelson et al.
2005/0004438 A1 1/2005 Ward et al.
2005/0004439 A1 1/2005 Shin et al.
2005/0004494 A1 1/2005 Perez et al.
2005/0006122 A1 1/2005 Burnette
2005/0008671 A1 1/2005 Van Antwerp
2005/0010265 A1 1/2005 Baru Fassio et al.
2005/0010269 A1 1/2005 Lebel et al.
2005/0013842 A1 1/2005 Qiu et al.
2005/0016325 A1 1/2005 Enokido
2005/0026689 A1 2/2005 Marks
2005/0027177 A1 2/2005 Shin et al.
2005/0027180 A1 2/2005 Goode et al.
2005/0027181 A1 2/2005 Goode et al.
2005/0027182 A1 2/2005 Siddiqui et al.
2005/0027462 A1 2/2005 Goode et al.
2005/0027463 A1 2/2005 Goode, Jr. et al.
2005/0031689 A1 2/2005 Shults et al.
2005/0033132 A1 2/2005 Shults et al.
2005/0038332 A1 2/2005 Saidara et al.
2005/0043598 A1 2/2005 Goode, Jr. et al.
2005/0049472 A1 3/2005 Manda et al.
2005/0051427 A1 3/2005 Brauker et al.
2005/0051440 A1 3/2005 Simpson et al.
2005/0054909 A1 3/2005 Petisce et al.
2005/0056551 A1 3/2005 White et al.
2005/0056552 A1 3/2005 Simpson et al.
2005/0059871 A1 3/2005 Gough et al.
2005/0065464 A1 3/2005 Talbot et al.
2005/0070770 A1 3/2005 Dirac et al.
2005/0077584 A1 4/2005 Uhland et al.
2005/0079200 A1 4/2005 Rathenow et al.
2005/0080345 A1 4/2005 Finburgh et al.
2005/0085839 A1 4/2005 Allen et al.
2005/0090607 A1 4/2005 Tapsak et al.
2005/0096512 A1 5/2005 Fox et al.
2005/0096519 A1 5/2005 DeNuzzio et al.
2005/0101847 A1 5/2005 Routt et al.
2005/0103624 A1 5/2005 Bhullar et al.
2005/0103625 A1 5/2005 Rhodes et al.
2005/0106713 A1 5/2005 Phan et al.
2005/0107677 A1 5/2005 Ward et al.
2005/0112169 A1 5/2005 Brauker et al.
2005/0113653 A1 5/2005 Fox et al.
2005/0114068 A1 5/2005 Chey et al.
2005/0115832 A1 6/2005 Simpson et al.
2005/0118344 A1 6/2005 Pacetti
2005/0119720 A1 6/2005 Gale et al.
2005/0121322 A1 6/2005 Say et al.
2005/0124873 A1 6/2005 Shults et al.
2005/0131305 A1 6/2005 Danielson et al.
2005/0131346 A1 6/2005 Douglas
2005/0133368 A1 6/2005 Davies et al.
2005/0139489 A1 6/2005 Davies et al.
2005/0143635 A1 6/2005 Kamath et al.
2005/0143636 A1 6/2005 Zhang et al.
2005/0143675 A1 6/2005 Neel et al.
2005/0154264 A1 7/2005 Lecompte et al.
2005/0154271 A1 7/2005 Rasdal et al.
2005/0154272 A1 7/2005 Dirac et al.
2005/0161346 A1 7/2005 Simpson et al.
2005/0173245 A1 8/2005 Feldman et al.
2005/0176136 A1 8/2005 Burd et al.
2005/0176678 A1 8/2005 Horres et al.
2005/0177036 A1 8/2005 Shults et al.
2005/0177398 A1 8/2005 Watanabe et al.
2005/0181012 A1 8/2005 Saint et al.
2005/0182306 A1 8/2005 Sloan
2005/0182451 A1 8/2005 Griffin et al.
2005/0183954 A1 8/2005 Hitchcock et al.
2005/0187720 A1 8/2005 Goode, Jr. et al.
2005/0192557 A1 9/2005 Brauker et al.
2005/0195930 A1 9/2005 Spital et al.
2005/0196747 A1 9/2005 Stiene
2005/0197554 A1 9/2005 Polcha
2005/0199494 A1 9/2005 Say et al.
2005/0203360 A1 9/2005 Brauker et al.
2005/0203364 A1 9/2005 Monfre et al.
2005/0209665 A1 9/2005 Hunter et al.
2005/0211571 A1 9/2005 Schulein et al.
2005/0215871 A1 9/2005 Feldman et al.
2005/0215872 A1 9/2005 Berner et al.
2005/0215979 A1 9/2005 Kornerup et al.
2005/0225361 A1 10/2005 Rhee
2005/0228238 A1 10/2005 Monitzer
2005/0233407 A1 10/2005 Pamidi et al.
2005/0236361 A1 10/2005 Ufer et al.
2005/0239154 A1 10/2005 Feldman et al.
2005/0239156 A1 10/2005 Drucker et al.
2005/0241957 A1 11/2005 Mao et al.
2005/0242479 A1 11/2005 Petisce et al.
2005/0245795 A1 11/2005 Goode, Jr. et al.
2005/0245799 A1 11/2005 Brauker et al.
2005/0251083 A1 11/2005 Carr-Brendel et al.
2005/0258037 A1 11/2005 Hajizadeh et al.
2005/0261563 A1 11/2005 Zhou et al.
2005/0267325 A1 12/2005 Bouchier et al.
2005/0267440 A1 12/2005 Herman et al.
2005/0271546 A1 12/2005 Gerber et al.
2005/0272989 A1 12/2005 Shah et al.
2005/0277164 A1 12/2005 Drucker et al.
2005/0282997 A1 12/2005 Ward et al.
2005/0284758 A1 12/2005 Funke et al.
2005/0287620 A1 12/2005 Heller et al.
2005/0288596 A1 12/2005 Eigler et al.
2006/0001538 A1 1/2006 Kraft et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2006/0001550 A1 1/2006 Mann et al.
2006/0003398 A1 1/2006 Heller et al.
2006/0007017 A1 1/2006 Mann et al.
2006/0008370 A1 1/2006 Massaro et al.
2006/0010098 A1 1/2006 Goodnow et al.
2006/0015020 A1 1/2006 Neale et al.
2006/0015024 A1 1/2006 Brister et al.
2006/0016700 A1 1/2006 Brister et al.
2006/0019327 A1 1/2006 Brister et al.
2006/0020186 A1 1/2006 Brister et al.
2006/0020187 A1 1/2006 Brister et al.
2006/0020188 A1 1/2006 Kamath et al.
2006/0020189 A1 1/2006 Brister et al.
2006/0020190 A1 1/2006 Kamath et al.
2006/0020191 A1 1/2006 Brister et al.
2006/0020192 A1 1/2006 Brister et al.
2006/0025663 A1 2/2006 Talbot et al.
2006/0036139 A1 2/2006 Brister et al.
2006/0036140 A1 2/2006 Brister et al.
2006/0036141 A1 2/2006 Kamath et al.
2006/0036142 A1 2/2006 Brister et al.
2006/0036143 A1 2/2006 Brister et al.
2006/0036144 A1 2/2006 Brister et al.
2006/0036145 A1 2/2006 Brister et al.
2006/0040402 A1 2/2006 Brauker et al.
2006/0042080 A1 3/2006 Say et al.
2006/0047215 A1 3/2006 Newman et al.
2006/0049359 A1 3/2006 Busta et al.
2006/0052745 A1 3/2006 Van Antwerp et al.
2006/0065527 A1 3/2006 Samproni
2006/0068208 A1 3/2006 Tapsak et al.
2006/0074564 A1 4/2006 Bartkowiak et al.
2006/0079740 A1 4/2006 Silver et al.
2006/0079809 A1 4/2006 Goldberger et al.
2006/0086624 A1 4/2006 Tapsak et al.
2006/0094946 A1 5/2006 Kellogg et al.
2006/0100588 A1 5/2006 Brunnberg et al.
2006/0118415 A1 6/2006 Say et al.
2006/0142651 A1 6/2006 Brister et al.
2006/0155180 A1 7/2006 Brister et al.
2006/0159718 A1 7/2006 Rathenow et al.
2006/0159981 A1 7/2006 Heller
2006/0166629 A1 7/2006 Reggiardo
2006/0169599 A1 8/2006 Feldman et al.
2006/0171980 A1 8/2006 Helmus et al.
2006/0173444 A1 8/2006 Choy et al.
2006/0183178 A1 8/2006 Gulati et al.
2006/0183871 A1 8/2006 Ward et al.
2006/0183984 A1 8/2006 Dobbles et al.
2006/0183985 A1 8/2006 Brister et al.
2006/0189856 A1 8/2006 Petisce et al.
2006/0189863 A1 8/2006 Peyser et al.
2006/0195029 A1 8/2006 Shults et al.
2006/0198864 A1 9/2006 Shults et al.
2006/0200019 A1 9/2006 Petisce et al.
2006/0200020 A1 9/2006 Brister et al.
2006/0200022 A1 9/2006 Brauker et al.
2006/0200970 A1 9/2006 Brister et al.
2006/0200981 A1 9/2006 Bhullar et al.
2006/0200982 A1 9/2006 Bhullar et al.
2006/0204536 A1 9/2006 Shults et al.
2006/0211921 A1 9/2006 Brauker et al.
2006/0222566 A1 10/2006 Brauker et al.
2006/0224108 A1 10/2006 Brauker et al.
2006/0224109 A1 10/2006 Steil et al.
2006/0224141 A1 10/2006 Rush et al.
2006/0226985 A1 10/2006 Goodnow et al.
2006/0229512 A1 10/2006 Petisce et al.
2006/0235285 A1 10/2006 Brister et al.
2006/0247508 A1 11/2006 Fennell
2006/0247710 A1 11/2006 Goetz et al.
2006/0253085 A1 11/2006 Geismar et al.
2006/0253086 A1 11/2006 Moberg et al.
2006/0257995 A1 11/2006 Simpson et al.
2006/0257996 A1 11/2006 Simpson et al.
2006/0258761 A1 11/2006 Boock et al.
2006/0258929 A1 11/2006 Goode, Jr. et al.
2006/0263673 A1 11/2006 Kim et al.
2006/0263763 A1 11/2006 Simpson et al.
2006/0263839 A1 11/2006 Ward et al.
2006/0264888 A1 11/2006 Moberg et al.
2006/0270922 A1 11/2006 Brauker et al.
2006/0270923 A1 11/2006 Brauker et al.
2006/0275859 A1 12/2006 Kjaer
2006/0281985 A1 12/2006 Ward et al.
2006/0287691 A1 12/2006 Drew
2006/0289307 A1 12/2006 Yu et al.
2006/0293576 A1 12/2006 Van Antwerp et al.
2007/0003588 A1 1/2007 Chinn et al.
2007/0007133 A1 1/2007 Mang et al.
2007/0016381 A1 1/2007 Kamath et al.
2007/0017805 A1 1/2007 Hodges et al.
2007/0027370 A1 2/2007 Brauker et al.
2007/0027381 A1 2/2007 Stafford
2007/0027384 A1 2/2007 Brister et al.
2007/0027385 A1 2/2007 Brister et al.
2007/0032706 A1 2/2007 Kamath et al.
2007/0032717 A1 2/2007 Brister et al.
2007/0032718 A1 2/2007 Shults et al.
2007/0038044 A1 2/2007 Dobbles et al.
2007/0045902 A1 3/2007 Brauker et al.
2007/0049865 A1 3/2007 Radmer et al.
2007/0049873 A1 3/2007 Hansen et al.
2007/0059196 A1 3/2007 Brister et al.
2007/0060814 A1 3/2007 Stafford
2007/0066873 A1 3/2007 Kamath et al.
2007/0073129 A1 3/2007 Shah et al.
2007/0078320 A1 4/2007 Stafford
2007/0078321 A1 4/2007 Mazza et al.
2007/0078322 A1 4/2007 Stafford
2007/0085995 A1 4/2007 Pesach et al.
2007/0093704 A1 4/2007 Brister et al.
2007/0106135 A1 5/2007 Sloan et al.
2007/0111196 A1 5/2007 Alarcon et al.
2007/0116600 A1 5/2007 Kochar et al.
2007/0129524 A1 6/2007 Sunkara
2007/0129619 A1 6/2007 Ward et al.
2007/0129621 A1 6/2007 Kellogg et al.
2007/0135698 A1 6/2007 Shah et al.
2007/0135699 A1 6/2007 Ward et al.
2007/0142584 A1 6/2007 Schorzman et al.
2007/0149875 A1 6/2007 Ouyang et al.
2007/0151869 A1 7/2007 Heller et al.
2007/0156033 A1 7/2007 Causey, III et al.
2007/0161070 A1 7/2007 Wilsey
2007/0161880 A1 7/2007 Say et al.
2007/0163880 A1 7/2007 Woo et al.
2007/0173706 A1 7/2007 Neinast et al.
2007/0173709 A1 7/2007 Petisce et al.
2007/0173710 A1 7/2007 Petisce et al.
2007/0173711 A1 7/2007 Shah et al.
2007/0179436 A1 8/2007 Braig et al.
2007/0191701 A1 8/2007 Feldman et al.
2007/0197889 A1 8/2007 Brister et al.
2007/0197890 A1 8/2007 Boock et al.
2007/0200254 A1 8/2007 Curry
2007/0200267 A1 8/2007 Tsai
2007/0202562 A1 8/2007 Curry
2007/0202672 A1 8/2007 Curry
2007/0203407 A1 8/2007 Hoss et al.
2007/0203410 A1 8/2007 Say et al.
2007/0203966 A1 8/2007 Brauker et al.
2007/0206193 A1 9/2007 Pesach
2007/0208244 A1 9/2007 Brauker et al.
2007/0208245 A1 9/2007 Brauker et al.
2007/0208246 A1 9/2007 Brauker et al.
2007/0213610 A1 9/2007 Say et al.
2007/0213611 A1 9/2007 Simpson et al.
2007/0215491 A1 9/2007 Heller et al.
2007/0218097 A1 9/2007 Heller et al.
2007/0219441 A1 9/2007 Carlin et al.
2007/0219496 A1 9/2007 Kamen et al.
2007/0222609 A1 9/2007 Duron et al.
2007/0225579 A1 9/2007 Lucassen et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2007/0225675 A1 9/2007 Robinson et al.
2007/0227907 A1 10/2007 Shah et al.
2007/0232876 A1 10/2007 Otto et al.
2007/0232879 A1 10/2007 Brister et al.
2007/0233013 A1 10/2007 Schoenberg
2007/0235331 A1 10/2007 Simpson et al.
2007/0240497 A1 10/2007 Robinson et al.
2007/0244381 A1 10/2007 Robinson et al.
2007/0244382 A1 10/2007 Robinson et al.
2007/0244383 A1 10/2007 Talbot et al.
2007/0249916 A1 10/2007 Pesach et al.
2007/0249922 A1 10/2007 Peyser et al.
2007/0255114 A1 11/2007 Ackermann et al.
2007/0255126 A1 11/2007 Moberg et al.
2007/0255531 A1 11/2007 Drew
2007/0258395 A1 11/2007 Jollota et al.
2007/0259217 A1 11/2007 Logan
2007/0275193 A1 11/2007 Desimone et al.
2007/0276211 A1 11/2007 Mir et al.
2007/0282180 A1 12/2007 Caduff et al.
2007/0299385 A1 12/2007 Santini, Jr. et al.
2007/0299409 A1 12/2007 Whitbourne et al.
2008/0001318 A1 1/2008 Schorzman et al.
2008/0009692 A1 1/2008 Stafford
2008/0017522 A1 1/2008 Heller et al.
2008/0021666 A1 1/2008 Goode, Jr. et al.
2008/0027245 A1 1/2008 Suri
2008/0029390 A1 2/2008 Roche et al.
2008/0029391 A1 2/2008 Mao et al.
2008/0030369 A1 2/2008 Mann et al.
2008/0033254 A1 2/2008 Kamath et al.
2008/0033268 A1 2/2008 Stafford
2008/0033269 A1 2/2008 Zhang
2008/0034972 A1 2/2008 Gough et al.
2008/0039702 A1 2/2008 Hayter et al.
2008/0045824 A1 2/2008 Tapsak et al.
2008/0064941 A1 3/2008 Funderburk et al.
2008/0064943 A1 3/2008 Talbot et al.
2008/0064944 A1 3/2008 Van Antwerp et al.
2008/0070231 A1 3/2008 Franciskovich et al.
2008/0071156 A1 3/2008 Brister et al.
2008/0071157 A1 3/2008 McGarraugh et al.
2008/0071158 A1 3/2008 McGarraugh et al.
2008/0071328 A1 3/2008 Haubrich et al.
2008/0072663 A1 3/2008 Keenan et al.
2008/0081969 A1 4/2008 Feldman et al.
2008/0083617 A1 4/2008 Simpson et al.
2008/0086040 A1 4/2008 Heller et al.
2008/0086041 A1 4/2008 Heller et al.
2008/0086042 A1 4/2008 Brister et al.
2008/0086043 A1 4/2008 Heller et al.
2008/0086044 A1 4/2008 Brister et al.
2008/0086273 A1 4/2008 Shults et al.
2008/0091094 A1 4/2008 Heller et al.
2008/0091095 A1 4/2008 Heller et al.
2008/0108942 A1 5/2008 Brister et al.
2008/0119703 A1 5/2008 Brister et al.
2008/0119704 A1 5/2008 Brister et al.
2008/0119705 A1 5/2008 Patel et al.
2008/0119706 A1 5/2008 Brister et al.
2008/0119707 A1 5/2008 Stafford
2008/0125751 A1 5/2008 Fjield et al.
2008/0139910 A1 6/2008 Mastrototaro et al.
2008/0154101 A1 6/2008 Jain et al.
2008/0161666 A1 7/2008 Feldman et al.
2008/0183061 A1 7/2008 Goode et al.
2008/0183399 A1 7/2008 Goode et al.
2008/0187655 A1 8/2008 Markle et al.
2008/0188722 A1 8/2008 Markle et al.
2008/0188725 A1 8/2008 Markle et al.
2008/0188731 A1 8/2008 Brister et al.
2008/0189051 A1 8/2008 Goode et al.
2008/0193936 A1 8/2008 Squirrell
2008/0194837 A1 8/2008 Kim et al.
2008/0194935 A1 8/2008 Brister et al.
2008/0194936 A1 8/2008 Goode et al.
2008/0194937 A1 8/2008 Goode et al.
2008/0194938 A1 8/2008 Brister et al.
2008/0195232 A1 8/2008 Carr-Brendel et al.
2008/0195967 A1 8/2008 Goode et al.
2008/0197024 A1 8/2008 Simpson et al.
2008/0200788 A1 8/2008 Brister et al.
2008/0200789 A1 8/2008 Brister et al.
2008/0200791 A1 8/2008 Simpson et al.
2008/0208025 A1 8/2008 Shults et al.
2008/0210557 A1 9/2008 Heller et al.
2008/0214915 A1 9/2008 Brister et al.
2008/0214918 A1 9/2008 Brister et al.
2008/0228051 A1 9/2008 Shults et al.
2008/0228054 A1 9/2008 Shults et al.
2008/0235469 A1 9/2008 Drew
2008/0242961 A1 10/2008 Brister et al.
2008/0255438 A1 10/2008 Saidara et al.
2008/0255440 A1 10/2008 Eilersen et al.
2008/0262334 A1 10/2008 Dunn et al.
2008/0262469 A1 10/2008 Brister et al.
2008/0275313 A1 11/2008 Brister et al.
2008/0275326 A1 11/2008 Kasielke et al.
2008/0287764 A1 11/2008 Rasdal et al.
2008/0287765 A1 11/2008 Rasdal et al.
2008/0287766 A1 11/2008 Rasdal et al.
2008/0296155 A1 12/2008 Shults et al.
2008/0305009 A1 12/2008 Gamsey et al.
2008/0305506 A1 12/2008 Suri
2008/0306368 A1 12/2008 Goode, Jr. et al.
2008/0306433 A1 12/2008 Cesaroni
2008/0306434 A1 12/2008 Dobbles et al.
2008/0306435 A1 12/2008 Kamath et al.
2008/0306444 A1 12/2008 Brister et al.
2008/0312397 A1 12/2008 Lai et al.
2008/0312518 A1 12/2008 Jina et al.
2008/0312859 A1 12/2008 Skyggebjerg et al.
2008/0313896 A1 12/2008 Shah et al.
2009/0005666 A1 1/2009 Shin et al.
2009/0012379 A1 1/2009 Goode, Jr. et al.
2009/0018418 A1 1/2009 Markle et al.
2009/0018424 A1 1/2009 Kamath et al.
2009/0018425 A1 1/2009 Ouyang et al.
2009/0018426 A1 1/2009 Markle et al.
2009/0020502 A1 1/2009 Bhullar et al.
2009/0030294 A1 1/2009 Petisce et al.
2009/0036758 A1 2/2009 Brauker et al.
2009/0036763 A1 2/2009 Brauker et al.
2009/0043181 A1 2/2009 Brauker et al.
2009/0043182 A1 2/2009 Brauker et al.
2009/0043525 A1 2/2009 Brauker et al.
2009/0043541 A1 2/2009 Brauker et al.
2009/0043542 A1 2/2009 Brauker et al.
2009/0045055 A1 2/2009 Rhodes et al.
2009/0054748 A1 2/2009 Feldman
2009/0054812 A1 2/2009 Mace
2009/0061528 A1 3/2009 Suri
2009/0062633 A1 3/2009 Brauker et al.
2009/0062635 A1 3/2009 Brauker et al.
2009/0062645 A1 3/2009 Fehre et al.
2009/0076356 A1 3/2009 Simpson et al.
2009/0076360 A1 3/2009 Brister et al.
2009/0076361 A1 3/2009 Kamath et al.
2009/0081803 A1 3/2009 Gamsey et al.
2009/0085768 A1 4/2009 Patel et al.
2009/0099434 A1 4/2009 Liu et al.
2009/0099436 A1 4/2009 Brister et al.
2009/0112478 A1 4/2009 Mueller, Jr. et al.
2009/0124877 A1 5/2009 Goode, Jr. et al.
2009/0124878 A1 5/2009 Goode, Jr. et al.
2009/0124879 A1 5/2009 Brister et al.
2009/0124964 A1 5/2009 Leach et al.
2009/0131768 A1 5/2009 Simpson et al.
2009/0131769 A1 5/2009 Leach et al.
2009/0131776 A1 5/2009 Simpson et al.
2009/0131777 A1 5/2009 Simpson et al.
2009/0137886 A1 5/2009 Shariati et al.
2009/0137887 A1 5/2009 Shariati et al.
2009/0143659 A1 6/2009 Li et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2009/0143660 A1 6/2009 Brister et al.
2009/0150186 A1 6/2009 Cohen et al.
2009/0156919 A1 6/2009 Brister et al.
2009/0156924 A1 6/2009 Shariati et al.
2009/0163790 A1 6/2009 Brister et al.
2009/0163791 A1 6/2009 Brister et al.
2009/0177059 A1 7/2009 Say et al.
2009/0177060 A1 7/2009 Say et al.
2009/0177143 A1 7/2009 Markle et al.
2009/0178459 A1 7/2009 Li et al.
2009/0182217 A1 7/2009 Li et al.
2009/0187090 A1 7/2009 Say et al.
2009/0187091 A1 7/2009 Say et al.
2009/0192366 A1 7/2009 Mensinger et al.
2009/0192380 A1 7/2009 Shariati et al.
2009/0192722 A1 7/2009 Shariati et al.
2009/0192724 A1 7/2009 Brauker et al.
2009/0192745 A1 7/2009 Kamath et al.
2009/0192751 A1 7/2009 Kamath et al.
2009/0203981 A1 8/2009 Brauker et al.
2009/0204340 A1 8/2009 Feldman et al.
2009/0204341 A1 8/2009 Brauker et al.
2009/0216103 A1 8/2009 Brister et al.
2009/0240120 A1 9/2009 Mensinger et al.
2009/0240128 A1 9/2009 Mensinger et al.
2009/0240193 A1 9/2009 Mensinger et al.
2009/0242399 A1 10/2009 Kamath et al.
2009/0242425 A1 10/2009 Kamath et al.
2009/0247855 A1 10/2009 Boock et al.
2009/0247856 A1 10/2009 Boock et al.
2009/0247857 A1 10/2009 Harper et al.
2009/0264719 A1 10/2009 Markle et al.
2009/0264856 A1 10/2009 Lebel et al.
2009/0287073 A1 11/2009 Boock et al.
2009/0287074 A1 11/2009 Shults et al.
2009/0299155 A1 12/2009 Yang et al.
2009/0299156 A1 12/2009 Simpson et al.
2009/0299162 A1 12/2009 Brauker et al.
2009/0299276 A1 12/2009 Brauker et al.
2009/0310743 A1 12/2009 Carpenter et al.
2010/0010324 A1 1/2010 Brauker et al.
2010/0010330 A1 1/2010 Rankers et al.
2010/0010331 A1 1/2010 Brauker et al.
2010/0010332 A1 1/2010 Brauker et al.
2010/0016687 A1 1/2010 Brauker et al.
2010/0016698 A1 1/2010 Rasdal et al.
2010/0022855 A1 1/2010 Brauker et al.
2010/0030038 A1 2/2010 Brauker et al.
2010/0030053 A1 2/2010 Goode, Jr. et al.
2010/0030484 A1 2/2010 Brauker et al.
2010/0030485 A1 2/2010 Brauker et al.
2010/0036215 A1 2/2010 Goode, Jr. et al.
2010/0036216 A1 2/2010 Goode, Jr. et al.
2010/0036222 A1 2/2010 Goode, Jr. et al.
2010/0036223 A1 2/2010 Goode, Jr. et al.
2010/0036224 A1 2/2010 Goode, Jr. et al.
2010/0036225 A1 2/2010 Goode, Jr. et al.
2010/0041971 A1 2/2010 Goode, Jr. et al.
2010/0045465 A1 2/2010 Brauker et al.
2010/0049024 A1 2/2010 Saint et al.
2010/0063373 A1 3/2010 Kamath et al.
2010/0069728 A1 3/2010 Funderburk et al.
2010/0076283 A1 3/2010 Simpson et al.
2010/0081908 A1 4/2010 Dobbles et al.
2010/0081910 A1 4/2010 Brister et al.
2010/0087721 A1 4/2010 Stafford
2010/0087724 A1 4/2010 Brauker et al.
2010/0096259 A1 4/2010 Zhang et al.
2010/0099970 A1 4/2010 Shults et al.
2010/0099971 A1 4/2010 Shults et al.
2010/0113897 A1 5/2010 Brenneman et al.
2010/0119693 A1 5/2010 Tapsak et al.
2010/0121169 A1 5/2010 Petisce et al.
2010/0161269 A1 6/2010 Kamath et al.
2010/0168543 A1 7/2010 Kamath et al.
2010/0168545 A1 7/2010 Kamath et al.
2010/0168546 A1 7/2010 Kamath et al.
2010/0174157 A1 7/2010 Brister et al.
2010/0174158 A1 7/2010 Kamath et al.
2010/0174163 A1 7/2010 Brister et al.
2010/0174164 A1 7/2010 Brister et al.
2010/0174165 A1 7/2010 Brister et al.
2010/0174166 A1 7/2010 Brister et al.
2010/0174167 A1 7/2010 Kamath et al.
2010/0174168 A1 7/2010 Goode, Jr. et al.
2010/0179399 A1 7/2010 Goode, Jr. et al.
2010/0179400 A1 7/2010 Brauker et al.
2010/0179401 A1 7/2010 Rasdal et al.
2010/0179402 A1 7/2010 Goode, Jr. et al.
2010/0179404 A1 7/2010 Kamath et al.
2010/0179405 A1 7/2010 Goode, Jr. et al.
2010/0179406 A1 7/2010 Goode, Jr. et al.
2010/0179407 A1 7/2010 Goode, Jr. et al.
2010/0179408 A1 7/2010 Kamath et al.
2010/0179409 A1 7/2010 Kamath et al.
2010/0185065 A1 7/2010 Goode, Jr. et al.
2010/0185069 A1 7/2010 Brister et al.
2010/0185070 A1 7/2010 Brister et al.
2010/0185071 A1 7/2010 Simpson et al.
2010/0185072 A1 7/2010 Goode, Jr. et al.
2010/0185073 A1 7/2010 Goode, Jr. et al.
2010/0185074 A1 7/2010 Goode, Jr. et al.
2010/0185075 A1 7/2010 Brister et al.
2010/0191082 A1 7/2010 Brister et al.
2010/0198035 A1 8/2010 Kamath et al.
2010/0204555 A1 8/2010 Shults et al.
2010/0212583 A1 8/2010 Brister et al.
2010/0214104 A1 8/2010 Goode, Jr. et al.
2010/0217106 A1 8/2010 Goode, Jr. et al.
2010/0217555 A1 8/2010 Kamath et al.
2010/0234707 A1 9/2010 Goode, Jr. et al.
2010/0234796 A1 9/2010 Kamath et al.
2010/0235106 A1 9/2010 Kamath et al.
2010/0240975 A1 9/2010 Goode, Jr. et al.
2010/0240976 A1 9/2010 Goode, Jr. et al.
2010/0256779 A1 10/2010 Brauker et al.
2010/0286496 A1 11/2010 Simpson et al.
2010/0305869 A1 12/2010 Brauker et al.
2010/0331648 A1 12/2010 Kamath et al.
2010/0331655 A1 12/2010 Kamath et al.
2010/0331656 A1 12/2010 Mensinger et al.
2010/0331657 A1 12/2010 Mensinger et al.
2011/0009727 A1 1/2011 Mensinger et al.
2011/0027127 A1 2/2011 Simpson et al.
2011/0028815 A1 2/2011 Simpson et al.
2011/0028816 A1 2/2011 Simpson et al.
2011/0046467 A1 2/2011 Simpson et al.
2011/0087196 A1 4/2011 Hunn et al.
2011/0118579 A1 5/2011 Goode, Jr. et al.
2011/0124992 A1 5/2011 Brauker et al.
2011/0124997 A1 5/2011 Goode, Jr. et al.
2011/0128052 A1 6/2011 Fujibe et al.
2011/0130639 A1 6/2011 Feldman
2011/0130970 A1 6/2011 Goode, Jr. et al.
2011/0136249 A1 6/2011 Stiene
2011/0137601 A1 6/2011 Goode, Jr. et al.
2011/0144465 A1 6/2011 Shults et al.
2011/0178378 A1 7/2011 Brister et al.
2011/0190614 A1 8/2011 Brister et al.
2011/0201910 A1 8/2011 Rasdal et al.
2011/0218414 A1 9/2011 Kamath et al.
2011/0231107 A1 9/2011 Brauker et al.
2011/0231140 A1 9/2011 Goode, Jr. et al.
2011/0231141 A1 9/2011 Goode, Jr. et al.
2011/0231142 A1 9/2011 Goode, Jr. et al.
2011/0253533 A1 10/2011 Shults et al.
2011/0257895 A1 10/2011 Brauker et al.
2011/0263958 A1 10/2011 Brauker et al.
2011/0270158 A1 11/2011 Brauker et al.
2011/0290645 A1 12/2011 Brister et al.
2011/0313543 A1 12/2011 Brauker et al.
2011/0319739 A1 12/2011 Kamath et al.
2012/0130214 A1 5/2012 Brister et al.
2012/0172691 A1 7/2012 Brauker et al.

(56) **References Cited**

U.S. PATENT DOCUMENTS

2012/0265042	A1	10/2012	Neinast et al.
2012/0277562	A1	11/2012	Brister et al.
2012/0277566	A1	11/2012	Kamath et al.
2013/0012798	A1	1/2013	Brister et al.
2013/0060112	A1	3/2013	Pryor et al.
2013/0131478	A1	5/2013	Simpson et al.
2013/0255570	A1	10/2013	Brister et al.
2013/0267808	A1	10/2013	Brister et al.
2013/0281931	A1	10/2013	Hunn et al.
2013/0296677	A1	11/2013	Pryor et al.
2014/0128704	A1	5/2014	Simpson et al.
2014/0142405	A1	5/2014	Brister et al.
2014/0257065	A1	9/2014	Brister et al.
2014/0288402	A1	9/2014	Brister et al.
2016/0008029	A1	1/2016	Brister et al.
2016/0051173	A1	2/2016	Brister et al.
2016/0310051	A1	10/2016	Brister et al.
2017/0196491	A1	7/2017	Brister et al.
2017/0367627	A1	12/2017	Brister et al.
2018/0014762	A1	1/2018	Brister et al.
2018/0049682	A1	2/2018	Brister et al.
2018/0055423	A1	3/2018	Pryor et al.
2018/0140236	A1	5/2018	Brister et al.
2018/0160949	A1	6/2018	Brister et al.
2018/0242894	A1	8/2018	Brauker et al.
2018/0303394	A1	10/2018	Brauker et al.
2019/0015020	A1	1/2019	Brister et al.
2019/0069817	A1	3/2019	Brister et al.
2019/0076071	A1	3/2019	Brister et al.
2019/0320951	A1	10/2019	Pryor et al.
2019/0320952	A1	10/2019	Pryor et al.
2019/0320953	A1	10/2019	Pryor et al.
2019/0320954	A1	10/2019	Pryor et al.
2019/0320955	A1	10/2019	Pryor et al.
2019/0320956	A1	10/2019	Pryor et al.
2019/0320957	A1	10/2019	Pryor et al.
2019/0343436	A1	11/2019	Pryor
2019/0350503	A1	11/2019	Pryor
2019/0350504	A1	11/2019	Pryor
2019/0357821	A1	11/2019	Brister et al.
2019/0357822	A1	11/2019	Pryor

FOREIGN PATENT DOCUMENTS

DE	2658734	A1	6/1978
DE	3144459	A1	10/1982
DE	4105222	A1	8/1992
DE	3933373	C2	9/1992
DE	4401400	A1	7/1995
DE	20110059	U1	8/2002
DE	10117285	A1	11/2002
EP	0098592	A2	1/1984
EP	0107634	A2	5/1984
EP	0127958	A2	12/1984
EP	0143517	A1	6/1985
EP	0282349	A2	9/1988
EP	0284518	A2	9/1988
EP	0286039	A2	10/1988
EP	0286118	A2	10/1988
EP	0288793	A2	11/1988
EP	0143517	B1	4/1989
EP	0314027	A1	5/1989
EP	0319277	A1	6/1989
EP	0320109	A1	6/1989
EP	0351892	A2	1/1990
EP	0352138	A2	1/1990
EP	0352610	A2	1/1990
EP	0352631	A2	1/1990
EP	0352708	A2	1/1990
EP	0353328	A1	2/1990
EP	0390390	A1	10/1990
EP	0396788	A1	11/1990
EP	0406473	A1	1/1991
EP	0420021	A2	4/1991
EP	0440044	A1	8/1991
EP	0441252	A2	8/1991
EP	0441394	A2	8/1991
EP	0457292	A2	11/1991
EP	0467078	A2	1/1992
EP	0471391	A2	2/1992
EP	0473065	A2	3/1992
EP	0476980	A2	3/1992
EP	0275139	B1	4/1992
EP	0477501	A2	4/1992
EP	0494704	A3	9/1992
EP	0494705	A3	9/1992
EP	0508388	A1	10/1992
EP	0512122	A1	11/1992
EP	0520430	A2	12/1992
EP	0520443	A2	12/1992
EP	0262328	B1	1/1993
EP	0319277	B1	3/1993
EP	0534074	A1	3/1993
EP	0535898	A1	4/1993
EP	0539625	A1	5/1993
EP	0264036	B1	6/1993
EP	0351891	B1	9/1993
EP	0563795	A1	10/1993
EP	0567725	A1	11/1993
EP	0323605	B1	1/1994
EP	0279069	B1	7/1994
EP	0595474	A3	7/1994
EP	0561966	B1	10/1994
EP	0286118	B1	1/1995
EP	0478550	B1	1/1995
EP	0647849	A2	4/1995
EP	0424634	B1	6/1995
EP	0677743	A2	10/1995
EP	0191640	B2	11/1995
EP	0424633	B1	1/1996
EP	0690134	A1	1/1996
EP	0534074	B1	3/1996
EP	0709677	A2	5/1996
EP	0532187	B1	10/1996
EP	0470652	B1	12/1996
EP	0476715	B1	12/1996
EP	0747069	A2	12/1996
EP	0776628	A2	6/1997
EP	0387696	B1	8/1997
EP	0593096	A3	12/1997
EP	0817809	A1	1/1998
EP	0838230	A2	4/1998
EP	0880936	A2	12/1998
EP	0885932	A2	12/1998
EP	0587008	B1	2/1999
EP	0967788	A2	12/1999
EP	0995805	A1	4/2000
EP	0678308	B1	5/2000
EP	1048264	A1	11/2000
EP	1077634	A1	2/2001
EP	1078258	A1	2/2001
EP	1028320	A3	3/2001
EP	1111378	A2	6/2001
EP	1112717	A1	7/2001
EP	1112718	A1	7/2001
EP	1120084	A1	8/2001
EP	1120085	A1	8/2001
EP	1120650	A2	8/2001
EP	1130386	A1	9/2001
EP	1153571	A1	11/2001
EP	0777122	B1	4/2002
EP	0817809	B1	7/2002
EP	1251137	A2	10/2002
EP	0958495	B1	11/2002
EP	1258728	A2	11/2002
EP	1266607	A2	12/2002
EP	1281351	A2	2/2003
EP	0824900	B1	4/2003
EP	1340980	A1	9/2003
EP	1340981	A2	9/2003
EP	1077636	B1	1/2004
EP	0674176	B1	2/2004
EP	1391728	A1	2/2004
EP	0846776	B1	3/2004

(56) **References Cited**

FOREIGN PATENT DOCUMENTS

EP	1413245	A2	4/2004
EP	1430831	A1	6/2004
EP	1498067	A1	1/2005
EP	1498428	A1	1/2005
EP	1614464	A1	1/2006
EP	0922959	B1	10/2006
EP	1717924	A2	11/2006
EP	0877252	B1	1/2007
EP	1234053	B1	4/2007
EP	1798542	A1	6/2007
EP	1286164	B1	7/2007
EP	1804650	A1	7/2007
EP	1612560	B1	10/2007
EP	1905514	A1	4/2008
EP	1582874	B1	7/2008
EP	1977829	A1	10/2008
EP	1982644	A1	10/2008
EP	1457913	B1	12/2008
EP	2223710	A1	9/2010
EP	2226086	A1	9/2010
EP	2228642	A1	9/2010
EP	2236077	A1	10/2010
EP	2327362	A1	6/2011
EP	2329770	B1	9/2014
EP	2407094	B1	10/2014
EP	2335584	B1	6/2015
FR	2656423	A1	6/1991
FR	2760962	A1	9/1998
GB	1442303	A	7/1976
GB	1556969	A	12/1979
GB	2149918	A	6/1985
GB	2230865	A	10/1990
JP	S5441190	A	4/1979
JP	S57156004	A	9/1982
JP	S58124912	A	7/1983
JP	S59211459	A	11/1984
JP	S61271418	A	12/1986
JP	S6283649	A	4/1987
JP	S6283849	A	4/1987
JP	H022913	A	1/1990
JP	H03293556	A	12/1991
JP	H06288853	A	10/1994
JP	H06307898	A	11/1994
JP	H0783871	A	3/1995
JP	H11258381	A	9/1999
JP	2000060826	A	2/2000
JP	2000149072	A	5/2000
JP	2002513602	A	5/2002
JP	2002189015	A	7/2002
JP	2003108679	A	4/2003
JP	2003297163	A	10/2003
WO	WO-8103614	A1	12/1981
WO	WO-8706342	A1	10/1987
WO	WO-8706706	A1	11/1987
WO	WO-8808137	A1	10/1988
WO	WO-8902720	A1	4/1989
WO	WO-8904302	A1	5/1989
WO	WO-8907263	A1	8/1989
WO	WO-9000738	A1	1/1990
WO	WO-9002938	A1	3/1990
WO	WO-9005296	A1	5/1990
WO	WO-9005301	A1	5/1990
WO	WO-9005302	A1	5/1990
WO	WO-9005910	A1	5/1990
WO	WO-9007525	A1	7/1990
WO	WO-9007575	A1	7/1990
WO	WO-9010716	A1	9/1990
WO	WO-9010861	A1	9/1990
WO	WO-9013021	A1	11/1990
WO	WO-9109302	A1	6/1991
WO	WO-9115993	A1	10/1991
WO	WO-9116416	A1	10/1991
WO	WO-9117259	A1	11/1991
WO	WO-9201315	A1	1/1992
WO	WO-9201928	A1	2/1992
WO	WO-9207525	A1	5/1992
WO	WO-9208985	A2	5/1992
WO	WO-9210584	A1	6/1992
WO	WO-9212255	A1	7/1992
WO	WO-9213271	A1	8/1992
WO	WO-9214138	A1	8/1992
WO	WO-9214139	A1	8/1992
WO	WO-9218887	A1	10/1992
WO	WO-9221772	A1	12/1992
WO	WO-9301308	A1	1/1993
WO	WO-9302703	A1	2/1993
WO	WO-9303362	A1	2/1993
WO	WO-9305701	A1	4/1993
WO	WO-9312256	A1	6/1993
WO	WO-9313048	A1	7/1993
WO	WO-9313408	A1	7/1993
WO	WO-9314185	A1	7/1993
WO	WO-9314693	A1	8/1993
WO	WO-9319370	A1	9/1993
WO	WO-9319701	A1	10/1993
WO	WO-9320240	A1	10/1993
WO	WO-9320440	A1	10/1993
WO	WO-9320441	A1	10/1993
WO	WO-9320443	A1	10/1993
WO	WO-9320444	A1	10/1993
WO	WO-9320450	A1	10/1993
WO	WO-9323744	A1	11/1993
WO	WO-9406011	A1	3/1994
WO	WO-9406012	A1	3/1994
WO	WO-9408236	A1	4/1994
WO	WO-9409506	A1	4/1994
WO	WO-9409507	A1	4/1994
WO	WO-9419695	A1	9/1994
WO	WO-9421642	A1	9/1994
WO	WO-9421643	A1	9/1994
WO	WO-9421644	A1	9/1994
WO	WO-9422367	A1	10/1994
WO	WO-9426414	A1	11/1994
WO	WO-9502357	A1	1/1995
WO	WO-9507109	A1	3/1995
WO	WO-9508774	A2	3/1995
WO	WO-9510044	A1	4/1995
WO	WO-9511454	A1	4/1995
WO	WO-9513838	A1	5/1995
WO	WO-9517966	A1	7/1995
WO	WO-9522597	A1	8/1995
WO	WO-9528878	A1	11/1995
WO	WO-9534814	A1	12/1995
WO	WO-9601611	A1	1/1996
WO	WO-9603117	A1	2/1996
WO	WO-9605501	A1	2/1996
WO	WO-9606947	A1	3/1996
WO	WO-9614026	A1	5/1996
WO	WO-9622991	A1	8/1996
WO	WO-9622992	A1	8/1996
WO	WO-9624690	A1	8/1996
WO	WO-9625088	A1	8/1996
WO	WO-9625089	A1	8/1996
WO	WO-9630431	A1	10/1996
WO	WO-9632076	A1	10/1996
WO	WO-9635370	A1	11/1996
WO	WO-9636296	A1	11/1996
WO	WO-9636870	A1	11/1996
WO	WO-9641179	A1	12/1996
WO	WO-9701986	A1	1/1997
WO	WO-9702811	A1	1/1997
WO	WO-9706727	A1	2/1997
WO	WO-9711080	A1	3/1997
WO	WO-9713874	A1	4/1997
WO	WO-9717884	A2	5/1997
WO	WO-9719188	A1	5/1997
WO	WO-9719344	A1	5/1997
WO	WO-9728737	A1	8/1997
WO	WO-9730628	A1	8/1997
WO	WO-9733176	A1	9/1997
WO	WO-9738625	A1	10/1997
WO	WO-9743633	A1	11/1997
WO	WO-9801071	A1	1/1998
WO	WO-9804902	A1	2/1998

(56) **References Cited**

FOREIGN PATENT DOCUMENTS

WO	WO-9806423 A1	2/1998
WO	WO-9819159 A1	5/1998
WO	WO-9824358 A2	6/1998
WO	WO-9824366 A2	6/1998
WO	WO-9833549 A1	8/1998
WO	WO-9834541 A1	8/1998
WO	WO-9835053 A2	8/1998
WO	WO-9838904 A1	9/1998
WO	WO-9838906 A1	9/1998
WO	WO-9841854 A1	9/1998
WO	WO-9842249 A1	10/1998
WO	WO-9844347 A1	10/1998
WO	WO-9844348 A1	10/1998
WO	WO-9845331 A2	10/1998
WO	WO-9845427 A2	10/1998
WO	WO-9852043 A1	11/1998
WO	WO-9856293 A1	12/1998
WO	WO-9856923 A1	12/1998
WO	WO-9858250 A2	12/1998
WO	WO-9904043 A1	1/1999
WO	WO-9912607 A1	3/1999
WO	WO-9913101 A1	3/1999
WO	WO-9913574 A1	3/1999
WO	WO-9927848 A1	6/1999
WO	WO-9927852 A1	6/1999
WO	WO-9929230 A1	6/1999
WO	WO-9929429 A1	6/1999
WO	WO-9929892 A1	6/1999
WO	WO-9933504 A1	7/1999
WO	WO-9940848 A1	8/1999
WO	WO-9948419 A1	9/1999
WO	WO-9949856 A2	10/1999
WO	WO-9956613 A1	11/1999
WO	WO-9958051 A1	11/1999
WO	WO-9958709 A1	11/1999
WO	WO-9958973 A1	11/1999
WO	WO-9959657 A1	11/1999
WO	WO-9964620 A2	12/1999
WO	WO-0007013 A2	2/2000
WO	WO-0012720 A2	3/2000
WO	WO-0013002 A2	3/2000
WO	WO-0013003 A1	3/2000
WO	WO-0018449 A2	4/2000
WO	WO-0019887 A1	4/2000
WO	WO-0020626 A1	4/2000
WO	WO-0030530 A1	6/2000
WO	WO-0032098 A1	6/2000
WO	WO-0033065 A1	6/2000
WO	WO-0035530 A1	6/2000
WO	WO-0045696 A1	8/2000
WO	WO-0049940 A2	8/2000
WO	WO-0049941 A1	8/2000
WO	WO-0049942 A2	8/2000
WO	WO-0059370 A1	10/2000
WO	WO-0059373 A1	10/2000
WO	WO-0074753 A1	12/2000
WO	WO-0078210 A1	12/2000
WO	WO-0078992 A2	12/2000
WO	WO-0079258 A1	12/2000
WO	WO-0100865 A2	1/2001
WO	WO-0109096 A2	2/2001
WO	WO-0112158 A1	2/2001
WO	WO-0116579 A1	3/2001
WO	WO-0120019 A2	3/2001
WO	WO-0120334 A1	3/2001
WO	WO-0121827 A1	3/2001
WO	WO-0134243 A1	5/2001
WO	WO-0136666 A1	5/2001
WO	WO-0143660 A2	6/2001
WO	WO-0152727 A1	7/2001
WO	WO-0152935 A1	7/2001
WO	WO-0154753 A2	8/2001
WO	WO-0158347 A1	8/2001
WO	WO-0158348 A2	8/2001
WO	WO-0159425 A1	8/2001
WO	WO-0164105 A1	9/2001
WO	WO-0168901 A2	9/2001
WO	WO-0169222 A2	9/2001
WO	WO-0173109 A2	10/2001
WO	WO-0188524 A1	11/2001
WO	WO-0188534 A2	11/2001
WO	WO-0191634 A2	12/2001
WO	WO-0202755 A2	1/2002
WO	WO-0205702 A2	1/2002
WO	WO-0207596 A1	1/2002
WO	WO-0207617 A2	1/2002
WO	WO-0216535 A2	2/2002
WO	WO-0216905 A2	2/2002
WO	WO-0217780 A1	3/2002
WO	WO-0224065 A1	3/2002
WO	WO-0233407 A1	4/2002
WO	WO-0243585 A1	6/2002
WO	WO-02056751 A2	7/2002
WO	WO-02058537 A2	8/2002
WO	WO-02062210 A1	8/2002
WO	WO-02066509 A2	8/2002
WO	WO-02066986 A2	8/2002
WO	WO-02074161 A2	9/2002
WO	WO-02082989 A1	10/2002
WO	WO-02087681 A2	11/2002
WO	WO-02089666 A2	11/2002
WO	WO-02097414 A2	12/2002
WO	WO-02099097 A1	12/2002
WO	WO-02099428 A1	12/2002
WO	WO-02100266 A1	12/2002
WO	WO-02100457 A2	12/2002
WO	WO-02100474 A2	12/2002
WO	WO-03000127 A2	1/2003
WO	WO-03008013 A2	1/2003
WO	WO-03008014 A2	1/2003
WO	WO-03009207 A1	1/2003
WO	WO-03009208 A1	1/2003
WO	WO-03011131 A2	2/2003
WO	WO-03012422 A1	2/2003
WO	WO-03022327 A2	3/2003
WO	WO-03028797 A1	4/2003
WO	WO-03032411 A2	4/2003
WO	WO-03033726 A1	4/2003
WO	WO-03035117 A1	5/2003
WO	WO-03036310 A1	5/2003
WO	WO-03044511 A2	5/2003
WO	WO-03053498 A2	7/2003
WO	WO-03057028 A2	7/2003
WO	WO-03063700 A1	8/2003
WO	WO-03072164 A2	9/2003
WO	WO-03072269 A1	9/2003
WO	WO-03076893 A2	9/2003
WO	WO-03076937 A2	9/2003
WO	WO-03082091 A2	10/2003
WO	WO-03085372 A2	10/2003
WO	WO-03088832 A1	10/2003
WO	WO-03094714 A1	11/2003
WO	WO-03097866 A1	11/2003
WO	WO-03101862 A1	12/2003
WO	WO-03106031 A1	12/2003
WO	WO-03106966 A2	12/2003
WO	WO-2004004905 A2	1/2004
WO	WO-2004010844 A2	2/2004
WO	WO-2004030726 A1	4/2004
WO	WO-2004036183 A2	4/2004
WO	WO-2004039265 A2	5/2004
WO	WO-2004052190 A1	6/2004
WO	WO-2004060455 A1	7/2004
WO	WO-2004061420 A2	7/2004
WO	WO-2004063718 A2	7/2004
WO	WO-2004071291 A2	8/2004
WO	WO-2004073138 A1	8/2004
WO	WO-2004086970 A1	10/2004
WO	WO-2004098685 A1	11/2004
WO	WO-2004105641 A2	12/2004
WO	WO-2004110256 A2	12/2004
WO	WO-2004113901 A1	12/2004
WO	WO-2005011489 A1	2/2005
WO	WO-2005012873 A2	2/2005

(56) **References Cited**

FOREIGN PATENT DOCUMENTS

WO	WO-2005013824	A1	2/2005
WO	WO-2005018443	A1	3/2005
WO	WO-2005026178	A2	3/2005
WO	WO-2005026689	A2	3/2005
WO	WO-2005026690	A2	3/2005
WO	WO-2005032362	A2	4/2005
WO	WO-2005032400	A2	4/2005
WO	WO-2005041766	A1	5/2005
WO	WO-2005048834	A1	6/2005
WO	WO-2005051440	A1	6/2005
WO	WO-2005057168	A2	6/2005
WO	WO-2005057173	A2	6/2005
WO	WO-2005057175	A2	6/2005
WO	WO-2005063115	A1	7/2005
WO	WO-2005065542	A2	7/2005
WO	WO-2005067797	A1	7/2005
WO	WO-2005070287	A1	8/2005
WO	WO-2005074811	A1	8/2005
WO	WO-2005078424	A1	8/2005
WO	WO-2005084530	A2	9/2005
WO	WO-2005084545	A1	9/2005
WO	WO-2005084546	A2	9/2005
WO	WO-2005089103	A2	9/2005
WO	WO-2005026689	A9	10/2005
WO	WO-2005094714	A1	10/2005
WO	WO-2005098431	A1	10/2005
WO	WO-2005107594	A2	11/2005
WO	WO-2005114218	A2	12/2005
WO	WO-2005119238	A1	12/2005
WO	WO-2005121355	A1	12/2005
WO	WO-2005121785	A2	12/2005
WO	WO-2005122296	A2	12/2005
WO	WO-2006001929	A1	1/2006
WO	WO-2006001973	A2	1/2006
WO	WO-2006002960	A1	1/2006
WO	WO-2006005503	A1	1/2006
WO	WO-2006010533	A2	2/2006
WO	WO-2006015922	A1	2/2006
WO	WO-2006017358	A1	2/2006
WO	WO-2006018425	A2	2/2006
WO	WO-2006019665	A1	2/2006
WO	WO-2006021430	A2	3/2006
WO	WO-2006024671	A1	3/2006
WO	WO-2006029293	A1	3/2006
WO	WO-2006050405	A1	5/2006
WO	WO-2006050843	A1	5/2006
WO	WO-2006060806	A2	6/2006
WO	WO-2006071770	A2	7/2006
WO	WO-2006072089	A1	7/2006
WO	WO-2006076412	A1	7/2006
WO	WO-2006079114	A2	7/2006
WO	WO-2006088576	A2	8/2006
WO	WO-2006098887	A2	9/2006
WO	WO-2006099151	A2	9/2006
WO	WO-2006102359	A2	9/2006
WO	WO-2006102412	A2	9/2006
WO	WO-2006104843	A1	10/2006
WO	WO-2006105146	A2	10/2006
WO	WO-2006108811	A1	10/2006
WO	WO-2006118713	A1	11/2006
WO	WO-2006118947	A2	11/2006
WO	WO-2006122048	A1	11/2006
WO	WO-2006122553	A1	11/2006
WO	WO-2006124759	A2	11/2006
WO	WO-2006130268	A1	12/2006
WO	WO-2006131288	A1	12/2006
WO	WO-2006132884	A2	12/2006
WO	WO-2006133171	A2	12/2006
WO	WO-2007002209	A2	1/2007
WO	WO-2007002579	A2	1/2007
WO	WO-2007005170	A2	1/2007
WO	WO-2007006454	A1	1/2007
WO	WO-2007009911	A1	1/2007
WO	WO-2007011587	A2	1/2007
WO	WO-2007016399	A2	2/2007
WO	WO-2007021892	A1	2/2007
WO	WO-2007021894	A2	2/2007
WO	WO-2007025088	A2	3/2007
WO	WO-2007027788	A2	3/2007
WO	WO-2007028138	A2	3/2007
WO	WO-2007028271	A2	3/2007
WO	WO-2007033010	A1	3/2007
WO	WO-2007037970	A1	4/2007
WO	WO-2007037989	A2	4/2007
WO	WO-2007040559	A2	4/2007
WO	WO-2007041069	A2	4/2007
WO	WO-2007041070	A2	4/2007
WO	WO-2007041072	A2	4/2007
WO	WO-2007041248	A2	4/2007
WO	WO-2007053832	A2	5/2007
WO	WO-2007056638	A2	5/2007
WO	WO-2007058921	A2	5/2007
WO	WO-2007059476	A2	5/2007
WO	WO-2007059478	A1	5/2007
WO	WO-2007061992	A2	5/2007
WO	WO-2007065285	A2	6/2007
WO	WO-2007070486	A2	6/2007
WO	WO-2007076303	A1	7/2007
WO	WO-2007079015	A2	7/2007
WO	WO-2007079025	A2	7/2007
WO	WO-2007081811	A2	7/2007
WO	WO-2007090037	A2	8/2007
WO	WO-2007097754	A1	8/2007
WO	WO-2007101223	A2	9/2007
WO	WO-2007101260	A2	9/2007
WO	WO-2007109372	A2	9/2007
WO	WO-2007111885	A2	10/2007
WO	WO-2007112006	A2	10/2007
WO	WO-2007114943	A2	10/2007
WO	WO-2007115094	A2	10/2007
WO	WO-2007120363	A2	10/2007
WO	WO-2007120552	A2	10/2007
WO	WO-2007126444	A2	11/2007
WO	WO-2007127606	A1	11/2007
WO	WO-2007127622	A2	11/2007
WO	WO-2007127879	A2	11/2007
WO	WO-2007127880	A2	11/2007
WO	WO-2007130239	A1	11/2007
WO	WO-2007137286	A2	11/2007
WO	WO-2007053832	A3	12/2007
WO	WO-2007143225	A2	12/2007
WO	WO-2008001091	A1	1/2008
WO	WO-2008003003	A2	1/2008
WO	WO-2008005780	A2	1/2008
WO	WO-2008013849	A2	1/2008
WO	WO-2008016486	A2	2/2008
WO	WO-2008021913	A2	2/2008
WO	WO-2008022021	A2	2/2008
WO	WO-2008028644	A1	3/2008
WO	WO-2008031106	A2	3/2008
WO	WO-2008031110	A2	3/2008
WO	WO-2008037485	A1	4/2008
WO	WO-2008039944	A2	4/2008
WO	WO-2008039946	A2	4/2008
WO	WO-2008039949	A2	4/2008
WO	WO-2008042760	A2	4/2008
WO	WO-2008048709	A1	4/2008
WO	WO-2008051407	A2	5/2008
WO	WO-2008051924	A2	5/2008
WO	WO-2008052199	A2	5/2008
WO	WO-2008055037	A2	5/2008
WO	WO-2008055128	A1	5/2008
WO	WO-2008055199	A1	5/2008
WO	WO-2008067314	A2	6/2008
WO	WO-2008069931	A1	6/2008
WO	WO-2008069932	A1	6/2008
WO	WO-2008073813	A1	6/2008
WO	WO-2008076868	A2	6/2008
WO	WO-2008079616	A1	7/2008
WO	WO-2008080591	A1	7/2008
WO	WO-2008083379	A1	7/2008
WO	WO-2008088490	A1	7/2008
WO	WO-2008094249	A1	8/2008
WO	WO-2008101211	A2	8/2008

(56) **References Cited**

FOREIGN PATENT DOCUMENTS

WO	WO-2008101217	A2	8/2008
WO	WO-2008103620	A2	8/2008
WO	WO-2006023241	A9	10/2008
WO	WO-2008116329	A1	10/2008
WO	WO-2008118257	A1	10/2008
WO	WO-2008119470	A1	10/2008
WO	WO-2008124597	A1	10/2008
WO	WO-2008134441	A1	11/2008
WO	WO-2008135453	A1	11/2008
WO	WO-2008137405	A1	11/2008
WO	WO-2008138006	A2	11/2008
WO	WO-2008150280	A1	12/2008
WO	WO-2008150917	A1	12/2008
WO	WO-2008150946	A1	12/2008
WO	WO-2008150949	A1	12/2008
WO	WO-2009006139	A2	1/2009
WO	WO-2009105709	A1	8/2009

OTHER PUBLICATIONS

Aalders, et al., "Development of a Wearable Glucose Sensor; Studies in Healthy Volunteers and in Diabetic Patients," The International Journal of Artificial Organs, 1991, vol. 14, No. 2, pp. 102-108.

Abe, et al., "Characterization of Glucose Microsensors for Intracellular Measurements," Analytical Chemistry, 1992, vol. 64, No. 18, pp. 2160-2163.

Abel, et al., "Biosensors for in Vivo Glucose Measurements: Can We Cross the Experimental Stage," Biosensors & Bioelectronics, 2002, vol. 17, pp. 1059-1070.

Abel, et al., "Experience With an Implantable Glucose Sensor as a Prerequisite of an Artificial Beta Cell," Biomed. Biochim. Actan, 1984, vol. 43, No. 5, pp. 577-584.

Adilman, et al., "Videogames: Knowing the Score, Creative Computing," Dec, 1983, Dialog: File 148; IAC Trade & Industry Database, vol. 9, p. 224(5) (9 pages).

Alberts B., et al., "Molecular Biology of the Cell," 3rd edition, 1994, p. G19 (3 pages).

Alcock S.J., et al., "Continuous Analyte Monitoring to Aid Clinical Practice," IEEE Engineering in Medicine & Biology, 1994, vol. 13, pp. 319-325.

Amato, et al., "Experience with the Polytetrafluoroethylene Surgical Membrane for Pericardial Closure in Operations for Congenital Cardiac Defects," Journal of Thoracic and Cardiovascular Surgery, 1989, vol. 97, pp. 929-934.

Amer M.M.B., "An Accurate Amperometric Glucose Sensor Based Glucometer with Eliminated Cross-Sensitivity," Journal of Medical Engineering & Technology, vol. 26 (5), Sep./Oct. 2002, pp. 208-213.

Amin, et al., "Hypoglycemia Prevalence in Prepubertal Children With Type 1 Diabetes on Standard Insulin Regimen: Use of Continuous Glucose Monitoring System," Diabetes Care, 2003, vol. 26, No. 3, pp. 662-667.

Armour J.C., et al., "Application of Chronic Intravascular Blood Glucose Sensor in Dogs," Diabetes, Dec. 1990, vol. 39, pp. 1519-1526.

Asberg P., et al., "Hydrogels of a Conducting Conjugated Polymer as 3-D Enzyme Electrode," Biosensors Bioelectronics, 2003, vol. 19, pp. 199-207.

Asker.Co.JP, "What is durometer?," /Product/, Kobunshi Keiki Co., Ltd. retrieved from http://www.asker.co.jp/en/products/durometer/analog/about/index.html on May 9, 2018, 2 pages.

Assolant-Vinet C.H., et al., "New Immoblized Enzyme Membranes for Tailor-Made Biosensors", Analytical Letters, 1986, vol. 19(7 &8), pp. 875-885.

ASTM International, Inc., "ASTM, Designation: D2240-05, Standard Test Method for Rubber Property-Durometer Hardness," 2005, 13 pages.

Atanasov P., et al., "Biosensor for Continuous Glucose Monitoring," Biotechnology and Bioengineering, John Wiley & sons Inc, 1994, vol. 43, pp. 262-266.

Atanasov P., et al., "Implantation of a Refillable Glucose Monitoring-Telemetry Device," Biosenors and Bioelectronics, vol. 12 (7), 1997, pp. 669-680.

Aussedat B., et al., "A User-Friendly Method for Calibrating a Subcutaneous Glucose Sensor-Based Hypoglycaemic Alarm," Elsevier Science Limited, Biosensors & Bioelectronic, 1997, vol. 12, No. 11, pp. 1061-1071.

Bailey T.S., et al., "Reduction in Hemoglobin A1C with Real-Time Continuous Glucose Monitoring: Results from a 12-Week Observational Study," Diabetes Technology & Therapeutics, vol. 9 (3), 2007, pp. 203-210.

Baker D.A., et al., "Dynamic Concentration Challenges for Biosensor Characterization," Biosensors & Bioelectronics, vol. 8, 1993, pp. 433-441.

Baker D.A., et al., "Dynamic Delay and Maximal Dynamic Error in Continuous Biosensors," Analytical Chemistry, vol. 68 (8), Apr. 15, 1996, pp. 1292-1297.

Bard A.J., et al., "Electrochemical Methods," Fundamentals and Applications, John Wiley & Sons, New York, 1980, pp. 173-175.

Bardeletti G., et al., "A Reliable L-Lactate Electrode with a New Membrane for Enzyme Immobilization for Amperometric Assay of Lactate," Analytica Chemica Acta, vol. 187, 1986, pp. 47-54.

Beach R.D., et al., "Subminiature Implantable Potentiostat and Modified Commercial Telemetry Device for Remote Glucose Monitoring," IEEE Transactions on Instrumentation and Measurement, vol. 48 (6), Dec. 1999, pp. 1239-1245.

Bellucci F., et al., "Electrochemical Behaviour of Graphite-Epoxy Composite Materials (GECM) in Aqueous Salt Solutions," Journal of Applied Electrochemistry, vol. 16 (1), Jan. 1986, pp. 15-22.

Bennion N., et al., "Alternate Site Glucose Testing: a Crossover Design," Diabetes Technology & Therapeutics, vol. 4 (1), 2002, pp. 25-33.

Berghe V.D., "Tight Blood Glucose Control with Insulin in "Real-Life" Intensive Care," Mayo Clinic Proceedings, vol. 79 (8), Aug. 2004, pp. 977-978.

Bergveld, et al., "Fabrication and Mass Production," Advances in Biosensors, Supplement 1, Chapter 6, 1993, pp. 165-186.

Bertrand C., et al., "Multipurpose Electrode with Different Enzyme Systems Bound to Collagen Films," Analytica Chemica Acta, 1981, vol. 126, pp. 23-34.

Bessman S.P., et al., "Progress toward a Glucose Sensor for the Artificial Pancreas," Proceedings of a Workshop on Ion-Selective Microelectrodes, Jun. 4-5, 1973, Boston University, 1973, pp. 189-197.

Biermann E., et al., "How Would Patients Behave if they were Continually Informed of their Blood Glucose Levels? A Simulation Study Using a "Virtual" Patient," Diabetes Technology & Therapeutics, vol. 10 (3), 2008, pp. 178-187.

Bindra D.S., et al., "Design and in Vitro Studies of a Needle-Type Glucose Sensor for Subcutaneous Monitoring," Analytical Chemistry, vol. 63, Sep. 1, 1991, pp. 1692-1696.

Bindra D.S., et al., "Pulsed Amperometric Detection of Glucose in Biological Fluids at a Surface-Modified Gold Electrode," Analytical Chemistry, vol. 61 (22), Nov. 15, 1989, pp. 2566-2570.

Bisenberger M., et al., "A Triple-Step Potential Waveform at Enzyme Multisensors with Thick-Film Gold Electrodes for Detection of Glucose and Sucrose," Sensors and Actuators B, vol. 28, 1995, pp. 181-189.

Bland J.M., et al., "A Note on the Use of the Intraclass Correlation Coefficient in the Evaluation of Agreement between Two Methods of Measurement," Computers in Biology and Medicine, vol. 20 (5), 1990, pp. 337-340.

Bland J.M., et al., "Statistical Methods for Assessing Agreement Between Two Methods of Clinical Measurement," The Lancet, Feb. 8, 1986, pp. 307-310.

Blank T.B., et al., "Clinical Results From a Non-Invasive Blood Glucose Monitor," Optical Diagnostics and Sensing of Biological Fluids and Glucose and Cholesterol Monitoring II, Proceedings of SPIE, vol. 4624, 2002, pp. 1-10.

(56) **References Cited**

OTHER PUBLICATIONS

Bobbioni-Harsch E., et al., "Lifespan of Subcutaneous Glucose Sensors and their Performances during Dynamic Glycaemia Changes in Rats," J. Biomed. Eng., vol. 15, 1993, pp. 457-463.

Bode B.W., "Clinical Utility of the Continuous Glucose Monitoring System," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S35-S41.

Bode B.W., et al., "Continuous Glucose Monitoring Used to Adjust Diabetes Therapy Improves Glycosylated Hemoglobin: A Pilot Study," Diabetes Research and Clinical Practice, vol. 46, 1999, pp. 183-190.

Bode B.W., et al., "Using the Continuous Glucose Monitoring System to Improve the Management of Type 1 Diabetes," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S43-S48.

Boedeker Plastics Inc, "Polyethylene Specifications," Polyethylene Data Sheet, Retrieved from http://www.boedeker.com/polye.sub.--p. htm on Aug. 19, 2009, 4 pages.

Boland E., et al., "Limitations of Conventional Methods of Self-Monitoring of Blood Glucose," Diabetes Care, vol. 24 (11), Nov. 2001, pp. 1858-1862.

Bolinder J., et al., "Microdialysis Measurement of the Absolute Glucose Concentration in Subcutaneous Adipose Tissue Allowing Glucose Monitoring in Diabetic Patients," Rapid Communication, Diabetologia, vol. 35, 1992, pp. 1177-1180.

Bolinder J., et al., "Self-Monitoring of Blood Glucose in Type I Diabetic Patients: Comparison with Continuous Microdialysis Measurements of Glucose in Subcutaneous Adipose Tissue during Ordinary Life Conditions," Diabetes Care, vol. 20 (1), Jan. 1997, pp. 64-70.

Bott A.W., "A Comparison of Cyclic Voltammetry and Cyclic Staircase Voltammetry," Current Separations, vol. 16 (1), 1997, pp. 23-26.

Bott A.W., "Electrochemical Methods for the Determination of Glucose," Current Separations, vol. 17 (1), 1998, pp. 25-31.

Bowman L., et al., "The Packaging of Implantable Integrated Sensors," IEEE Transactions in Biomedical Engineering, vol. BME-33 (2), Feb. 1986, pp. 248-255.

Brauker J H., et al., "Neovascularization of Synthetic Membranes Directed by Membrane Microarchitecture," Journal of Biomedical Material Research, 1995, vol. 29, pp. 1517-1524.

Brauker J., "Unraveling Mysteries at the Biointerface: Molecular Mediator of Inhibition of Blood Vessel Formation in the Foreign Body Capsule Revealed," SurFACTS in Biomaterials, vol. 6 (3), 2001, 2 pages.

Brauker J., et al., "Local Inflammatory Response Around Diffusion Chambers Containing Xenografts," Transplantation, vol. 61 (12), Jun. 27, 1996, pp. 1671-1677.

Brauker, et al., "Sustained Expression of High Levels of Human Factor IX from Human Cells Implanted within an Immunoisolation Device into Athymic Rodents," Human Gene Therapy, Apr. 10, 1998, vol. 9, pp. 879-888.

Braunwald E., "Biomarkers in Heart Failure," Medical Progress, The New England Journal of Medicine, vol. 358, May 15, 2008, pp. 2148-2159.

Bremer T., et al., "Is Blood Glucose Predictable from Previous Values? A Solicitation for Data," Perspectives in Diabetes, vol. 48, Mar. 1999, pp. 445-451.

Bremer T.M., et al., "Benchmark Data from the Literature for Evaluation of New Glucose Sensing Technologies," Diabetes Technology & Therapeutics, vol. 3 (3), 2001, pp. 409-418.

Brooks S.L., et al., "Development of an On-line Glucose Sensor for Fermentation Monitoring," Biosensors, vol. 3, 1987/1988, pp. 45-56.

Bruckel J., et al., "In Vivo Measurement of Subcutaneous Glucose Concentrations with an Enzymatic Glucose Sensor and a Wick Method," Klin Wochenschr, vol. 67, 1989, pp. 491-495.

Brunner G.A., et al., "Validation of Home Blood Glucose Meters with Respect to Clinical and Analytical Approaches," Diabetes Care, vol. 21(4), Apr. 1998, pp. 585-590.

Brunstein E., et al., "Preparation and Validation of Implantable Electrodes for the Measurement of Oxygen and Glucose," Biomed Biochim. Acta, vol. 48 (11/12), 1989, pp. 911-917.

Cai Q., et al., "A Wireless, Remote Query Glucose Biosensor Based on a pH-Sensitive Polymer," Analytical Chemistry, vol. 76 (14), Jul. 15, 2004, pp. 4038-4043.

Cameron T., et al., "Micromodular Implants to Provide Electrical Stimulation of Paralyzed Muscles and Limbs," IEEE Transactions on Biomedical Engineering, vol. 44 (9), Sep. 1997, pp. 781-790.

Campanella L., et al., "Biosensor for Direct Determination of Glucose and Lactate in Undiluted Biological Fluids," Biosensors & Bioelectronics, vol. 8, 1993, pp. 307-314.

Candas B., et al., "An Adaptive Plasma Glucose Controller Based on a Nonlinear Insulin/Glucose Model," IEEE Transactions on Biomedical Engineering, vol. 41 (2), Feb. 1994, pp. 116-124.

Cass A.E.G., et al., "Ferrocene-Mediated Enzyme Electrodes for Amperometric Determination of Glucose," Analytical Chemistry, vol. 56 (4), Apr. 1984, pp. 667-671.

Cassidy J.F., et al., "Novel Electrochemical Device for the Detection of Cholesterol or Glucose," Analyst, vol. 118, Apr. 1993, pp. 415-418.

Chase H.P., et al., "Continuous Subcutaneous Glucose Monitoring in Children with Type 1 Diabetes," Pediatrics, vol. 107 (2), Feb. 2001, pp. 222-226.

Chatterjee G., et al., "Poly(ether urethane) and Poly(ether urethane urea) Membranes with High H2S/CH4 Selectivity," Journal of Membrane Science, vol. 135, 1997, pp. 99-106.

Chen C., et al., "A Noninterference Polypyrrole Glucose Biosensor," Biosensors and Bioelectronics, vol. 22, 2006, pp. 639-643.

Chen T., et al., "Defining the Period of Recovery of the Glucose Concentration after its Local Perturbation by the Implantation of a Miniature Sensor," Clinical Chemistry and Laboratory Medicine, vol. 40 (8), 2002, pp. 786-789.

Chia C.W., et al., "Glucose Sensors: Toward Closed Loop Insulin Delivery," Endocrinology and Metabolism Clinics of North America, vol. 33, 2004, pp. 175-195.

Choleau C., et al., "Calibration of a Subcutaneous Amperometric Glucose Sensor Implanted for 7 Days in Diabetic Patients Part 2. Superiority of the One-point Calibration Method," Biosensors and Bioelectronics, vol. 17 (8), 2002, pp. 647-654.

Choleau C., et al., "Calibration of a Subcutaneous Amperometric Glucose Sensor Part 1. Effect of Measurement Uncertainties on the Determination of Sensor Sensitivity and Background Current," Biosensors and Bioelectronics, vol. 17, 2002, pp. 641-646.

Ciba Specialty Chemicals, "Ciba® IRGACURE® 2959," Coating Effects Segment, Photoinitiator Product Description, Basel Switzerland, Apr. 2, 1998, 3 pages.

Claremont D.J., et al., "Potentially-Implantable, Ferrocene-Mediated Glucose Sensor," Journal of Biomedical Engineering, vol. 8, Jul. 1986, pp. 272-274.

Claremont D.J., et al., "Subcutaneous Implantation of a Ferrocene-Mediated Glucose Sensor in Pigs," Diabetologia, vol. 29, 1986, pp. 817-821.

Clark L.C., et al., "Configurational Cyclic Voltammetry: Increasing the Specificity and Reliability of Implanted Electrodes," IEEE/Ninth Annual Conference of the Engineering in Medicine and Biollogy Society, 1987, pp. 0782-0783.

Clark L.C., et al., "Long-Term Stability of Electroenzymatic Glucose Sensors Implanted in Mice," vol. XXXIV, Transactions—American Society for Artificial Internal Organs, 1988, vol. 34, pp. 259-265.

Clark L.C., et al., "One-Minute Electrochemical Enzymic Assay for Cholesterol in Biological Materials," Clinical Chemistry, vol. 27 (12), 1981, pp. 1978-1982.

Clarke W.L., et al., "Evaluating Clinical Accuracy of Systems for Self Monitoring of Blood Glucose," Technical Articles, Diabetes Care, vol. 10 (5), Sep.-Oct. 1987, pp. 622-628.

Clarke W.L., et al., "Evaluating the Clinical Accuracy of Two Continuous Glucose Sensors Using Continuous Glucose-Error Grid Analysis," Emerging Treatment and Technologies, Diabetes Care, vol. 28(10), Oct. 2005, pp. 2412-2417.

(56) **References Cited**

OTHER PUBLICATIONS

Colangelo V.J., et al., "Corrosion Rate Measurements in Vivo," Journal of Biomedical Materials Research, vol. 1, 1967, pp. 405-414.

Colowick S.P., et al., "Methods in Enzymology," vol. XLIV, Immobilized Enzymes, Edited by Mosbach K, New York Academic Press, 1976, 11 pages.

Copeland J.G., et al., "Synthetic Membrane Neo-Pericardium Facilitates Total Artificial Heart Explanation," The Journal of Heart Lung Transplantation, vol. 20(6), Jun. 2001, pp. 654-656.

Coulet P.R., "Polymeric Membranes and Coupled Enzymes in the Design of Biosensors," Journal of Membrane Science, 1992, vol. 68, pp. 217-228.

Coulet P.R., et al., "Enzymes Immobilized on Collagen Membranes: A Tool for Fundamental Research and Enzyme Engineering," Journal of Chromatography, vol. 215, 1981, pp. 65-72.

Cox D.J., et al., "Accuracy of Perceiving Blood Glucose in IDDM," Diabetes Care, vol. 8 (6), Nov.-Dec. 1985, pp. 529-536.

Csoregi E., et al., "Amperometric Microbiosensors for Detection of Hydrogen Peroxide and Glucose Based on Peroxidase-Modified Carbon Fibers," Electroanalysis, vol. 6, 1994, pp. 925-933.

Csoregi E., et al., "Design and Optimization of a Selective Subcutaneously Implantable Glucose Electrode Based on 'Wired' Glucose Oxidase," Analytical Chemistry, vol. 67 (7), Apr. 1, 1995, pp. 1240-1244.

Csoregi E., et al., "Design, Characterization and One-Point in Vivo Calibration of a Subcutaneously Implanted Glucose Electrode," American Chemical Society, Analytical Chemistry, vol. 66 (19), Oct. 1, 1994, pp. 3131-3138.

Currie J.F., et al., "Novel Non-Intrusive Trans-Dermal Remote Wireless Micro-Fluidic Monitoring System Applied to Continuous Glucose and Lactate Assays for Casualty Care and Combat Readiness Assessment," RTO HFM Symposium, RTO-MP-HFM-109, Aug. 16-18, 2004, pp. 24-1-24-18.

Dai W.S., et al., "Hydrogel Membranes with Mesh Size Asymmetry based on the Gradient Crosslinking of Poly(Vinyl Alcohol)," Journal of Membrane Science, 1999, vol. 156, pp. 67-79.

Danielsson B., et al., "Enzyme Thermistors," Methods in Enzymology, vol. 137, 1988, pp. 181-197.

D'Arrigo, et al., "Porous-Si Based Bio Reactors for Glucose Monitoring and Drugs Production," Proceedings of SPIE, 2003, vol. 4982, pp. 178-184.

Dassau E., et al., "In Silico Evaluation Platform for Artificial Pancreatic β- Cell Development a Dynamic Simulator for Closed-Loop Control with Hardware-in-the-loop," Diabetes Technology & Therapeutics, vol. 11 (3), 2009, pp. 1-8.

Dassau E., et al., "Real-Time Hypoglycemia Prediction Suite Using Continuous Glucose Monitoring," Emerging Treatment and Technologies, Diabetes Care, vol. 33 (6), Jun. 2010, pp. 1249-1254.

Davies M.L., et al., "Polymer Membranes in Clinical Sensor Applications," An overview of membrane function, Biomaterials, vol. 13 (14), 1992, pp. 971-978.

Davis G., et al., "Bioelectrochemical Fuel Cell and Sensor Based on a Quinoprotein, Alcohol Dehydrogenase," Enzyme and Microbial Technology, vol. 5 (5), Sep. 1983, pp. 383-388.

Declaration presented during Opposition proceedings EP2407094, PACE L., Nov. 13, 2015. 6 pages.

Declaration presented during Opposition proceedings EP2407094, Schoonmaker R., Dec. 14, 2015, 10 pages.

Definition of plunger: https://www.merriam-webster.com/dictionary/plunger,dated Nov. 7, 2016, 2 pages.

Deutsch T., et al., "Time Series Analysis and Control of Blood Glucose Levels in Diabetic Patients," Computer Methods and Programs in Biomedicine, Elsevier Scientific Publishers, vol. 41, 1994, pp. 167-182.

Dixon B.M., et al., "Characterization in Vitro and in Vivo of the Oxygen Dependence of an Enzyme/Polymer Biosensor for Monitoring Brain Glucose," Journal of Neuroscience Methods, vol. 119, 2002, pp. 135-142.

Dobson, et al.,"1-Butyrul-Glycerol: A Novel Angiogenesis Factor Secreted by Differentiating Adipocytes," Cell, Apr. 20, 1990, vol. 61 (2), pp. 223-230.

DuPont, "Dimension® AR Clinical Chemistry System," The Chemistry Analyzer that Makes the most of your Time, Money and Effort, Dade International, Chemistry Systems, Newark, 1998, 18 pages.

Durliat H., et al., "Spectrophotometric and Electrochemical Determinations of L(+)-Lactate in Blood by Use of Lactate Dehydrogenase from Yeast," Clinical Chemistry, vol. 22 (11), 1976, pp. 1802-1805.

Edwards Lifesciences, "Accuracy for You and Your Patients," Marketing materials, 2002, 4 pages.

El Degheidy M.M., et al., "Optimization of an Implantable Coated Wire Glucose Sensor," Journal of Biomedical Engineering, vol. 8, Apr. 1986, pp. 121-129.

Elco Diagnostics Company, "Direct 30/30® Blood Glucose Sensor," Markwell Medical Catalog, 1990, 1 pages.

El-Khatib F.H., et al., "Adaptive Closed-Loop Control Provides Blood-Glucose Regulation Using Dual Subcutaneous Insulin and Glucagon Infusion in Diabetic Swine," Journal of Diabetes Science and Technology, Diabetes Technology Society, vol. 1 (2), 2007, pp. 181-192.

El-Sa'ad L., et al., "Moisture Absorption by Epoxy Resins: The Reverse Thermal Effect," Journal of Materials Science, vol. 25, 1990, pp. 3577-3582.

English D., et al., "Platelet-Released Phospholipids Link Haemostasis and Angiogenesis," Cardiovascular Research, 2001, vol. 49, pp. 588-599.

Ernst H., et al., "Reliable Glucose Monitoring Through the Use of Microsystem Technology," Analytical Bioanalytical Chemistry, vol. 373, 2002, pp. 758-761.

European Electronic File History for EP Application No. 10195447.7, filed Jul. 13, 2005—withdrawn, 200 pages.

European Electronic File History for EP Application No. 10195504.5, filed Jul. 13, 2005—withdrawn, 186 pages.

European Electronic File History for EP Application No. 11182615.2, filed Feb. 22, 2006, 344 pages.

European Electronic File History for EP Application No. 11182630.1, filed Sep. 23, 2011, 336 pages.

European Electronic File History for EP Application No. 14184330.0, filed Sep. 10, 2014, 377 pages.

European Electronic File History for EP Application No. 17198022.0, filed Oct. 24, 2017, 339 pages.

European Electronic File History for EP Patent Application No. 06773682.7 filed Jun. 20, 2006, 1697 pages.

European Electronic File History for EP Patent No. 1804650 (05771643.3) granted Mar. 14, 2012, 3733 pages.

European Electronic File History for EP Patent No. 1855588 (06748336.2), granted Nov. 21, 2007, 862 pages.

European Electronic File History for EP Patent No. 1986543 (06736006.5) granted Dec. 13, 2011, 1895 pages.

European Electronic File History for EP Patent No. 2322094 (10195496.4) granted Mar. 5, 2014, 1778 pages.

European Electronic File History for EP Patent No. 2327362 (10195517.7) granted Nov. 13, 2013, 423 pages.

European Electronic File History for EP Patent No. 2327984 (10195521.9) granted Feb. 25, 2015, 394 pages.

European Electronic File History for EP Patent No. 2329770 (10195509.4) granted Sep. 10, 2014, 1274 pages.

European Electronic File History for EP Patent No. 2332466 (10195483.2) granted Apr. 9, 2014, 484 pages.

European Electronic File History for EP Patent No. 2335582 (10195511.0) granted Sep. 10, 2014, 608 pages.

European Electronic File History for EP Patent No. 2335583 (10195520.1) granted Mar. 18, 2015, 677 pages.

European Electronic File History for EP Patent No. 2335584 (10195519.3) granted Jun. 17, 2015, 796 pages.

European Electronic File History for EP Patent No. 2335585 (10195508.6) granted Sep. 7, 2016, 722 pages.

European Electronic File History for EP Patent No. 2335586 (10195514.4) granted Feb. 19, 2014, 526 pages.

European Electronic File History for EP Patent No. 2335587 (10195518.5) granted Feb. 19, 2014, 5531 pages.

(56) **References Cited**

OTHER PUBLICATIONS

European Electronic File History for EP Patent No. 2407094 (11182622.8) granted Oct. 22, 2014, 2417 pages.
European Electronic File History for EP Patent No. 2499969 (12151826.0) granted Jun. 4, 2014, 814 pages.
European Electronic File History for EP Patent No. 2517623 (12151823.7) granted Mar. 2, 2016, 864 pages.
European Electronic File History for EP Patent No. 2532302 (12151819.5) granted Nov. 18, 2015, 845 pages.
European Electronic File History for EP Patent No. 2532305 (12171365.5) granted Aug. 13, 2014, 504 pages.
European Electronic File History for EP Patent No. 2561807 (12193934.2) granted Oct. 5, 2016, revoked Jan. 31, 2019, 2496 pages.
European Electronic File History for EP Patent No. 2596747 (13156245.6) granted Oct. 25, 2017, 1091 pages.
European Electronic File History for EP Publication No. 3001952 (15195173.8), published Apr. 16, 2016, 237 pages.
European Electronic File History for EP Publication No. 3111832 (16179208.0), published Jan. 4, 2017—pending, 191 pages.
Examination Report for European Application No. 09729717.0, dated Jun. 6, 2013, 3 pages.
Extended European Search Report for Application No. 06718980.3 dated Mar. 9, 2010, 8 pages.
Extended European Search Report for Application No. 19185160.9 dated Sep. 23, 2019, 8 pages.
Extended European Search Report for Application No. 98908875.2 dated Apr. 29, 2004, 5 pages.
Fabietti P.G., et al., "Clinical Validation of a New Control-Oriented Model of Insulin and Glucose Dynamics in Subjects with Type 1 Diabetes," Diabetes Technology & Therapeutics, vol. 9 (4), 2007, pp. 327-338.
Fahy B.G., et al., "An Analysis: Hyperglycemic Intensive Care Patients Need Continuous Glucose Monitoring-Easier Said Than Done," Journal of Diabetes Science and Technology, Diabetes Technology Society, vol. 2 (2), Mar. 2008, pp. 201-204.
Fare T.L., et al., "Functional Characterization of a Conducting Polymer-Based Immunoassay System," Biosonsors & Bioelectronics, vol. 13 (3-4), 1998, pp. 459-470.
Farlex, Inc, Definition of term "elastomeric", Free Dictionary, Copyright 2008, retrieved from http://www.thefreedictionary.com/elastomeric, 3 pages.
Farlex, Inc., "Statistical Distribution—Definition of Statistical Distribution by the Free Dictionary—Thesaurus," 2003-2016, 2 pages.
Feldman B., et al., "A Continuous Glucose Sensor Based on Wired EnzymeTM Technology-Results from a 3-Day Trial in Patients with Type 1 Diabetes," Diabetes Technology & Therapeutics, vol. 5 (5), 2003, pp. 769-779.
Feldman B., et al., "Correlation of Glucose Concentrations in Interstitial Fluid and Venous Blood During Periods of Rapid Glucose Change," Abbott Diabetes Care, Inc. Freestyle Navigator Continuous Glucose Monitor Pamphlet, 2004, 1 page.
File History of European Patent Application No. 19185160.9, filed on Mar. 10, 2006, downloaded Oct. 10, 2019, 541 pages.
File History of U.S. Appl. No. 09/334,996, filed Jun. 17, 1999, 36 pages.
File History of U.S. Appl. No. 09/447,227, filed Nov. 22, 1999, 1184 pages.
File History of U.S. Appl. No. 09/489,588, filed Jan. 21, 2000, 412 pages.
File History of U.S. Appl. No. 09/636,369, filed Aug. 11, 2000, 91 pages
File History of U.S. Appl. No. 09/916,711, filed Jul. 27, 2001, 400 pages.
File History of U.S. Appl. No. 09/916,858, filed Jul. 27, 2001, 243 pages.
File History of U.S. Appl. No. 10/153,356, filed May 22, 2002, 326 pages.
File History of U.S. Appl. No. 10/632,537, filed Aug. 1, 2003, 211 pages.
File History of U.S. Appl. No. 10/633,329, filed Aug. 1, 2003, 711 pages.
File History of U.S. Appl. No. 10/633,367, filed Aug. 1, 2003, 432 pages.
File History of U.S. Appl. No. 10/633,404, filed Aug. 1, 2003, 270 pages.
File History of U.S. Appl. No. 10/646,333, filed Aug. 22, 2003, 303 pages.
File History of U.S. Appl. No. 10/647,065, filed Aug. 22, 2003, 203 pages.
File History of U.S. Appl. No. 10/648,849, filed Aug. 22, 2003, 803 pages.
File History of U.S. Appl. No. 10/695,636, filed Oct. 28, 2003, 194 pages.
File History of U.S. Appl. No. 10/789,359, filed Feb. 26, 2004, 361 pages.
File History of U.S. Appl. No. 10/838,658, filed May 3, 2004, 748 pages.
File History of U.S. Appl. No. 10/838,909, filed May 3, 2004, 356 pages.
File History of U.S. Appl. No. 10/838,912, filed May 3, 2004, 1288 pages.
File History of U.S. Appl. No. 10/842,716, filed May 10, 2004, 670 pages.
File History of U.S. Appl. No. 10/846,150, filed May 14, 2004, 382 pages.
File History of U.S. Appl. No. 10/885,476, filed Jul. 6, 2004, 226 pages.
File History of U.S. Appl. No. 10/896,637, filed Jul. 21, 2004, 295 pages.
File History of U.S. Appl. No. 10/896,639, filed Jul. 21, 2004, 337 pages.
File History of U.S. Appl. No. 10/896,772, filed Jul. 21, 2004, 210 pages.
File History of U.S. Appl. No. 10/897,312, filed Jul. 21, 2004, 238 pages.
File History of U.S. Appl. No. 10/897,377, filed Jul. 21, 2004, 178 pages.
File History of U.S. Appl. No. 11/039,269, filed Jan. 19, 2005, 209 pages.
File History of U.S. Appl. No. 11/077,643, filed Mar. 10, 2005, 253 pages.
File History of U.S. Appl. No. 11/077,693 filed Mar. 10, 2005, 755 pages.
File History of U.S. Appl. No. 11/077,713, filed Mar. 10, 2005, 1025 pages.
File History of U.S. Appl. No. 11/077,714, filed Mar. 10, 2005, 320 pages.
File History of U.S. Appl. No. 11/077,715, filed Mar. 10, 2005, 717 pages.
File History of U.S. Appl. No. 11/077,739, filed Mar. 10, 2005, 778 pages.
File History of U.S. Appl. No. 11/077,740, filed Mar. 10, 2005, 921 pages.
File History of U.S. Appl. No. 11/077,763, filed Mar. 10, 2005, 515 pages.
File History of U.S. Appl. No. 11/077,765, filed Mar. 10, 2005, 932 pages.
File History of U.S. Appl. No. 11/077,883, filed Mar. 10, 2005, 1159 pages.
File History of U.S. Appl. No. 11/078,072, filed Mar. 10, 2005, 1334 pages.
File History of U.S. Appl. No. 11/078,230, filed Mar. 10, 2005, 535 pages.
File History of U.S. Appl. No. 11/078,232, filed Mar. 10, 2005, 256 pages.
File History of U.S. Appl. No. 11/157,365, filed Jun. 21, 2005, 977 pages.
File History of U.S. Appl. No. 11/157,746, filed Jun. 21, 2005, 603 pages.
File History of U.S. Appl. No. 11/158,227, filed Jul. 21, 2004, 474 pages.

(56) **References Cited**

OTHER PUBLICATIONS

File History of U.S. Appl. No. 11/201,445, filed Jul. 21, 2004, 120 pages.
File History of U.S. Appl. No. 11/280,102, filed Nov. 16, 2005, 109 pages.
File History of U.S. Appl. No. 11/280,672, filed Nov. 16, 2005, 572 pages.
File History of U.S. Appl. No. 11/333,837, filed Jan. 17, 2006, 672 pages.
File History of U.S. Appl. No. 11/334,107, filed Jan. 17, 2006.
File History of U.S. Appl. No. 11/335,879, filed Jan. 18, 2006, 499 pages.
File History of U.S. Appl. No. 11/360,262, filed Feb. 22, 2006, 766 pages.
File History of U.S. Appl. No. 11/360,299, filed Feb. 22, 2006, 676 pages.
File History of U.S. Appl. No. 11/360,819, filed Feb. 22, 2006, 778 pages.
File History of U.S. Appl. No. 11/373,628, filed Mar. 9, 2006, 532 pages.
File History of U.S. Appl. No. 11/411,656, filed Apr. 26, 2006, 285 pages.
File History of U.S. Appl. No. 11/415,593, filed May 2, 2006, 160 pages.
File History of U.S. Appl. No. 11/415,999, filed May 2, 2006, 154 pages.
File History of U.S. Appl. No. 11/416,375, filed May 2, 2006, 153 pages.
File History of U.S. Appl. No. 11/439,559, filed May 23, 2006, 155 pages.
File History of U.S. Appl. No. 11/439,630, filed May 23, 2006, 1237 pages.
File History of U.S. Appl. No. 11/439,800, filed May 23, 2006, 157 pages.
File History of U.S. Appl. No. 11/445,792 filed Jun. 1, 2006, 811 pages.
File History of U.S. Appl. No. 11/503,367, filed Aug. 10, 2006, 1169 pages.
File History of U.S. Appl. No. 11/618,706, filed Dec. 29, 2006, 288 pages.
File History of U.S. Appl. No. 11/681,145, filed Mar. 1, 2007, 684 pages.
File History of U.S. Appl. No. 11/690,752, filed Mar. 23, 2007, 528 pages.
File History of U.S. Appl. No. 11/695,607, filed Mar. 4, 2007, 615 pages.
File History of U.S. Appl. No. 11/734,178, filed Apr. 11, 2007, 512 pages.
File History of U.S. Appl. No. 11/734,184, filed Apr. 11, 2007, 531 pages.
File History of U.S. Appl. No. 11/734,203, filed Apr. 11, 2007, 606 pages.
File History of U.S. Appl. No. 11/742,546, filed Apr. 30, 2007, 348 pages.
File History of U.S. Appl. No. 11/766,747, filed Jun. 21, 2007, 789 pages.
File History of U.S. Appl. No. 11/797,520, filed May 3, 2007, 406 pages.
File History of U.S. Appl. No. 11/797,521, filed May 3, 2007, 374 pages.
File History of U.S. Appl. No. 11/842,139, filed Aug. 21, 2007.
File History of U.S. Appl. No. 11/842,142, filed Aug. 21, 2007.
File History of U.S. Appl. No. 11/842,143, filed Aug. 21, 2007, 235 pages.
File History of U.S. Appl. No. 11/842,146, filed Aug. 20, 2007.
File History of U.S. Appl. No. 11/842,148, filed Aug. 21, 2007.
File History of U.S. Appl. No. 11/842,149, filed Aug. 21, 2007, 256 pages.
File History of U.S. Appl. No. 11/842,151 filed Aug. 21, 2007, 252 pages.
File History of U.S. Appl. No. 11/842,154, filed Aug. 21, 2007.
File History of U.S. Appl. No. 11/842,157, filed Aug. 21, 2007, 167 pages.
File History of U.S. Appl. No. 11/925,603, filed Oct. 26, 2007, 453 pages.
File History of U.S. Appl. No. 11/928,968, filed Oct. 30, 2007, 741 pages.
File History of U.S. Appl. No. 12/101,790, filed Apr. 11, 2008, 431 pages.
File History of U.S. Appl. No. 12/101,806, filed Apr. 11, 2008, 611 pages.
File History of U.S. Appl. No. 12/101,810, filed Apr. 11, 2008, 569 pages.
File History of U.S. Appl. No. 12/137,396, filed Jun. 11, 2008, 278 pages.
File History of U.S. Appl. No. 12/175,391, filed Jul. 17, 2008, 564 pages.
File History of U.S. Appl. No. 12/245,618, filed Oct. 3, 2008, 214 pages.
File History of U.S. Appl. No. 12/250,918, filed Oct. 14, 2008, 333 pages.
File History of U.S. Appl. No. 12/273,359, filed Nov. 18, 2008, 312 pages.
File History of U.S. Appl. No. 12/329,496, filed Dec. 5, 2008, 438 pages.
File History of U.S. Appl. No. 12/353,870, filed Jan. 14, 2009, 425 pages.
File History of U.S. Appl. No. 12/359,207, filed Jan. 23, 2009, 456 pages.
File History of U.S. Appl. No. 12/364,786, filed Feb. 3, 2009, 1230 pages.
File History of U.S. Appl. No. 12/391,148, filed Feb. 23, 2009, 484 pages.
File History of U.S. Appl. No. 12/393,887, filed Feb. 26, 2009, 502 pages.
File History of U.S. Appl. No. 12/405,883, filed Mar. 17, 2009, 215 pages.
File History of U.S. Appl. No. 12/437,436, filed May 7, 2009, 524 pages.
File History of U.S. Appl. No. 12/537,245, filed Aug. 6, 2009, 107 pages.
File History of U.S. Appl. No. 12/610,866, filed Nov. 2, 2009, 682 pages.
File History of U.S. Appl. No. 12/683,724, filed Jan. 7, 2010, 497 pages.
File History of U.S. Appl. No. 12/683,755, filed Jan. 7, 2010, 694 pages.
File History of U.S. Appl. No. 12/728,032, filed Mar. 19, 2010, 819 pages.
File History of U.S. Appl. No. 12/728,060, filed Mar. 19, 2010, 323 pages.
File History of U.S. Appl. No. 12/728,061, filed Mar. 19, 2010, 437 pages.
File History of U.S. Appl. No. 12/728,082, filed Mar. 19, 2010, 802 pages.
File History of U.S. Appl. No. 12/729,035, filed Mar. 22, 2010, 466 pages.
File History of U.S. Appl. No. 12/729,048, filed Mar. 22, 2010, 488 pages.
File History of U.S. Appl. No. 12/729,058, filed Mar. 22, 2010, 502 pages.
File History of U.S. Appl. No. 12/730,072, filed Mar. 23, 2010, 278 pages.
File History of U.S. Appl. No. 12/748,154, filed Mar. 26, 2010, 419 pages.
File History of U.S. Appl. No. 12/749,139, filed Mar. 29, 2010, 489 pages.
File History of U.S. Appl. No. 12/749,981, filed Mar. 30, 2010, 662 pages.
File History of U.S. Appl. No. 12/775,315, filed May 6, 2010, 447 pages.
File History of U.S. Appl. No. 12/780,606, filed May 14, 2010, 372 pages.

(56) **References Cited**

OTHER PUBLICATIONS

File History of U.S. Appl. No. 12/780,723, filed May 14, 2010, 519pages.
File History of U.S. Appl. No. 12/780,725, filed May 14, 2010, 415 pages.
File History of U.S. Appl. No. 12/780,739, filed May 14, 2010, 346 pages.
File History of U.S. Appl. No. 12/780,759, filed May 14, 2010, 348 pages.
File History of U.S. Appl. No. 12/853,235, filed Aug. 9, 2010, 374 pages.
File History of U.S. Appl. No. 13/077,884, filed Mar. 31, 2011, 635 pages.
File History of U.S. Appl. No. 13/086,160, filed Apr. 13, 2011, 967 pages.
File History of U.S. Appl. No. 13/116,871, filed May 26, 2011, 475 pages.
File History of U.S. Appl. No. 13/157,031, filed Jun. 9, 2011, 514 pages.
File History of U.S. Appl. No. 13/172,640, filed Jun. 29, 2011, 436 pages.
File History of U.S. Appl. No. 13/361,820, filed Jan. 30, 2012, 383 pages.
File History of U.S. Appl. No. 13/415,721, filed Mar. 8, 2012, 473 pages.
File History of U.S. Appl. No. 13/547,952, filed Jul. 12, 2012, 539 pages.
File History of U.S. Appl. No. 13/548,627, filed Jul. 13, 2012, 361 pages.
File History of U.S. Appl. No. 13/549,313, filed Jul. 13, 2012, 354 pages.
File History of U.S. Appl. No. 13/595,673, filed Aug. 27, 2012, 311 pages.
File History of U.S. Appl. No. 13/607,162, filed Sep. 7, 2012, 420 pages.
File History of U.S. Appl. No. 13/620,574, filed Sep. 14, 2012, 318 pages.
File History of U.S. Appl. No. 13/893,237, filed May 13, 2013, 569 pages.
File History of U.S. Appl. No. 13/903,609, filed May 28, 2013, 401 pages.
File History of U.S. Appl. No. 13/909,962, filed Jun. 4, 2013, 723 pages.
File History of U.S. Appl. No. 13/938,103, filed Jul. 9, 2013, 393 pages.
File History of U.S. Appl. No. 14/144,523, filed Dec. 30, 2013, 405 pages.
File History of U.S. Appl. No. 14/163,346, filed Jan. 24, 2014, 583 pages.
File History of U.S. Appl. No. 14/283,153, filed May 20, 2014, 568 pages.
File History of U.S. Appl. No. 14/293,298, filed Jun. 2, 2014, 565 pages.
File History of U.S. Appl. No. 14/296,735, filed Jun. 5, 2014, 637 pages.
File History of U.S. Appl. No. 14/552,398, filed Nov. 24, 2014, 422 pages.
File History of U.S. Appl. No. 14/590,483, filed Jan. 6, 2015, 556 pages.
File History of U.S. Appl. No. 14/743,777, filed Jun. 18, 2015, 392 pages.
File History of U.S. Appl. No. 14/860,602, filed Sep. 21, 2015, 429 pages.
File History of U.S. Appl. No. 14/923,350, filed Oct. 26, 2015, 476 pages.
File History of U.S. Appl. No. 14/924,030, filed Oct. 27, 2015, 403 pages.
File History of U.S. Appl. No. 15/065,623, filed Mar. 9, 2016, 401 pages.
File History of U.S. Appl. No. 15/201,313, filed Jul. 1, 2016, 800 pages.
File History of U.S. Appl. No. 15/470,766, filed Mar. 27, 2017, 336 pages.
File History of U.S. Appl. No. 15/686,650, filed Aug. 25, 2017, 296 pages.
File History of U.S. Appl. No. 15/719,298, filed Sep. 28, 2017, 258 pages.
File History of U.S. Appl. No. 15/787,595, filed Oct. 18, 2017, 230 pages.
File History of U.S. Appl. No. 15/797,986, filed Oct. 30, 2017, 342 pages.
File History of U.S. Appl. No. 15/798,097, filed Oct. 30, 2017, 323 pages.
File History of U.S. Appl. No. 15/877,311, filed Jan. 22, 2018, 208 pages.
File History of U.S. Appl. No. 15/891,201, filed Feb. 7, 2018, 364 pages.
File History of U.S. Appl. No. 15/967,338, filed Apr. 30, 2018, 191 pages.
File History of U.S. Appl. No. 16/017,668, filed Jun. 25, 2018, 308 pages.
File History of U.S. Appl. No. 16/133,469, filed Sep. 17, 2018, 178 pages.
File History of U.S. Appl. No. 16/137,411, filed Sep. 20, 2018, 174 pages.
File History of U.S. Appl. No. 16/179,662, filed Nov. 2, 2018, 272 pages.
File History of U.S. Appl. No. 16/181,678, filed Nov. 6, 2018,155 pages.
File History of U.S. Appl. No. 60/362,899, filed Mar. 8, 2002, 67 pages.
File History of U.S. Appl. No. 60/527,892, filed Dec. 8, 2003, 25 pages.
File History of U.S. Appl. No. 60/528,382, filed Dec. 9, 2003, 182 pages.
File History of U.S. Appl. No. 60/587,787, filed Jul. 13, 2004, 70 pages.
File History of U.S. Appl. No. 60/587,800, filed Jul. 13, 2004, 29 pages.
File History of U.S. Appl. No. 60/614,683, filed Sep. 30, 2004, 465 pages.
File History of U.S. Appl. No. 60/614,764, filed Sep. 30, 2004, 657 pages.
File History of U.S. Appl. No. 60/660,743, filed Mar. 10, 2005, 104 pages.
File History of U.S. Appl. No. 90/011,086, filed Jul. 8, 2010, 169 pages.
File History of U.S. Appl. No. 90/011,351, filed Nov. 22, 2010, 260 pages.
File History of U.S. Appl. No. 90/011,663, filed Apr. 29, 2011, 306 pages.
File History of U.S. Appl. No. 90/011,887, filed Oct. 7, 2011, 769 pages.
File History of U.S. Appl. No. 90/012,558, filed Sep. 13, 2012, 294 pages.
File History of U.S. Appl. No. 95/001,038, filed Apr. 17, 2008, 1332 pages.
File History of U.S. Appl. No. 95/001,039, filed Apr. 17, 2008, 1409 pages.
File History of U.S. Appl. No. 95/002,333, filed Sep. 14, 2012, 556 pages.
File History of U.S. Appl. No. 11/334,876, filed Jan. 18, 2006, 751 pages.
File History of U.S. Appl. No. 11/360,250, filed Feb. 22, 2006, 950 pages.
File History of U.S. Appl. No. 11/360,252, filed Feb. 22, 2006, 594 pages.
File History of U.S. Appl. No. 90/011,333, filed Nov. 15, 2010, 315 pages.
File History of U.S. Appl. No. 90/011,720, filed Nov. 15, 2010, 361 pages.

(56) **References Cited**

OTHER PUBLICATIONS

Fischer U., et al., "Assessment of Subcutaneous Glucose Concentration: Validation of the Wick Technique as a Reference for Implanted Electrochemical Sensors in Normal and Diabetic Dogs," Diabetologia, vol. 30, 1987, pp. 940-945.
Fischer U., et al., "Hypoglycaemia-Warning by Means of Subcutaneous Electrochemical Glucose Sensors: An Animal Study," Horm. Metab. Res, vol. 27, 1995, p. 53. (Abstract Only).
Fischer U., et al., "Oxygen Tension at the Subcutaneous Implantation Site of Glucose Sensors," Biomed. Biochim. Acta, vol. 48 (11/12), 1989, pp. 965-971.
Freedman D., et al., "Statistics," Second Edition, W.W. Norton & Company, New York & London, 1991, p. 74 (3 pages).
Freiberger P., "Video Game Takes on Diabetes Superhero 'Captain Novolin' Offers Treatment Tips," Fourth Edition, Jun. 26, 1992, Business Section, 2 pages.
Frohnauer M.K., et al., "Graphical Human Insulin Time-Activity Profiles Using Standardized Definitions," Diabetes Technology & Therapeutics, vol. 3 (3), 2001, pp. 419-429.
Frost M.C., et al., "Implantable Chemical Sensors for Real-Time Clinical Monitoring: Progress and Challenges," Current Opinion in Chemical Biology, Analytical Techniques, vol. 6, 2002, pp. 633-641.
Gabby R.A., et al., "Optical Coherence Tomography-Based Continuous Noninvasive Glucose Monitoring in Patients with Diabetes," Diabetes Technology & Therapeutics, vol. 10, Nov. 3, 2008, pp. 188-193.
Ganesan N., et al., "Gold Layer-Based Dual Crosslinking Procedure of Glucose Oxidase with Ferrocene Monocarboxylic Acid Provides a Stable Biosensor," Analytical Biochemistry, Notes & Tips, vol. 343, 2005, pp. 188-191.
Ganesh A., et al., "Evaluation of the VIA® Blood Chemistry Monitor for Glucose in Healthy and Diabetic Volunteers," Journal of Diabetes Science and Technology, vol. 2 (2), Mar. 2008, pp. 182-193.
Gao S., et al., "Determination of Interfacial Parameters of Cellulose Acetate Membrane Materials by HPLC," Journal of Liquid Chromatography, 1989, vol. 12(11), pp. 2083-2092.
Garg S.K., et al., "Correlation of Fingerstick Blood Glucose Measurements With GlucoWatch Biographer Glucose Results in Young Subjects With Type 1 Diabetes," Emerging Treatments and Technologies, Diabetes Care, vol. 22 (10), Oct. 1999, pp. 1708-1714.
Garg S.K., et al., "Improved Glucose Excursions Using an Implantable Real-Time Continuous Glucose Sensor in Adults With Type 1 Diabetes," Emerging Treatments and Technologies, Diabetes Care, vol. 27 (3), 2004, pp. 734-738.
Geller R.I., et al., "Use of an Immunoisolation Device for Cell Transplantation and Tumor Immunotherapy," Annals of the New York Academy of Science, 1997, vol. 831, pp. 438-451.
Gerritsen M., et al., "Influence of Inflammatory Cells and Serum on the Performance of Implantable Glucose Sensors," Journal of Biomedical Material Research, 2001, vol. 54, pp. 69-75.
Gerritsen M., et al., "Performance of Subcutaneously Implanted Glucose Sensors for Continuous Monitoring," The Netherlands Journal of Medicine, vol. 54, 1999, pp. 167-179.
Gerritsen M., et al., "Problems Associated with Subcutaneously Implanted Glucose Sensors," Diabetes Care, vol. 23 (2), Feb. 2000, pp. 143-145.
Gilligan B.J., et al., "Evaluation of a Subcutaneous Glucose Sensor Out to 3 Months in a Dog Model" Diabetes Care, vol. 17 (8), Aug. 1994, pp. 882-887.
Gilligan B.J., et al., "Feasibility of Continuous Long-Term Glucose Monitoring from a Subcutaneous Glucose Sensor in Humans," Diabetes Technology & Therapeutics, vol. 6 (3), 2004, pp. 378-386.
Godsland I.F., et al., "Maximizing the Success Rate of Minimal Model Insulin Sensivity Measurement in Humans: The Importance of Basal Glucose Levels," The Biochemical Society and the Medical Research Society, Clinical Science, vol. 101, 2001, pp. 1-9.
Gore Preclude®, Pericardial Membrane Brochure, Jun. 2009. W.L. Gore & Associates Inc., Flagstaff, AZ 86004.
Gore Preclude®, Pericardial Membrane Brochure, Nov. 2001, W.L. Gore & Associates Inc., Flagstaff, AZ—86004, 4 pages.
Gouda M.D., et al., "Thermal Inactivation of Glucose Oxidase," The Journal of Biological Chemistry, vol. 278 (27), Issue of Jul. 4, 2003, pp. 24324-24333.
Gough D.A., "The implantable Glucose Sensor: An Example of Bioengineering Design," Introduction to Bioengineering, 2001, Chapter 3, pp. 57-66.
Gough D.A., et al., "Frequency Characterization of Blood Glucose Dynamics," Annals of Biomedical Engineering, vol. 31, 2003, pp. 91-97.
Gough D.A., et al., "Immobilized Glucose Oxidase in Implantable Glucose Sensor Technology," Diabetes Technology & Therapeutics, vol. 2 (3), 2000, pp. 377-380.
Gregg B A., et al., "Cross-Linked Redox Gels Containing Glucose Oxidase for Amperometric Biosensor Applications," Anal Chem, 1990, vol. 62, pp. 258-263.
Gross T.M., et al., "Efficacy and Reliability of the Continuous Glucose Monitoring System," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S19-S26.
Gross T.M., et al., "Performance Evaluation of the Minimed® Continuous Glucose Monitoring System During Patient Home Use," Diabetes Technology & Therapeutics, vol. 2(1), 2000, pp. 49-56.
Gross, et al., "Diabetes Technology & Therapeutics," Letters to the Editor, Diabetes Technology & Therapeutics, vol. 3 (1), 2001, pp. 129-131.
Guerci B., et al., "Clinical Performance of CGMS in Type 1 Diabetic Patients Treated by Continuous Subcutaneous Insulin Infusion Using Insulin Analogs," Diabetes Care, vol. 26, 2003, pp. 582-589.
Guerra S., et al., "Enhancing the Accuracy of Subcutaneous Glucose Sensors: A Real-Time Deconvolution-Based Approach," IEEE Transactions on Biomedical Engineering, vol. 59(6), Jun. 2012, pp. 1658-1669.
Guo M., et al., "Modification of Cellulose Acetate Ultrafiltration Membrane by Gamma Ray Radiation," Shuichuli Jishi Bianji Weiyuanhui, 1998, vol. 23(6), pp. 315-318. (Abstract only).
Hall S.B., et al., "Electrochemical Oxidation of Hydrogen Peroxide at Platinum Electrodes. Part 1. An Adsorption-controlled Mechanism," Electrochimica Acta, vol. 43, Nos. 5/6, 1998, pp. 579-588.
Hall S.B., et al., "Electrochemical Oxidation of Hydrogen Peroxide at Platinum Electrodes. Part II: Effect of potential," Electrochimica Acta, vol. 43 (14-15), 1998, pp. 2015-2024.
Hall S.B., et al., "Electrochemical Oxidation of Hydrogen Peroxide at Platinum Electrodes. Part III: Effect of Temperature," Electrochimica Acta, vol. 44, 1999, pp. 2455-2462.
Hall S.B., et al., "Electrochemical Oxidation of Hydrogen Peroxide at Platinum Electrodes. Part IV: Phosphate Buffer Dependence," Electrochimica Acta, vol. 44, 1999, pp. 4573-4582.
Hall S.B., et al., "Electrochemical Oxidation of Hydrogen Peroxide at Platinum Electrodes. Part V: Inhibition by Chloride," Electrochimica Acta, vol. 45, 2000, pp. 3573-3579.
Halvorsen C., et al., "Vasodilation of Rat Retinal Microvessels Induced by Monobutyrin," Journal of Clinical Investigation, Dec. 1993, vol. 92, pp. 2872-2876.
Hamilton, "Complete Guide to Selecting the Right Hamilton Gastight, Microliter, and Specialty Syringe for your Application," Syringe Selection, www.hamiltoncompany.com, 2006, 20 pages.
Harada, et al., "Long-Term Results of the Clinical Use of an Expanded Polytetrafluoroethylene Surgical Membrane as a Pericardial Substitute," Journal of Thoracic and Cardiovascular Surgery, 1988, vol. 96(5), pp. 811-815.
Harrison, et al., "Characterization of Perfluorosulfonic Acid Polymer Coated Enzyme Electrodes and a Miniaturized Integrated Potentiostat for Glucose Analysis in Whole Blood," Analytical Chemistry, 1988, vol. 60, pp. 2002-2007.
Hashiguchi Y., et al., "Development of a Miniaturized Glucose Monitoring System by Combining a Needle-Type Glucose Sensor with Microdialysis Sampling Method: Long-term subcutaneous tissue glucose monitoring in ambulatory diabetic patients," Diabetes Care, vol. 17, No. 5, May 1994, pp. 387-396.

(56) **References Cited**

OTHER PUBLICATIONS

Heise T., et al., "Hypoglycemia warning signal and glucose sensors: Requirements and concepts," Diabetes Technology & Therapeutics, vol. 5, No. 4, 2003, pp. 563-571.
Heller A., "Electrical Connection of Enzyme Redox Centers to Electrodes," J. Phys. Chem., vol. 96, 1992, pp. 3579-3587.
Heller A., "Electrical Wiring of Redox Enzymes," Ace. Chem. Res., vol. 23, 1990, pp. 128-134.
Heller A., "Implanted Electrochemical Glucose Sensors for the Management of Diabetes," Annu. Rev., Biomed Eng., vol. 1, 1999, pp. 153-175.
Heller A., "Plugging Metal Connectors into Enzymes," Nature Biotechnology, vol. 21, No. 6, Jun. 2003, pp. 631-632.
Heller, et al., "In vivo Glucose Monitoring with Miniature "Wired" Glucose Oxidase Electrodes," Analytical Sciences, 2001, vol. 17 Supplement, pp. i297-i300.
Heydron W H., et al., "A New Look at Pericardial Substitutes", Journal of Thoracic and Cardiovascular Surgery, 1987, vol. 94(2), pp. 291-296.
Hicks J.M., "In Situ Monitoring," Clinical Chemistry, vol. 31 (12), 1985, pp. 1931-1935.
Hitchman M.L., "Measurement of Dissolved Oxygen," Edited by Elving P.J et al., Chemical Analysis, New York, John Wiley & Sons, vol. 49, Chapter 3, 1978, pp. 34-49 and 59-123.
Hoel P.G., "Elementary Statistics," Fourth Edition, John Wiley & Sons, Inc., 1976, pp. 113-114.
Houghton Mifflin Company, "American Heritage Dictionary," 4th Edition, 2000, pp. 82.
Houghton Mifflin Company, "Xenogenic, the American Heritage Stedman's Medical Dictionary," 2002, Answers.Com, retrieved from http://www.answers.com/topic/xenogenic, on Nov. 7, 2006, 2 Pages.
Hrapovic S., et al., "Picoamperometric Detection of Glucose at Ultrasmall Platinum-Based Biosensors Preparation and Characterization," Anal. Chem, vol. 75, 2003, pp. 3308-3315.
Hu Y., et al., "A Needle-Type Enzyme-Based Lactate Sensor for In Vivo Monitoring," Analytica Chimica Acta, vol. 281, 1993, pp. 503-511.
Huang C., et al., "Electrochemical Generation of Oxygen. 1: The Effects of Anions and Cations on Hydrogen Chemisorption and Aniodic Oxide Film Formation on Platinum Electrode. 2: The Effects of Anions and Cations on Oxygen Generation on Platinum Electrode," U.S. Department of Commence/NTIS, 1975, 126 pages.
Huang Q., et al., "A 0.5mW Passive Telemetry IC for Biomedical Applications," Proceedings of the 23rd European Solid-State Circuits Conference (ESSCIRC '97), Southampton, UK, Sep. 16-18, 1997, pp. 172-175.
Hunter I., et al., "Minimally Invasive Glucose Sensor and Insulin Delivery System," MIT Home Automation and Healthcare Consortium, Mar. 31, 2000, Progress Report No. 25, 17 pages.
International Preliminary Examination Report for Application No. PCT/US2002/023903 dated Apr. 18, 2005, 5 pages.
International Preliminary Report on Patentability for Application No. PCT/US2004/023455 dated Jan. 23, 2006, 7 pages.
International Preliminary Report on Patentability for Application No. PCT/US2004/040476 dated Dec. 8, 2006, 4 pages.
International Preliminary Report on Patentability for Application No. PCT/US2005/014696 dated Nov. 7, 2006, 6 pages.
International Preliminary Report on Patentability for Application No. PCT/US2005/024993 dated Jan. 16, 2007, 7 pages.
International Preliminary Report on Patentability for Application No. PCT/US2005/024994 dated Jan. 16, 2007, 8 pages.
International Preliminary Report on Patentability for Application No. PCT/US2006/006574 dated Aug. 26, 2008, 10 pages.
International Preliminary Report on Patentability for Application No. PCT/US2006/008616 dated Mar. 3, 2009, 7 pages.
International Preliminary Report on Patentability for Application No. PCT/US2006/024132 dated Dec. 24, 2007, 6 pages.
International Preliminary Report on Patentability for Application No. PCT/US2006/031496 dated Nov. 27, 2008, 8 pages.
International Preliminary Report on Patentability for Application No. PCT/US2007/005422 dated Sep. 1, 2009, 5 pages.
International Preliminary Report on Patentability for Application No. PCT/US2008/058158 dated Sep. 29, 2009, 9 pages.
International Preliminary Report on Patentability for Application No. PCT/US2009/040285, dated Oct. 21, 2011, 8 pages.
International Search Report and Written Opinion for Application No. PCT/US2004/023455 dated Dec. 23, 2004, 8 pages.
International Search Report and Written Opinion for Application No. PCT/US2004/040476 dated Nov. 16, 2006, 4 pages.
International Search Report and Written Opinion for Application No. PCT/US2005/014696 dated Jun. 29, 2006, 8 pages.
International Search Report and Written Opinion for Application No. PCT/US2005/024993, dated Nov. 4, 2005, 12 pages.
International Search Report and Written Opinion for Application No. PCT/US2005/024994 dated Nov. 15, 2005, 11 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/001998 dated Jul. 25, 2006, 5 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/006574 dated Aug. 4, 2006, 13 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/008616 dated Mar. 13, 2008, 9 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/019889 dated Feb. 20, 2007, 6 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/024132 dated Jul. 20, 2007, 6 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/031496 dated Sep. 20, 2007, 9 pages.
International Search Report and Written Opinion for Application No. PCT/US2006/038820 dated Jun. 20, 2007, 7 pages.
International Search Report and Written Opinion for Application No. PCT/US2007/005422 dated May 20, 2008, 5 pages.
International Search Report and Written Opinion for Application No. PCT/US2008/058158 dated Aug. 8, 2008, 10 pages.
International Search Report and Written Opinion for Application No. PCT/US2009/040285, dated May 29, 2009, 8 pages.
International Search Report for Application No. PCT/US2001/023850 dated Jan. 16, 2002, 3 pages.
International Search Report for Application No. PCT/US2002/023903 dated Feb. 27, 2003, 4 pages.
International Search Report for Application No. PCT/US2003/015816 dated Sep. 22, 2003, 4 pages.
Isermann R., "Supervision, Fault-Detection and Fault-Diagnosis Methods—An Introduction," Control Engineering Practice, vol. 5 (5), 1997, pp. 639-652.
Isermann R., et al., "Trends in the Application of Model-Based Fault Detection and Diagnosis of Technical Processes", Control Engineering Practice, vol. 5 (5), 1997, pp. 709-719.
Ishikawa M., et al., "Initial Evaluation of a 290-Mm Diameter Subcutaneous Glucose Sensor: Glucose Monitoring With a Biocompatible, Flexible-Wire, Enzyme-Based Amperometric Microsensor in Diabetic and Nondiabetic Humans," Journal of Diabetes and Its Complications, vol. 12, 1998, pp. 295-301.
Jablecki M., et al., "Simulations of the Frequency Response of Implantable Glucose Sensors," Analytical Chemistry, vol. 72, 2000, 1853-1859.
Jaffari S.A., et al., "Recent Advances in Amperometric Glucose Biosensors for In Vivo Monitoring," Physiological Measurement, 1995, vol. 16, pp. 1-15.
Jaremko J., et al., "Advances Toward the Implantable Artificial Pancreas for Treatment of Diabetes," Diabetes Care, vol. 21 (3), Mar. 1998, pp. 444-450.
Jensen M.B., et al., "Fast Wave Forms for Pulsed Electrochemical Detection of Glucose by Incorporation of Reductive Desorption of Oxidation Products," Analytical Chemistry, vol. 69 (9), May 1997, pp. 1776-1781.
Jeong R.A., et al., "In Vivo Calibration of the Subcutaneous Amperometric Glucose Sensors Using a Non-Enzyme Electrode," Biosensors and Bioelectronics, Elsevier, vol. 19, 2003, pp. 313-319.
Jeutter D.C., "A Transcutaneous Implanted Battery Recharging and Biotelemeter Power Switching System," IEEE Transactions on Biomedical Engineering, vol. BME-29 (5), May 1982, pp. 314-321.

(56) **References Cited**

OTHER PUBLICATIONS

Jeutter D.C., et al., "Design of a Radio-Linked Implantable Cochlear Prosthesis Using Surface Acoustic Wave Devices," IEEE Transactions on Ultrasonics, Ferroelectrics and Frequency Control, vol. 40 (5), Sep. 1993, pp. 469-477.

Jobst G., et al., "Thin-Film Microbiosensors for Glucose-Lactate Monitoring," Anal Chem, Sep. 15, 1996, vol. 68(18), pp. 3173-3179.

Johnson K.W., "Reproducible Electrodeposition of Biomolecules for the Fabrication of Miniature Electroenzymatic Biosensors," Sensors and Actuators B, vol. 5, 1991, pp. 85-89.

Johnson K.W., et al., "In Vivo Evaluation of an Electroenzymatic Glucose Sensor Implanted in Subcutaneous Tissue," Biosensors and Bioelectronics, 1992, vol. 7, pp. 709-714.

Johnson P.C., "Peripheral Circulation," John Wiley & Sons, 1978, p. 198 (5 pages).

Johnson R C., et al., "Abstract: Neovascularization of Cell Transplantation Devices: Role of Membrane Architecture and Encapsulated Tissue," Abstracts of Papers, American Chemical Society, Sep. 7-11, 1997, 214th ACS National Meeting, Part 2, 305-PMSE, 2 pages.

Joung G.B., et al., "An Energy Transmission System for an Artificial Heart Using Leakage Inductance Compensation of Transcutaneous Transformer," IEEE Transactions on Power Electronics, vol. 13 (6), Nov. 1998, pp. 1013-1022.

Jovanovic L.M.D., "The Role of Continuous Glucose Monitoring in Gestational Diabetes Mellitus," Diabetes Technology and Therapeutics, vol. 2 (1), 2000, pp. S67-S71.

Jungheim K., et al., "How Rapid Does Glucose Concentration Change in Daily Life of Patients with Type 1 Diabetes?," Diabetologia, 2002, vol. 45, pp. A250-A276.

Jungheim K., et al., "Risky Delay of Hypoglycemia Detection by Glucose Monitoring at the Arm," Diabetes Care, vol. 24 (7), Jul. 2001, pp. 1303-1304.

Kacaniklic V., et al., "Amperometric Biosensors for Detection of L- and D-Amino Acids Based on Coimmoblized Peroxidase and L- and D-Amino Acid Oxidases in Carbon Paste Electrodes," Electroanalysis, vol. 6, May-Jun. 1994, pp. 381-390.

Kamath A., et al., "Calibration of a Continuous Glucose Monitor: Effect of Glucose Rate of Change," Eighth Annual Diabetes Technology Meeting, Nov. 13-15, 2008, pp. A88.

Kamath, et al., "Analysis of Time Lags and Other Sources of Error of the DexCom SEVEN Continuous Glucose Monitor," Diabetics Technology and Therapeutic, Nov. 2009, vol. 11, No. 11, pp. 689-695.

Kang S.K., et al., "In Vitro and Short-Term in Vivo Characteristics of a Kel-F Thin Film Modified Glucose Sensor," Analytical Sciences, vol. 19, Nov. 2003, pp. 1481-1486.

Kaplan S.M., "Wiley Electrical and Electronics Engineering Dictionary," IEEE Press, John Wiley & Sons, Inc., 2004, pp. 141, 142, 548 & 549.

Kargol M., et al., "Studies on the Structural Properties of Porous Membranes: Measurement of Linear Dimensions of Solutes," Biophysical Chemistry, 2001, vol. 91, pp. 263-271.

Karube I., et al., "Microbiosensors for Acetylcholine and Glucose," Biosensors & Bioelectronics, 1993, vol. 8, pp. 219-228.

Kaufman F.R., "Role of the Continuous Glucose Monitoring System in Pediatric Patients," Diabetes Technology and Therapeutics, vol. 2 (1), 2000, S49-S52.

Kaufman F.R., et al., "A Pilot Study of the Continuous Glucose Monitoring System," Diabetes Care, vol. 24 (12), Dec. 2001, pp. 2030-2034.

Kawagoe J.L., et al., "Enzyme-Modified Organic Conducting Salt Microelectrode," Analytical Chemistry, vol. 63, 1991, pp. 2961-2965.

Keedy F.H., et al., "Determination of Urate in Undiluted Whole Blood by Enzyme Electrode," Biosensors and Bioelectronics, vol. 6, 1991, pp. 491-499.

Kerner W., "Implantable Glucose Sensors: Present Status and Future Developments," Experimental and Clinical Endocrinol Diabetes, vol. 109 (2), 2001, pp. S341-S346.

Kerner W., et al., "The Function of a Hydrogen Peroxide-Detecting Electroenzymatic Glucose Electrode is Markedly Impaired in Human Sub-Cutaneous Tissue and Plasma," Biosensors and Bioelectronics, vol. 8, 1993, pp. 473-482.

Kerner, et al., "A Potentially Implantable Enzyme Electrode for Amperometric Measurement of Glucose," Hormone and Metabolic Research Supplement, vol. 20, 1988, pp. 8-13.

Kidd K R., et al., "Angiogenesis and Neovascularization Associated with Extracellular Matrix Modified Porous Implants," Journal of Biomedical Materials Research, 2001, vol. 59(2), pp. 366-377.

Kiechle F.L., "The Impact of Continuous Glucose Monitoring on Hospital Point-of-Care Testing Programs," Diabetes Technology and Therapeutics, vol. 3 (4), 2001, pp. 647-650.

Klonoff D., et al., "Performance Metrics for Continuous Interstitial Glucose Monitoring; Approved Guideline," Clinical and Laboratory Standards Institute, POCT05-A, vol. 28 (33), 2008, 72 pages.

Klueh U., et al., "Inflammation and Glucose Sensors: Use of Dexamethasone to Extend Glucose Sensor Function and Life Span in Vivo," Journal of Diabetes Science and Technology, vol. 1 (4), Jul. 2007, pp. 496-504.

Klueh U., et al., "Use of Vascular Endothelial Cell Growth Factor Gene Transfer to Enhance Implantable Sensor Function in Vivo," Biosensor Function and VEGF-Gene Transfer, vol. 67 (4), 2003, pp. 1072-1086.

Kondo T., et al., "A Miniature Glucose Sensor, Implantable in the Blood Stream," Diabetes Care, vol. 5 (3), May-Jun. 1982, 218-221.

Koschinsky T., et al., "New Approach to Technical and Clinical Evaluation of Devices for Self-Monitoring of Blood Glucose," Diabetes Care, vol. 11 (8), Sep. 1988, pp. 619-629.

Koschinsky T., et al., "Sensors for Glucose Monitoring: Technical and Clinical Aspects," Diabetes Metabolism Research and Reviews, vol. 17, 2001, pp. 113-123.

Kost J., et al., "Glucose-Sensitive Membranes Containing Glucose Oxidase: Activity, Swelling, and Permeability Studies," Journal of Biomedical Materials Research, vol. 19, 1985, pp. 1117-1133.

Koudelka M., et al., "In Vivo Response of Microfabricated Glucose Sensors to Glycemia Changes in Normal Rats," Biomed. Biochim. Acta, vol. 48 (11/12), Nov.-Dec. 1989, pp. 953-956.

Koudelka M., et al., "In-Vivo Behaviour of Hypodermically Implanted Microfabricated Glucose Sensors," Biosensors and Bioelectronics, vol. 6, 1991, pp. 31-36.

Kovatchev B.P., et al., "Evaluating the Accuracy of Continuous Glucose-Monitoring Sensors: Continuous Glucose-Error Grid Analysis Illustrated by TheraSense Freestyle Navigator Data," Diabetes Care, vol. 27 (8), Aug. 2004, pp. 1922-1928.

Kraver., et al., "A Mixed-Signal Sensor Interface Microinstrument," Sensors and Actuators A, Physical 2001, vol. 91, pp. 266-277.

Krouwer J.S., "Setting Performance Goals and Evaluating Total Analytical Error for Diagnostic Assays," Clinical Chemistry, vol. 48 (6), 2002, pp. 919-927.

Kruger D., et al., "Psychological Motivation and Patient Education: A Role for Continuous Glucose Monitoring," Diabetes Technology and Therapeutics, vol. 2 (1), 2000, pp. S93-S97.

Kugler J D., et al., "A New Steroid-Eluting Epicardial Lead: Experience with Atrial and Ventricular Implantation in the Immature Swine," PACE, Aug. 1990, vol. 13, pp. 976-981.

Kulys J., et al., "Carbon-Paste Biosensors Array for Long-Term Glucose Measurement," Biosensors & Bioelectronics, vol. 9, 1994, pp. 491-500.

Kunjan K., et al., "Automated Blood Sampling and Glucose Sensing in Critical Care Settings," Journal of Diabetes Science and Technology, vol. 2 (2), Mar. 2008, pp. 194-200.

Kunzler J F., et al., "Contact Lens Materials," Chemistry & Industry, Aug. 21, 1995, pp. 651-655.

Kunzler J., et al.," Hydrogels based on Hydrophilic Side Chain Siloxanes," Poly Mat Sci and Eng, 1993, vol. 69, pp. 226-227.

Kurnik R.T., et al., "Application of the Mixtures of Experts Algorithm for Signal Processing in a Noninvasive Glucose Monitoring System," Sensors and Actuators B, vol. 60, 1999, pp. 19-26.

(56) **References Cited**

OTHER PUBLICATIONS

Kurtz T.W., et al., "Recommendations for Blood Pressure Measurement in Humans and Experimental Animals, Part 2: Blood Pressure Measurement in Experimental Animals: A Statement for Professionals From the Subcommittee of Professional and Public Education of the American Heart Association Council on High Blood Pressure Research," Hypertension, Feb. 2005, vol. 45, pp. 299-310.
Kusano H., "Glucose Enzyme Electrode with Percutaneous Interface which Operates Independently of Dissolved Oxygen," Clinical Physics and Physiological Measurement, 1989, vol. 10, No. 1, pp. 1-9.
Lacourse W.R., et al., "Optimization of Waveforms for Pulsed Amperometric Detection of Carbohydrates Based on Pulsed Voltammetry," Analytical Chemistry, vol. 65, 1993, pp. 50-52.
Ladd M.F.C., et al., "Structure Determination by X-Ray Crystallography," 3rd Edition, Plenum Press, 1994, Ch. 1, pp. xxi-xxiv and 1-58.
Lee E., et al., "Effects of Pore Size, Void Volume, and Pore Connectivity on Tissue Responses to Porous Silicone Implants," Society for Biomaterials, 25th Annual Meeting, 1999, p. 171.
Lehmann E.D., et al., Retrospective Validation of a Physiological Model of Glucose-Insulin Interaction in Type 1 Diabetes Mellitus. Medical Engineering & Physics, vol. 16, May 1994, pp. 193-202.
Leprince P., et al., "Expanded Polytetrafluoroethylene Membranes to Wrap Surfaces of Circulatory Support Devices in Patients Undergoing Bridge to Heart Transplantation," European Journal of Cardiothoracic Surgery, 2001, vol. 19, pp. 302-306.
Lerner., et al., "An Implantable Electrochemical Glucose Sensor," Ann. N. Y. Acad. Sci., vol. 428, May 1984, pp. 263-278.
Lewandowski J.J., et al., "Evaluation of a Miniature Blood Glucose Sensor," Transactions—American Society for Artificial Internal Organs, vol. 34, 1988, pp. 255-258.
Leypoldt J.K., et al., "Model of a Two-Substrate Enzyme Electrode for Glucose," Analytical Chemistry, vol. 56, 1984, pp. 2896-2904.
Linke B., et al., "Amperometric Biosensor for In Vivo Glucose Sensing Based on Glucose Oxidase Immobilized in a Redox Hydrogel," Biosensors and Bioelectronics, vol. 9, 1994, pp. 151-158.
Loebe M., et al., "Use of Polytetrafluoroethylene Surgical Membranes as a Pericardial Substitute," PTFE Membrane in Correction of Congenital Heart Defects, Texas Heart Institute Journal, 1993, vol. 20, No. 3, pp. 213-217.
Loffler P., et al., "Separation and Determination of Traces of Ammonia in Air by Means of Chromatomembrane Cells," Fresenius Journal of Analytical Chemistry, 1995, vol. 352, pp. 613-614.
Lohn A., et al., "A Knowledge-Based System for Real-Time Validation of Calibrations and Measurements," Chemometrics and Intelligent Laboratory Systems, vol. 46, 1999, pp. 57-66.
Lortz J., et al., "What is Bluetooth? We Explain the Newest Short-Range Connectivity Technology," In Smart Computing Learning Series, Wireless Computing, vol. 8 (5), 2002, pp. 72-74.
Lowe C.R., "Biosensors," Trends in Biotechnology, vol. 2 (3), 1984, pp. 59-65.
Luong J.H.T., et al., "Solubilization of Multiwall Carbon Nanotubes by 3-Aminopropyltriethoxysilane towards the Fabrication of Electrochemical Biosensors with Promoted Electron Transfer," Electroanalysis, vol. 16 (1-2), 2004, pp. 132-139.
Lyandres O., et al. "Progress toward an In Vivo Surface-Enhanced Raman Spectroscopy Glucose Sensor," Diabetes Technology and Therapeutics, vol. 10 (4), 2008, pp. 257-265.
Lyman D J., "Polyurethanes. I. The Solution Polymerization of Diisocyanates with Ethylene Glycol," Journal of Polymer Science, 1960, vol. XLV, pp. 45-49.
Lynch S.M., et al., "Estimation-Based Model Predictive Control of Blood Glucose in Type I Diabetics: A Simulation Study," Proceedings of the IEEE 27th Annual Northeast Bioengineering Conference, 2001, pp. 79-80.
Lynn P.A., "Recursive Digital Filters for Biological Signals," Med. & Biol. Engineering, vol. 9, 1971, pp. 37-43.
Madaras M B., et al., "Microfabricated Amperometric Creatine and Creatinine Biosensors," Analytica Chimica Acta, 1996, vol. 319, pp. 335-345.
Maidan R., et al., "Elimination of Electrooxidizable Interferent-Produced Currents in Amperometric Biosensors," Analytical Chemistry, vol. 64, 1992, pp. 2889-2896.
Makale M.T., et al., "Tissue Window Chamber System for Validation of Implanted Oxygen Sensors," American Journal of Physiology-Heart and Circulatory Physiology, vol. 284, Feb. 21, 2003, pp. 1-27.
Malin S.F., et al., "Noninvasive Prediction of Glucose by Near-Infrared Diffuse Reflectance Spectroscopy," Clinical Chemistry, vol. 45 (9), 1999, pp. 1651-1658.
Mancy K.H., et al., "A Galvanic Cell Oxygen Analyzer," Journal of Electroanalytical Chemistry, vol. 4, 1962, pp. 65-92.
Maran A., et al., "Continuous Subcutaneous Glucose Monitoring in Diabetic Patients," A Multicenter Analysis, Diabetes Care, vol. 25 (2), Feb. 2002, pp. 347-352.
March W.F., "Dealing with the Delay," Diabetes Technology & Therapeutics, vol. 4 (1), 2002, pp. 49-50.
Marena S., et al., "The Artificial Endocrine Pancreas in Clinical Practice and Research," Panminerva Medica, vol. 35 (2), 1993, pp. 67-74.
Martin R.F., "General Deming Regression for Estimating Systematic Bias and its Confidence Interval in Method-Comparison Studies," Clinical Chemistry, vol. 46 (1), 2000, pp. 100-104.
Mascini M., et al., "Glucose Electrochemical Probe with Extended Linearity for Whole Blood," Journal Pharmaceutical and Biomedical Analysis, vol. 7 (12), 1989, pp. 1507-1512.
Mastrototaro J.J., "The MiniMed Continuous Glucose Monitoring System," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S13-S18.
Mastrototaro J.J., et al., "An Electroenzymatic Glucose Sensor Fabricated on a Flexible Substrate," Sensors and Actuators B, vol. 5, 1991, pp. 139-144.
Mastrototaro J.J., et al., "Reproducibility of the Continuous Glucose Monitoring System Matches Previous Reports and the Intended Use of the Product," Diabetes Care, vol. 26 (1), Jan. 2003, pp. 256-257.
Mathivanar R., et al., "In Vivo Elution Rate of Drug Eluting Ceramic Leads with a Reduced Dose of Dexamethasone Sodium Phosphate," PACE, vol. 13, Part II, Dec. 1990, pp. 1883-1886.
Matsuki H., "Energy Transfer System Utilizing Amorphous Wires for Implantable Medical Devices," IEEE Transactions on Magnetics, vol. 31 (2), 1994, pp. 1276-1282.
Matsumoto T., et al., "A Micro-Planar Amperometric Glucose Sensor Unsusceptible to Interference Species," Sensors and Actuators B, 49, 1998, pp. 68-72.
Matthews D.R., et al., "An Amperometric Needle-Type Glucose Sensor Testing in Rats and Man," Diabetic Medicine, vol. 5, 1988, pp. 248-252.
Mazze R.S., et al., "Characterizing Glucose Exposure for Individuals with Normal Glucose Tolerance Using Continuous Glucose Monitoring and Ambulatory Glucose Profile Analysis," Diabetes Technology & Therapeutics, vol. 10 (3), 2008, pp. 149-159.
Mazzola F., et al., "Video Diabetes: A Teaching Tool for Children with Insulin-Dependent Diabetes," IEEE, Proceedings 7th Annual Symposium on Computer Applications in Medical Care, Oct. 1983, 1 page Abstract.
McCartney L.J., et al., "Near-Infrared Fluorescence Lifetime Assay for Serum Glucose Based on Allophycocyanin-Labeled Concanavalin A," Analytical Biochemistry, vol. 292, 2001, pp. 216-221.
McGarraugh G., et al., "Glucose Measurements Using Blood Extracted From the Forearm and the Finger," TheraSense, Inc., 2001, pp. 1-8.
McGarraugh G., et al., "Physiological Influences on Off-Finger Glucose Testing", Diabetes Technology & Therapeutics, vol. 3 (3), 2001, pp. 367-376.
McGrath M.J., et al., "The Use of Differential Measurements with a Glucose Biosensor for Interference Compensation During Glucose Determinations by Flow Injection Analysis," Biosens Bioelectron, vol. 10, 1995, pp. 937-943.
McKean B.D., et al., "A Telemetry Instrumentation System for Chronically Implanted Glucose and Oxygen Sensors," IEEE Transactions on Biomedical Engineering, vol. 35 (7), Jul. 1988, pp. 526-532.

(56) **References Cited**

OTHER PUBLICATIONS

Memoli a., et al., "A Comparison between Different Immobilised Glucoseoxidase-Based Electrodes," Journal of Pharmaceutical and Biomedical Analysis, vol. 29, 2002, pp. 1045-1052.
Merriam Webster Online Dictionary, Definition for "Aberrant," retrieved from https://www.merriam-webster.com/dictionary/aberrant, Aug. 19, 2008, 1 page.
Merriam-Webster Online Dictionary, Definition of "Acceleration" retrieved from http://www.merriam-webster.com/dictionary/Acceleration, Jan. 11, 2010, 1 page.
Merriam-Webster Online Dictionary, Definition of "Nominal" retrieved from http://www.merriam-webster.com/dictionary/nominal, Apr. 23, 2007, 1 page.
Merriam-Webster Online Dictionary, Definition of "System". http://www.merriamwebster.com/dictionary/System, Jan. 11, 2010, 2 pages.
Metzger M., et al., "Reproducibility of Glucose Measurements using the Glucose Sensor," Diabetes Care, vol. 25 (6), Jul. 2002, pp. 1185-1191.
Meyerhoff C., et al., "On Line Continuous Monitoring of Subcutaneous Tissue Glucose in Men by Combining Portable Glucosensor With Microdialysis," Diabetologia, vol. 35 (11), 1992, pp. 1087-1092.
Mid-West Innovators, Inc., "Durometer Product Description," 2014, 1 page.
Miller J.A., et al., "Development of an Autotuned Transcutaneous Energy Transfer System," ASAIO Journal, vol. 39, 1993, pp. M706-M710.
Miller K.M., et al., "Generation of IL-1 like Activity in Response to Biomedical Polymer Implants: a Comparison of in Vitro and in Vivo Models," Journal of Biomedical Materials Research, vol. 23(9), 1989, pp. 1007-1026.
Miller K.M., et al., "Human monocyte/macrophage activation and interleukin 1 generation by biomedical polymers," Journal of Biomedical Materials Research, vol. 22 (8), 1988, pp. 713-731.
Miller K.M., et al., "In Vitro Stimulation of Fibroblast Activity by Factors Generated from Human Monocytes Activated by Biomedical Polymers," Journal of Biomedical Materials Research, vol. 23(8), 1989, pp. 911-930.
Minale C., et al., "Clinical Experience with Expanded Polytetrafluoroethylene Gore-Tex Surgical Membrane for Pericardial Closer: A Study of 110 Cases," Journal of Cardiac Surgery, vol. 3, Sep. 1988, pp. 193-201.
Moatti-Sirat D., et al., "Evaluating In Vitro and In Vivo the Interference of Ascorbate and Acetaminophen on Glucose Detection by a Needle-Type Glucose Sensor," Biosensors and Bioelectronics, vol. 7, 1992, pp. 345-352.
Moatti-Sirat D., et al., "Reduction of Acetaminophen Interference in Glucose Sensors by a Composite Nafion Membrane: Demonstration in Rats and Man," Diabetologia, vol. 37 (6), Jun. 1994, pp. 610-616.
Moatti-Sirat., et al., "Towards Continuous Glucose Monitoring: In Vivo Evaluation of a Miniaturized Glucose Sensor Implanted for Several Days in Rat Subcutaneous Tissue," Diabetologia, vol. 35, 1992, pp. 224-230.
Mond H.G., et al., "The Electrode-Tissue Interface: The Revolutionary Role of Steroid Elution," PACE, vol. 15, Jan. 1992, pp. 95-107.
Monsod T.P., et al., "Do Sensor Glucose Levels Accurately Predict Plasma Glucose Concentrations During Hypoglycemia and Hyperinsulinemia? ,"Diabetes Care, vol. 25 (5), 2002, pp. 889-893.
Morff R.J., et al., "Microfabrication of Reproducible, Economical, Electroenzymatic Glucose Sensors," Annual International Conference of the IEEE Engineering in Medicine and Biology Society, vol. 12 (2), 1990, pp. 0483-0484.
Mosbach K., et al., "Determination of Heat Changes in the Proximity of Immobilized Enzymes with an Enzyme Thermistor and its Use for the Assay of Metabolites," Biochimica Biophysica Acta, vol. 403, 1975, pp. 256-265.
Motonaka J., et al., "Determination of Cholesterol and Cholesterol Ester with Novel enzyme Microsensors," Anal. Chem., vol. 65, 1993, pp. 3258-3261.
Moussy F., "Implantable Glucose Sensor: Progress and Problems," IEEE, Nov. 2002, pp. 270-273.
Moussy F., et al., "A Miniaturized Nafion-Based Glucose Sensor: In Vitro and In Vivo Evaluation in Dogs," International Journals of Artificial Organs, vol. 17 (2), 1994, pp. 88-94.
Moussy F., et al., "Biomaterials community examines biosensor biocompatibility," Diabetes Technology & Therapeutics, vol. 2(3), 2000, pp. 473-477.
Moussy F., et al., "Performance of Subcutaneously Implanted Needle-Type Glucose Sensors Employing a Novel Trilayer Coating," Analytical Chemistry, vol. 65, Aug. 1, 1993, pp. 2072-2077.
Mowery K.A., et al., "Preparation and Characterization by Hydrophobic Polymeric Films that are Thromboresistant via Nitric Oxide Release," Biomaterials, vol. 21, 2000, pp. 9-21.
Murphy S.M., et al., "Polymer Membranes in Clinical Sensor Applications, II. The Design and Fabrication of Permselective Hydrogels for Electrochemical Devices," Biomaterials, 1992, vol. 13 (14), pp. 979-990.
Muslu, "Trickling Filter Performance," Applied Biochemistry and Biotechnology, vol. 37, 1992, pp. 211-224.
Myler S., et al., "Ultra-Thin-Polysiloxane-Film-Composite Membranes for the Optimisation of Amperometric Oxidase Enzyme Electrodes," Biosensors & Bioelectronics, vol. 17, 2002, pp. 35-43.
Nakayama Y., et al., "Surface Fixation of Hydrogels: Heparin and Glucose Oxidase Hydrogelated Surfaces" ASAIO Journal, 1992, pp. M421-M424.
Nam Y.S., et al., "A Novel Fabrication Method of Macroporous Biodegradable Polymer Scaffolds Using Gas Foaming Salt as a Porogen Additive," J Biomed Mater Res, 2000, vol. 53, pp. 1-7.
Neuburger G.G., et al., "Pulsed Amperometric Detection of Carbohydrates at Gold Electrodes with a Two-Step Potential Waveform," Anal. Chem., vol. 59, 1987, pp. 150-154.
Nintendo Healthcare, Wired, Dec. 1993, 1 page.
Notice of Allowance for U.S. Appl. No. 12/422,223, dated Jun. 22, 2012, 8 pages.
Novo Nordisk Pharmaceuticals Inc., "Diabetes Educational Video Game Recognized by Software Publishers Association," Press Release, Mar. 14, 1994, 4 pages.
Office Action for European Application No. 10193214.3 dated May 2, 2013 and Applicant Response filed Aug. 28, 2013, 12 pages.
Office Action for Japanese Application No. 10-538680, dated Nov. 20, 2007, 13 pages.
Office Action for U.S. Appl. No. 08/811,473, dated Dec. 7, 1998, 5 pages.
Office Action for U.S. Appl. No. 09/916,386, dated Apr. 9, 2003, 8 pages.
Office Action for U.S. Appl. No. 10/657,843, dated Sep. 21, 2004, 8 pages.
Office Action for U.S. Appl. No. 10/768,889, dated Feb. 4, 2009, 9 pages.
Office Action for U.S. Appl. No. 10/991,966, dated Jul. 22, 2008, 12 pages.
Office Action for U.S. Appl. No. 10/991,966, dated Nov. 28, 2007, 13 pages.
Office Action for U.S. Appl. No. 11/007,635, dated Jan. 27, 2006, 9 pages.
Office Action for U.S. Appl. No. 11/007,920, dated Jun. 24, 2008, 10 pages.
Office Action for U.S. Appl. No. 11/021,046, dated Aug. 19, 2009, 6 pages.
Office Action for U.S. Appl. No. 11/021,046, dated Dec. 26, 2007, 6 pages.
Office Action for U.S. Appl. No. 11/021,046, dated Feb. 4, 2009, 7 pages.
Office Action for U.S. Appl. No. 11/021,046, dated Jun. 23, 2008, 6 pages.
Office Action for U.S. Appl. No. 11/021,162, dated Jun. 19, 2008, 8 pages.
Office Action for U.S. Appl. No. 11/034,343, dated Dec. 30, 2008, 4 pages.
Office Action for U.S. Appl. No. 11/034,343, dated Jul. 10, 2008, 6 pages.

(56) **References Cited**

OTHER PUBLICATIONS

Office Action for U.S. Appl. No. 11/034,343, dated Nov. 1, 2007, 5 pages.
Office Action for U.S. Appl. No. 11/034,344, dated Jan. 15, 2008, 5 pages.
Office Action for U.S. Appl. No. 11/038,340, dated Feb. 2, 2010, 18 pages.
Office Action for U.S. Appl. No. 11/038,340, dated Jan. 5, 2009, 13 pages.
Office Action for U.S. Appl. No. 11/038,340, dated Jun. 7, 2010, 18 pages.
Office Action for U.S. Appl. No. 11/038,340, dated Jun. 17, 2008, 11 pages.
Office Action for U.S. Appl. No. 11/038,340, dated May 19, 2009, 14 pages.
Office Action for U.S. Appl. No. 11/038,340, dated Nov. 9, 2009, 16 pages.
Office Action for U.S. Appl. No. 11/055,779, dated May 23, 2007, 13 pages.
Office Action for U.S. Appl. No. 11/055,779, dated Oct. 24, 2007, 15 pages.
Office Action for U.S. Appl. No. 11/077,759, dated Jul. 10, 2008, 10 pages.
Office Action for U.S. Appl. No. 11/077,759, dated Mar. 31, 2008, 16 pages.
Office Action for U.S. Appl. No. 11/077,759, dated May 17, 2007, 13 pages.
Office Action for U.S. Appl. No. 11/077,759, dated May 26, 2009, 8 pages.
Office Action for U.S. Appl. No. 11/415,631, dated Mar. 4, 2010, 6 pages.
Office Action for U.S. Appl. No. 11/416,058, dated Mar. 5, 2010, 6 pages.
Office Action for U.S. Appl. No. 11/416,346, dated Mar. 4, 2010, 7 pages.
Office Action for U.S. Appl. No. 11/416,734, dated May 17, 2010, 10 pages.
Office Action for U.S. Appl. No. 11/416,734, dated Oct. 14, 2009, 11 pages.
Office Action for U.S. Appl. No. 11/416,825, dated Oct. 22, 2009, 11 pages.
Office Action for U.S. Appl. No. 11/543,539, dated Dec. 12, 2007, 8 pages.
Office Action for U.S. Appl. No. 11/543,539, dated May 23, 2007, 6 pages.
Office Action for U.S. Appl. No. 11/543,683, dated Dec. 12, 2007, 8 pages.
Office Action for U.S. Appl. No. 11/543,683, dated May 18, 2007, 7 pages.
Office Action for U.S. Appl. No. 11/543,707, dated Dec. 12, 2007, 8 pages.
Office Action for U.S. Appl. No. 11/543,707, dated May 18, 2007, 6 pages.
Office Action for U.S. Appl. No. 11/543,734, dated Dec. 17, 2007, 11 pages.
Office Action for U.S. Appl. No. 11/543,734, dated Jun. 5, 2007, 7 pages.
Office Action for U.S. Appl. No. 11/546,157, dated Feb. 26, 2010, 6 pages.
Office Action for U.S. Appl. No. 11/691,424, dated Dec. 8, 2009, 10 pages.
Office Action for U.S. Appl. No. 11/691,424, dated Jun. 11, 2009, 21 pages.
Office Action for U.S. Appl. No. 11/691,424, dated Nov. 12, 2009, 18 pages.
Office Action for U.S. Appl. No. 11/691,424, dated Sep. 25, 2008, 15 pages.
Office Action for U.S. Appl. No. 11/691,432, dated Feb. 18, 2010, 11 pages.
Office Action for U.S. Appl. No. 11/691,432, dated Jun. 10, 2009, 17 pages.
Office Action for U.S. Appl. No. 11/691,432, dated Nov. 30, 2009, 15 pages.
Office Action for U.S. Appl. No. 11/691,432, dated Sep. 19, 2008, 11 pages.
Office Action for U.S. Appl. No. 11/691,466, dated Oct. 3, 2008, 15 pages.
Office Action for U.S. Appl. No. 11/692,154, dated Jan. 22, 2009, 9 pages.
Office Action for U.S. Appl. No. 11/692,154, dated Jul. 8, 2009, 6 pages.
Office Action for U.S. Appl. No. 12/037,812, dated Apr. 1, 2009, 5 pages.
Office Action for U.S. Appl. No. 12/037,812, dated Jul. 24, 2009, 6 pages.
Office Action for U.S. Appl. No. 12/037,812, dated Sep. 29, 2008, 7 pages.
Office Action for U.S. Appl. No. 12/037,830, dated Aug. 7, 2009, 7 pages.
Office Action for U.S. Appl. No. 12/037,830, dated Feb. 23, 2010, 7 pages.
Office Action for U.S. Appl. No. 12/037,830, dated Feb. 26, 2009, 7 pages.
Office Action for U.S. Appl. No. 12/037,830, dated Sep. 29, 2008, 6 pages.
Office Action for U.S. Appl. No. 12/052,489, dated Aug. 25, 2010, 12 pages.
Office Action for U.S. Appl. No. 12/055,098, dated Oct. 5, 2010, 12 pages.
Office Action for U.S. Appl. No. 12/098,359, dated Jul. 7, 2010, 18 pages.
Office Action for U.S. Appl. No. 12/102,654, dated Jul. 30, 2009, 9 pages.
Office Action for U.S. Appl. No. 12/102,654, dated Mar. 10, 2010, 6 pages.
Office Action for U.S. Appl. No. 12/102,729, dated Jul. 7, 2009, 7 pages.
Office Action for U.S. Appl. No. 12/102,745, dated Dec. 23, 2008, 4 pages.
Office Action for U.S. Appl. No. 12/111,062, dated Mar. 3, 2010, 8 pages.
Office Action for U.S. Appl. No. 12/113,508, dated Feb. 23, 2010, 9 pages.
Office Action for U.S. Appl. No. 12/113,724, dated Jun. 24, 2010, 12 pages.
Office Action for U.S. Appl. No. 12/133,738, dated Sep. 10, 2010, 11 pages.
Office Action for U.S. Appl. No. 12/133,761, dated Sep. 7, 2010, 11 pages.
Office Action for U.S. Appl. No. 12/139,305, dated Jan. 13, 2010, 12 pages.
Office Action for U.S. Appl. No. 12/182,073, dated Jun. 28, 2010, 20 pages.
Office Action for U.S. Appl. No. 12/182,083, datd Jun. 24, 2010, 8 pages.
Office Action for U.S. Appl. No. 12/264,160, dated Jun. 3, 2010, 5 pages.
Office Action for U.S. Appl. No. 12/422,223, dated Dec. 22, 2011, 11 pages.
Office Action for U.S. Appl. No. 12/536,852, dated Jun. 25, 2010, 8 pages.
Office Action for U.S. Appl. No. 12/536,852, dated Oct. 18, 2010, 10 pages.
Office Action for U.S. App. No. 12/619,502, dated Sep. 7, 2010, 6 pages.
Office Action for U.S. Appl. No. 12/829,337, dated Oct. 5, 2012, 10 pages.
Office Action from European Patent Application No. 05771646.6, dated Aug. 16, 2011, 3 pages.
Office Action from European Patent Application No. 05771646.6, dated Aug. 19, 2009, 4 pages.

(56) **References Cited**

OTHER PUBLICATIONS

Office Action from European Patent Application No. 05771646.6, dated Jun. 2, 2010, 5 pages.
Office Action from European Patent Application No. 05856669.6, dated Aug. 4, 2011, 7 pages.
Office Action from European Patent Application No. 05856669.6, dated Jun. 7, 2010, 5 pages.
Office Action from European Patent Application No. 05856669.6, dated Nov. 8, 2010, 6 pages.
Office Action from Japanese Patent Application No. 2007-521636 dated Mar. 1, 2011, 4 pages.
Office Action from Japanese Patent Application No. 2011-121598 dated May 22, 2012, 5 pages.
Office Action from Japanese Patent Application No. 2011-121598 dated Oct. 18, 2011, 9 pages.
Office Action from Japanese Patent Application No. 2012-006893 dated May 22, 2012, 4 pages.
Office Action from Japanese Patent Application No. 2012-169323 dated Oct. 16, 2012, 4 pages.
Office Action from Japanese Patent Application No. 2013-000959 dated Jan. 31, 2014, 6 pages.
Ohara T.J., et al., ""Wired" Enzyme Electrodes for Amperometric Determination of Glucose or Lactate in the Presence of Interfering Substances," Anal Chem, vol. 66, 1994, pp. 2451-2457.
Ohara T.J., et al., "Glucose Electrodes Based on Cross-Linked [Os(bpy)2Cl](+/2+) Complexed Poly(1-Vinylimidazole) Films," Analytical Chemistry, vol. 65, Dec. 1993, pp. 3512-3517.
Okuda, et al., "Mutarotase Effect on Micro Determinations of D-Glucose and its Anomers with β D-Glucose Oxidase," Anal Biochem, vol. 43 (1), 1971, pp. 312-315.
Oxford English Dictionary Online, Definition of "Impending," http://www.askoxford.com/results/?view=devdict&field-12668446_Impending&branch=, Jan. 11, 2010, 1 page.
Palmisano F., et al., "Simultaneous Monitoring of Glucose and Lactate by an Interference and Cross-Talk Free Dual Electrode Amperometric Biosensor Based on Electropolymerized Thin Films," Biosensors & Bioelectronics, vol. 15, 2000, pp. 531-539.
Panetti T.S., "Differential Effects of Sphingosine 1-Phosphate and Lysophosphatidic Acid on Endothelial Cells," Biochimica et Biophysica Acta, vol. 1582, 2002, pp. 190-196.
Panteleon A.E., et al., "The Role of the Independent Variable to Glucose Sensor Calibration," Diabetes Technology & Therapeutics, vol. 5 (3), 2003, pp. 401-410.
Paramount PDS., "Durometer Made Easy (R) / Durometer Hardness Scales—General Reference Guide," Paramount Industries Inc., 2008, 1 page.
Park I.B., et al., "Gas Separation Properties of Polysiloxane/Polyether Mixed Soft Segment Urethane Urea Membranes," Journal of Membrane science, vol. 204, 2002, pp. 257-269.
Parker R.S., et al., "A Model-Based Algorithm for Blood Glucose Control in Type I Diabetic Patients," IEEE Trans Biomed Engg (BME), vol. 46(2), 1999, pp. 148-157.
Patel H., et al., "Amperometric Glucose Sensors Based on Ferrocene Containing Polymeric Electron Transfer Systems—A Preliminary Report," Biosensors & Bioelectronics, vol. 18, 2003, pp. 1073-1076.
Peacock W.F., et al., "Cardiac Troponin and Outcome in Acute Heart Failure," N. Engl. J. Med., vol. 358, 2008, pp. 2117-2126.
Pegoraro M., et al., "Gas Transport Properties of Siloxane Polyurethanes," Journal of Applied Polymer Science, vol. 57, 1995, pp. 421-429.
Peguin S., et al., "Pyruvate Oxidase and Oxaloacetate Decarbozylase Enzyme Electrodes—Simultaneous Determination of Transaminases with a Two-electrode-based Analyzer," Analytica Chimica Acta, vol. 222, 1989, pp. 83-93.
Pfeiffer E.F., "The Glucose Sensor: The Missing Link in Diabetes Therapy," Horm Metab Res Suppl., vol. 24, 1990, pp. 154-164.
Pfeiffer E.F., et al., "On Line Continuous Monitoring of Subcutaneous Tissue Glucose is Feasible by Combining Portable Glucosensor with Microdialysis," Horm. Metab. Res., vol. 25, 1993, pp. 121-124.
Phillips R.E., et al., "Biomedical Applications of Polyurethanes: Implications of Failure Mechanisms," Journal of Biomedical application, vol. 3, Oct. 1988, pp. 206-227.
Phillips R.P., "A High Capacity Transcutaneous Energy Transmission System," ASIAO Journal, vol. 41, 1995, pp. M259-M262.
Pichert J.W., et al., "Issues for the Coming Age of Continuous Glucose Monitoring," Diabetes Educator, vol. 26 (6), Nov.-Dec. 2000, pp. 969-980.
Pickup J.C., et al., "Developing Glucose Sensors for In Vivo Use," Elsevier Science Publishers Ltd (UK), Tibtech, vol. 11, 1993, pp. 285-291.
Pickup J.C., et al., "Implantable Glucose Sensors: Choosing the Appropriate Sensor Strategy," Biosensors, vol. 3, (1987/1988), pp. 335-346.
Pickup J.C., et al., "In Vivo Molecular Sensing in Diabetes Mellitus: An Implantable Glucose Sensor with Direct Electron Transfer," Diabetologia, vol. 32, 1989, pp. 213-217.
Pickup J.C., et al., "Potentially-Implantable, Amperometric Glucose Sensors with Mediated Electron Transfer: Improving the Operating Stability," Biosensors, vol. 4, 1989, pp. 109-119.
Pickup J.C., et al., "Progress Towards in Vivo Glucose Sensing with a Ferrocene-Mediated Amperometric Enzyme Electrode," Horm Metab Res Suppl, vol. 20, 1988, pp. 34-36.
Pickup J.C., et al., "Responses and Calibration of Amperometric Glucose Sensors Implanted in the Subcutaneous Tissue of Man," ACTA Diabetol, vol. 30, 1993, pp. 143-148.
Pineda L.M., et al., "Bone Regeneration with Resorbable Polymeric Membranes. III. Effect of Poly(L-lactide) Membrane Pore Size on the Bone Healing Process in Large Defects," Journal of Biomedical Materials Research, vol. 31, 1996, pp. 385-394.
Pinner S.H., et al., "Cross-Linking of Cellulose Acetate by Ionizing Radiation," Nature, vol. 184, Oct. 24, 1959, pp. 1303-1304.
Pishko M.V., et al., "Amperometric Glucose Microelectrodes Prepared Through Immobilization of Glucose Oxidase in Redox Hydrogels," Analytical Chemistry, vol. 63 (20), 1991, pp. 2268-2272.
Pitzer K.R., et al., "Detection of Hypogylcemia with the Glucowatch Biographer," Diabetes Care, vol. 24 (5), 2001, pp. 881-885.
Poirier J.Y., et al., "Clinical and Statistical Evaluation of Self-Monitoring Blood Glucose Meters," Diabetes Care, vol. 21 (11), Nov. 1998, pp. 1919-1924.
Poitout V., et al., "A Glucose Monitoring System for on Line Estimation in Man of Blood Glucose Concentration Using a Miniaturized Glucose Sensor Implanted in the Subcutaneous Tissue and a Wearable Control Unit," Diabetologia, vol. 36, 1993, pp. 658-663.
Poitout V., et al., "Development of a Glucose Sensor for Glucose Monitoring in Man: The Disposable Implant Concept," Clinical Materials, vol. 15, 1994, pp. 241-246.
Poitout V., et al., "In Vitro and In Vivo Evaluation in Dogs of a Miniaturized Glucose Sensor," ASAIO Transactions, vol. 37, 1991, pp. M298-M300.
Postlethwaite T.A., et al., "Interdigitated Array Electrode as an Alternative to the Rotated Ring-Disk Electrode for Determination of the Reaction Products of Dioxygen Reduction," Analytical Chemistry, vol. 68 (17), Sep. 1996, pp. 2951-2958.
Prabhu V.G., et al., "Electrochemical Studies of Hydrogen Peroxide at a Platinum Disc Electrode," Electrochimica Acta, vol. 26 (6), 1981, pp. 725-729.
Preliminary Amendment for U.S. Appl. No. 12/052,489, filed Mar. 20, 2008, 8 pages.
Quinn C.A.P., et al., "Biocompatible, Glucose-Permeable Hydrogel for In situ Coating of Implantable Biosensors," Biomaterials, vol. 18 (24), 1997, pp. 1665-1670.
Quinn C.P., et al., "Kinetics of Glucose Delivery to Subcutaneous Tissue in Rats Measured with 0.3-mm Amperometric Microsensors," The American Physiological Society, vol. 269, 1995, pp. E155-E161.

(56) **References Cited**

OTHER PUBLICATIONS

Rabah M.A., et al., "Electrochemical Wear of Graphite Anodes During Electrolysis of Brine," Carbon, vol. 29 (2), 1991, pp. 165-171.

Radovsky A.N., et al., "Effects of Dexamethasone Elution on Tissue Reaction Around Stimulating Electrodes of Endocardial Pacing Leads in Dogs," American Heart Journal, vol. 117 (6), Jun. 1989, pp. 1288-1298.

Rafael E., "Cell Transplantation and Immunoisolation: Studies on a Macroencapsulation Device," Departments of Transplantation Surgery and Pathology, Karolinska Institutet, Huddinge Hospital, Stockholm, Sweden, 1999, pp. 1-82.

Ratner B.D., "Reducing Capsular Thickness and Enhancing Angiogenesis around Implant Drug Release Systems," Journal of Controlled Release, vol. 78, 2002, pp. 211-218.

Rawlings R.A., et al., "Translating Glucose Variability Metrics into the Clinic via Continuous Glucose Monitoring: A Graphical User Interface for Diabetes Evaluation (CGM-Guide)," Diabetes Technology & Therapeutics, vol. 13 (12), 2011, pp. 1241-1248.

Raya Systems Pioneers, "Raya Systems Pioneers Healthy Video Games," PlayRight, Nov. 1993, pp. 14-15.

Reach G., "A Method for Evaluating in vivo the Functional Characteristics of Glucose Sensors," Biosensors, vol. 2, 1986, pp. 211-220.

Reach G., "Which Threshold to Detect Hypoglycemia? Value of Receiver-Operator Curve Analysis to Find a Compromise Between Sensitivity and Specificity," Diabetes Care, vol. 24 (5), May 2001, pp. 803-804.

Reach G., et al., "Can Continuous Glucose Monitoring Be Used for the Treatment of Diabetes?," Analytical Chemistry, vol. 64 (6), Mar. 15, 1992, pp. 381A-386A.

Reach, et al., "Clinical Sensors for In Vivo Monitoring," Advances in Biosensors, COMAC Biomedical Engineering Concerted Action on Chemical Sensors for In Vivo Monitoring, Supplement 1: Chapter 1, 1993, pp. 7-28.

Rebrin K., et al., "Automated Feedback Control of Subcutaneous Glucose Concentration in Diabetic Dogs," Diabetologia, vol. 32, 1989, pp. 573-576.

Rebrin K., et al., "Subcutaenous Glucose Monitoring by Means of Electrochemical Sensors: Fiction or Reality?," Journal of Biomedical Engineering, vol. 14, Jan. 1992, pp. 33-40.

Rebrin K., et al., "Subcutaneous Glucose Predicts Plasma Glucose Independent of Insulin: Implications for Continuous Monitoring," The American Physiological Society, vol. 277, 1999, pp. E561-E571.

Reush, "Organometallic Compounds," Chemical Reactivity, Virtual Textbook of Organic Chemistry, Retrieved from http://www.cem.msu.edu/-reuschlVirtualText/orgmetal.htm, 2004, pp. 1-16.

Revuelta J.M., et al., "Expanded Polytetrafluoroethylene Surgical Membrane for Pericardial Closure," The Journal of Thoracic and cardiovascular Surgery, vol. 89 (3), Mar. 1985, pp. 451-455.

Rhodes R.K., et al., "Prediction of Pocket-Portable and Implantable Glucose Enzyme Electrode Performance from Combined Species Permeability and Digital Simulation Analysis," Analytical Chemistry, vol. 66 (9), May 1, 1994, pp. 1520-1529.

Rigla M., et al., "Real-Time Continuous Glucose Monitoring Together with Telemedical Assistance Improves Glycemic Control and Glucose Stability in Pump-Treated Patients," Diabetes Technology & Therapeutics, vol. 10 (3), 2008, pp. 194-199.

Rinken T., et al., "Calibration of Glucose Biosensors by Using Pre-Steady State Kinetic Data," Biosensors & Bioelectronics, vol. 13, 1998, pp. 801-807.

Rivers E.P., et al., "Central Venous Oxygen Saturation Monitoring in the Critically Ill Patient," Current Opinion in Critical Care, 2001, vol. 7, pp. 204-211.

Roe J.N., et al., "Bloodless Glucose Measurements," Critical Reviews™ in Therapeutic Drug Carrier Systems, vol. 15 (3), 1998, pp. 199-241.

Sachlos E., et al., "Making Tissue Engineering Scaffolds Work Review on the Application of Sold Freeform Fabrication Technology to the Production of Tissue Engineering Scaffolds," European Cells and Materials, vol. 5, 2003, pp. 29-40.

Sakakida M., et al., "Development of Ferrocene-Mediated Needle-Type Glucose Sensor as a Measure of True Subcutaneous Tissue Glucose Concentrations," Artif. Organs Today, vol. 2 (2), 1992, pp. 145-158.

Sakakida M., et al., "Ferrocene-Mediated Needle Type Glucose Sensor Covered with Newly Designed Biocompatible Membrane," Sensors and Actuators B, vol. 13-14, 1993, pp. 319-322.

Salardi S., et al., "The Glucose Area Under the Profiles Obtained with Continuous Glucose Monitoring System Relationships with HbA1C in Pediatric Type 1 Diabetic Patients," Diabetes Care, vol. 25 (10), Oct. 2002, pp. 1840-1844.

Salehi C., et al., "A Telemetry-Instrumentation System for Long-Term Implantable Glucose and Oxygen Sensors," Analytical Letters, vol. 29 (13), 1996, pp. 2289-2308.

Samuels M.P., "The Effects of Flight and Altitude," Arch Dis Child, vol. 89, 2004, pp. 448-455.

San Diego Plastics Inc, "Polyethylene," Datasheet, Retrieved from http://www.sdplastics.com/polyeth.html on Aug. 19, 2009, 7 pages.

Sanders E., et al., "Fibrous Encapsulation of Single Polymer Microfibers Depends on their Vertical Dimension in Subcutaneous Tissue Polymer Microfibers," Journal of Biomedical Material Research, vol. 67A, 2003, pp. 1181-1187.

Sansen W., et al., "A Smart Sensor for the Voltammetric Measurement of Oxygen or Glucose Concentrations," Sensors and Actuators B1, 1990, pp. 298-302.

Sansen W., et al., "Glucose Sensor with Telemetry System," In Implantable Sensors for Closed Loop Prosthetic Systems edited by Ko W.H, Chapter 12, 1985, pp. 167-175.

Schaffar B.P.H., "Thick Film Biosensors for Metabolites in Undiluted Whole Blood and Plasma Samples," Analytical Bioanalytical Chemistry, Dec. 2001, vol. 372, pp. 254-260.

Schmidt F.J., et al., "Calibration of a Wearable Glucose Sensor," The International Journal of Artificial Organs, Wichtig Publishing, IT, vol. 15 (1), Jan. 1, 1992, pp. 55-61.

Schmidt F.J., et al., "Glucose Concentration in Subcutaneous Extracellular Space," Diabetes Care, vol. 16 (5), May 1993, pp. 695-700.

Schmidtke D.W., et al., "Accuracy of the One-Point in Vivo Calibration of "Wired" Glucose Oxidase Electrodes Implanted in Jugular Veins of Rats in Periods of Rapid Rise and Decline of the Glucose Concentration," Analytical Chemistry, vol. 70 (10), May 15, 1998, pp. 2149-2155.

Schmidtke D.W., et al., "Measurement and Modeling of the Transient Difference Between Blood and Subcutaneous Glucose Concentrations in the Rat After Injection of Insulin," Proceedings of the National Academy of Sciences, vol. 95, Jan. 1998, pp. 294-299.

Schoemaker M., et al., "The SCGMI System: Subcutaneous Continuous Glucose Monitoring Based on Microdialysis Technique," Diabetes Technology & Therapeutics, vol. 5 (4), 2003, pp. 599-608.

Schoonen A.J.M., et al., "Development of a Potentially Wearable Glucose Sensor for Patients with Diabetes Mellitus: Design and In-vitro Evaluation," Biosensors & Bioelectronics, vol. 5, 1990, pp. 37-46.

Schuler, et al., "Modified Gas-Permeable Silicone Rubber Membranes for Covalent Immobilisation of Enzymes and their Use in Biosensor Development," Analyst, 1999, vol. 124, pp. 1181-1184.

Selam J.L., "Management of Diabetes with Glucose Sensors and Implantable Insulin Pumps," From the Dream of the 60s to the Realities of the 90s, ASAIO Journal 1997, vol. 43, pp. 137-141.

Service F.J., et al., "Mean Amplitude of Glycemic Excursions, A Measure of Diabetic Instability," Diabetes, vol. 19 (9), Sep. 1970, pp. 644-655.

Service F.J., et al., "Measurements of Glucose Control," Diabetes Care, vol. 10 (2), Mar.-Apr. 1987, pp. 225-237.

Service R.F., "Can Sensors Make a Home in the Body?," Science, Materials Science: Soft Surface, Volume. 297, Aug. 9, 2002, pp. 962-963.

Sharkawy A.A., et al., "Engineering the Tissue Which Encapsulates Subcutaneous Implants. I. Diffusion Properties," Journal of Biomedical Materials Research, vol. 37, 1997, pp. 401-412.

(56) **References Cited**

OTHER PUBLICATIONS

Shaw G.W., et al., "In Vitro Testing of a Simply Constructed, Highly Stable Glucose Sensor Suitable for Implantation in Diabetic Patients," Biosensors & Bioelectronics, vol. 6, 1991, pp. 401-406.

Shichiri M., et al., "Glycaemic Control in Pancreatectomized Dogs with a Wearable Artificial Endocrine Pancreas," Diabetologia, vol. 24, 1983, pp. 179-184.

Shichiri M., et al., "Membrane Design for Extending the Long-Life of an Implantable Glucose Sensor," Diabetes Nutrition & Metabolism, vol. 2 (4), 1989, pp. 309-313.

Shichiri M., et al., "Needle Type Glucose Sensor for Wearable Artificial Endocrine Pancreas," In Implantable Sensors for Closed-Loop Prosthetic Systems edited by Ko W.H, Chapter 15, 1985, pp. 197-210.

Shichiri M., et al., "Telemetry Glucose Monitoring Device with Needle-Type Glucose Sensor: A Useful Tool for Blood Glucose Monitoring in Diabetic Individuals," Diabetes Care, vol. 9 (3), May-Jun. 1986, pp. 298-301.

Shichiri M., et al., "Wearable Artificial Endocrine Pancreas with Needle-Type Glucose Sensor," Preliminary Communication, Lancet, vol. 2, Nov. 20, 1982, pp. 1129-1131.

Shichiri, et al., "In Vivo Characteristics of Needle-Type Glucose Sensor-Measurements of Subcutaneous Glucose Concentrations in Human Volunteers," Implantable Glucose Sensors—The State of the Art, Hormone and Metabolic Research Supplement Series, 1988, vol. 20, pp. 17-20.

Shults M.C., et al., "A Telemetry-Instrumentation System for Monitoring Multiple Subcutaneously Implanted Glucose Sensors," IEEE Transactions on Biomedical Engineering, vol. 41 (10), Oct. 1994, pp. 937-942.

Sieminski, et al., "Biomaterial-Microvasculature Interactions," Biomaterials, 2000, vol. 21, pp. 2233-2241.

Sigma-Aldrich Corp "Nafion® 117 Solution Product Description, Product No. 70160," retrieved from https//:www.sigmaaldrich.com/cgi-bin/hsrun/Suite7/Suite/HAHTpage/Suite.HsExternalProd on Apr. 7, 2005, 1 page.

Sigma-Aldrich Corp., "Cellulose Acetate," Product Description, Product No. 419028, St. Louis, MO, 2005, 1 page.

Skyler J.S., "The Economic Burden of Diabetes and the Benefits of Improved Glycemic Control: The Potential Role of a Continuous Glucose Monitoring System," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S7-S12.

Slater-Maclean L., et al., "Accuracy of Glycemic Measurements in the Critically Ill" Diabetes Technology and Therapeutics, vol. 10 (3), 2008, pp. 169-177.

Smith B., et al., "An Externally Powered, Multichannel, Implantable Stimulator-Telemeter for Control of Paralyzed Muscle," IEEE Transactions on Biomedical Engineering, vol. 45 (4), Apr. 1998, pp. 463-475.

Smith, "The Scientist and Engineer's Guide to Digital Signal Processing," California Technical Publishing, 1997-2007, retrieved from http://www.dspguide.com/ch19.htm on Jan. 1, 2009, 2 pages.

Smith, et al.,"A Comparison of Islet Transplantation and Subcutaneous Insulin Injections for the Treatment of Diabetes Mellitus," Computers in Biology and Medicine, 1991, vol. 21 (6), pp. 417-427.

Smooth-On, "Durometer Shore Hardness Scale," downloaded from https://www.smooth-on.com/page/durometer-shore-hardness-scale/ on May 19, 2016, 1 page.

Sokol L., et al., "Immobilized-Enzyme Rate-Determination Method for Glucose Analysis," Clinical Chemistry, vol. 26 (1), 1980, pp. 89-92.

Sokolov S., et al., "Metrological Opportunities of the Dynamic Mode of Operating an Enzyme Amperometric Biosensor," Medical Engineering & Physics, vol. 17 (6), 1995, pp. 471-476.

Sparacino G., et al., "Continuous Glucose Monitoring Time Series and Hypo-Hyperglycemia Prevention: Requirements, Methods, Open Problems," Current Diabetes Reviews, vol. 4 (3), 2008, pp. 181-192.

Sproule B.A., et al., "Fuzzy Pharmacology: Theory and Applications," Trends in Pharmacological Sciences, vol. 23 (9), Sep. 2002, pp. 412-417.

Sriyudthsak M., et al., "Enzyme-Epoxy Membrane Based Glucose Analyzing System and Medical Applications," Biosensors & Bioelectronics, vol. 11 (8), 1996, pp. 735-742.

Steil G.M., et al., "Determination of Plasma Glucose During Rapid Glucose Excursions with a Subcutaneous Glucose Sensor," Diabetes Technology & Therapeutics, vol. 5 (1), 2003, pp. 27-31.

Stern M., et al., "Electrochemical Polarization: I. A Theoretical Analysis of the Shape of Polarization Curves," Journal of the Electrochemical Society, vol. 104 (1), Jan. 1957, pp. 56-63.

Sternberg F., et al., "Does Fall in Tissue Glucose Precede Fall in Blood Glucose?," Diabetologia, vol. 39, 1996, pp. 609-612.

Sternberg R., et al., "Study and Development of Multilayer Needle-type Enzyme Based Glucose Microsensors," Biosensors, vol. 4, 1988, pp. 27-40.

Sternberg, et al., "Covalent Enzyme Coupling on Cellulose Acetate Membranes for Glucose Sensor Development," Anal Chem, Dec. 1988, vol. 60(24), pp. 2781-2786.

Stokes, "Polyether Polyurethanes: Biostable or Not," Journal of Biomaterials Applications, Oct. 1988, vol. 3, pp. 228-259.

Street J.O., et al., "A Note on Computing Robust Regression Estimates Via Iteratively Reweighted Least Squares," The American Statistician, vol. 42 (2), May 1988, pp. 152-154.

Street, et al., "Islet Graft Assessment in the Edmonton Protocol: Implications for Predicting Long-Term Clinical Outcome," Diabetes, 2004, vol. 53, pp. 3107-3114.

Suh, et al., "Behavior of Fibroblasts on a Porous Hyaluronic Acid Incorporated Collagen Matrix," Yonsei Medical Journal, 2002, vol. 43 (2), pp. 193-202.

Sumino T., et al., "Preliminary Study of Continuous Glucose Monitoring with a Microdialysis Technique," Proceedings of the 20th Annual International Conference of the IEEE Engineering in Medicine and Biology Society, vol. 20 (4), 1998, pp. 1775-1778.

Supplementary European Search Report for Application No. 09729717.0, dated Oct. 21, 2011, 6 pages.

Takatsu, et al., "Solid State Biosensors Using Thin-Film Electrodes," Sensors and Actuators, 1997, vol. 11, pp. 309-317.

Takegami S., et al., "Pervaporation of Ethanol/Water Mixtures Using Novel Hydrophobic Membranes Containing Polydimethylsiloxane," Journal of Membrane Science, vol. 75, 1992, pp. 93-105.

Tamura T., et al., "Preliminary Study of Continuous Glucose Monitoring with a Microdialysis Technique and a Null Method—A Numerical Analysis," Frontiers of Medical & Biological Engineering, vol. 10 (2), 2000, pp. 147-156.

Tanenberg R.J., et al., "Continuous Glucose Monitoring System: A New Approach to the Diagnosis of Diabetic Gastroparesis," Diabetes Technology & Therapeutics, vol. 2, Supplement 1, 2000, pp. S73-S80.

Tang, et al., "Fibrin(ogen) Mediates Acute Inflammatory Responses to Biomaterials," J.Exp.Med, 1993, vol. 178, pp. 2147-2156.

Tang, et al., "Inflammatory Responses to Biomaterials," Am J Clin Pathol, 1995, vol. 103, pp. 466-471.

Tang, et al., "Mast Cells Mediate Acute Inflammatory Responses to Implanted Biomaterials," Proceedings of the National Academy of Sciences of the USA, 1998, vol. 95, pp. 8841-8846.

Tang, et al., "Molecular Determinants of Acute Inflammatory Responses to Biomaterials," J Clin Invest, 1996, vol. 97, pp. 1329-1334.

Tatsuma T., et al., "Oxidase/Peroxidase Bilayer-Modified Electrodes as Sensors for Lactate, Pyruvate, Cholesterol and Uric Acid," Analytica Chimica Acta, vol. 242, 1991, pp. 85-89.

Taub M.B., et al., "Numerical Simulation of the Effect of Rate of Change of Glucose on Measurement Error of Continuous Glucose Monitors," Journal of Diabetes Science and Technology, vol. 1 (5), Sep. 2007, pp. 685-694.

Thijssen P.C.,"A Kalman Filter for Calibration, Evaluation of Unknown Samples and Quality Control in Drifting Systems," Part 2,Optimal Designs, Analytica chimica Acta, vol. 162, 1984, pp. 253-262.

(56) **References Cited**

OTHER PUBLICATIONS

Thijssen, et al., "A Kalman Filter for Calibration, Evaluation of Unknown Samples and Quality Control in Drifting Systems," Part 1,Theory and Simulations, Analytica chimica Acta, 1984, vol. 156, pp. 87-101.
Thijssen, et al., "A Kalman Filter for Calibration, Evaluation of Unknown Samples and Quality Control in Drifting Systems," Part 3,Variance Reduction ,Analytica chimica Acta, 1985, vol. 173, pp. 265-272.
Thijssen, et al., "A Kalman Filter for Calibration, Evaluation of Unknown Samples and Quality Control in Drifting Systems," Part 4,Flow Injection Analysis, Analytica chimica Acta, 1985, vol. 174, pp. 27-40.
Thome V., et al., "(Abstract) Can the Decrease in Subcutaneous Glucose Concentration Precede the Decrease in Blood Glucose Level? Proposition for a Push-Pull Kinetics Hypothesis," Horm. metab. Res., vol. 27, 1995, p. 53.
Thome-Duret V., et al., "Continuous Glucose Monitoring in the Free-Moving Rat," Metabolism, vol. 47 (7), Jul. 1998, pp. 799-803.
Thome-Duret V., et al., "Modification of the Sensitivity of Glucose Sensor Implanted into Subcutaneous Tissue," Diabetes & Metabolism, vol. 22, 1996, pp. 174-178.
Thome-Duret V., et al., "Use of a Subcutaneous Glucose Sensor to Detect Decreases in Glucose Concentration Prior to Observation in Blood," Analytical Chemistry, vol. 68 (21), Nov. 1, 1996, pp. 3822-3826.
Thompson M., et al., "In Vivo Probes: Problems and Perspectives," Clinical Biochemistry, vol. 19 (5), Oct. 1986, pp. 255-261.
Tibell, et al., "Survival of Macroencapsulated Allogeneic Parathyroid Tissue One Year after Transplantation in Nonimmunosuppressed Humans," Cell Transplantation, 2001, vol. 10, pp. 591-599.
Tierney M.J., et al., "Effect of Acetaminophen on the Accuracy of Glucose Measurements Obtained with the GlucoWatch Biographer," Diabetes Technology & Therapeutics, vol. 2 (2), 2000, pp. 199-207.
Tierney M.J., et al., "The Gluco Watch® Biographer: A Frequent, Automatic and Noninvasive Glucose Monitor," Annals of Medicine, vol. 32, 2000, pp. 632-641.
Tilbury J.B., et al., "Receiver Operating Characteristic Analysis for Intelligent Medical Systems—A New Approach for Finding Confidence Intervals," IEEE Transactions on Biomedical Engineering, vol. 47 (7), Jul. 2000, pp. 952-963.
Torjman M.C., et al., "Glucose Monitoring in Acute Care: Technologies on the Horizon," Journal of Diabetes Science and Technology, vol. 2 (2), Mar. 2008, pp. 178-181.
Trajanoski Z., et al., "Neural Predictive Controller for Insulin Delivery Using the Subcutaneous Route," IEEE Transactions on Biomedical Engineering, vol. 45(9), 1998, pp. 1122-1134.
Trecroci D., "A Glimpse into the Future-Continuous Monitoring of Glucose with a Microfiber," Diabetes Interview, Jul. 2002, pp. 42-43.
Tse P.S.H., et al., "Time-Dependent Inactivation of Immobilized Glucose Oxidase and Catalase," Biotechnology & Bioengineering, vol. 29, 1987, pp. 705-713.
Turner A.P.F., "Amperometric Biosensor based on Mediator-Modified Electrodes," Methods in Enzymology, 1988, vol. 137, pp. 90-103.
Turner A.P.F., et al., "Carbon Monoxide: Acceptor Oxidoreductase from Pseudomonas Thermocarboxydovorans Strain C2 and its Use in a Carbon Monoxide Sensor," Analytica Chimica Acta, vol. 163, 1984, pp. 161-174.
Turner A.P.F., et al., "Diabetes Mellitus: Biosensors for Research and Management," Biosensors, vol. 1, 1985, pp. 85-115.
Unger J., et al., "Glucose Control in the Hospitalized Patient," Emergency Medicine, vol. 36 (9), 2004, pp. 12-18.
Updike S.J., et al., "A Subcutaneous Glucose Sensor with Improved Longevity, Dynamic Range, and Stability of Calibration," Diabetes Care, vol. 23 (2), Feb. 2000, pp. 208-214.
Updike S.J., et al., "Continuous Glucose Monitor Based on an Immobilized Enzyme Electrode Detector," Journal of Laboratory and Clinical Medicine, vol. 93(4), 1979, pp. 518-527.
Updike S.J., et al., "Enzymatic Glucose Sensor: Improved Long-Term Performance in Vitro and In Vivo," ASAIO Journal, vol. 40 (2), Apr.-Jun. 1994, pp. 157-163.
Updike S.J., et al., "Implanting the Glucose Enzyme Electrode: Problems, Progress, and Alternative Solutions," Diabetes Care, vol. 5 (3), May-Jun. 1982, pp. 207-212.
Updike S.J., et al., "Laboratory Evaluation of New Reusable Blood Glucose Sensor," Diabetes Care, vol. 11 (10), Nov.-Dec. 1988, pp. 801-807.
Updike S.J., et al., "Principles of Long-Term Fully Implanted Sensors with Emphasis on Radiotelemetric Monitoring of Blood Glucose Form Inside a Subcutaneous Foreign Body Capsule (FBC)," Edited by Fraser D M, Biosensors in the Body: Continuous in vivo Monitoring, John Wiley & Sons Ltd., New York, 1997, Chapter 4, pp. 117-137.
Updike S.J., et al., "The Enzyme Electrode," Nature, vol. 214, Jun. 3, 1967, pp. 986-988.
Utah Medical Products Inc., "Deltran—Disposable Blood Pressure Tranducers," Product Specifications, 2003-2006, 6 pages.
Vadgama P., "Diffusion Limited Enzyme Electrodes," NATO ASI Series: Series C, Math and Phys. Sci, vol. 226, 1988, pp. 359-377.
Vadgama P., "Enzyme Electrodes as Practical Biosensors," Journal of Medical Engineering & Technology, vol. 5 (6), Nov. 1981, pp. 293-298.
Valdes T.I., et al., "In Vitro and In Vivo Degradation of Glucose Oxidase Enzyme used for an Implantable Glucose Biosensor," Diabetes Technology & Therapeutics, vol. 2 (3), 2000, pp. 367-376.
Velho G., et al., "In Vitro and In Vivo Stability of Electrode Potentials in Needle-Type Glucose Sensors," Influence of Needle Material, Diabetes, vol. 38, Feb. 1989, pp. 164-171.
Velho G., et al., "Strategies for Calibrating a Subcutaneous Glucose Sensor," Biomed Biochim Acta, vol. 48 (11/12), 1989, pp. 957-964.
Vig, et al., "A Review of Sensor Sensitivity and Stability," IEEE/EIA International Frequency Control Symposium and Exhibition, 2000, pp. 30-33.
Von Woedtke T., et al., "In Situ Calibration of Implanted Electrochemical Glucose Sensors," Biomed. Biochim. Acta 48 vol. 11/12, 1989, pp. 943-952.
Von Woedtke T., et al., "Sterilization of Enzyme Glucose Sensors: Problems and Concepts," Biosensors & Bioelectronics, vol. 17, 2002, pp. 373-382.
Wade L.G., "Reactions of Aromatic Compounds," Organic Chemistry, Chapter 17, 5th edition, 2003, pp. 762-763.
Wagner, et al., "Continuous Amperometric Monitoring of Glucose in a Brittle Diabetic Chimpanzee with a Miniature Subcutaneous Electrode," Proc. Natl. Acad. Sci. USA, vol. 95, May 1998, pp. 6379-6382.
Wang J., "Electrochemical Glucose Biosensors," American Chemical Society, Chemical Reviews, Published on Web Dec. 23, 2007, pp. 1-12.
Wang J., et al., "Highly Selective Membrane-Free Mediator-Free Glucose Biosensor," Analytical Chemistry, vol. 66 (21), Nov. 1, 1994, pp. 3600-3603.
Wang X., et al., "Improved Ruggedness for Membrane-Based Amperometric Sensors using a Pulsed Amperometric Method," Analytical Chemistry, vol. 69 (21), Nov. 1, 1997, pp. 4482-4489.
Ward W.K., et al., "A New Amperometric Glucose Microsensor: In Vitro and Short-Term In Vivo Evaluation," Biosensors & Bioelectronics, vol. 17, 2002, pp. 181-189.
Ward W.K., et al., "Assessment of Chronically Subcutaneous Glucose Sensors in Dogs: The Effect of Surrounding Fluid Masses," ASAIO Journal, 1999, vol. 45 (6), pp. 555-561.
Ward W.K., et al., "Rise in Background Current Over Time in a Subcutaneous Glucose Sensor in the Rabbit," Relevance to Calibration and Accuracy, Biosensors & Bioelectronics, vol. 15, 2000, pp. 53-61.
Ward W.K., et al., "Understanding Spontaneous Output Fluctuations of an Amperometric Glucose Sensor: Effect of Inhalation Anesthesia and Use of a Nonenzyme Containing Electrode," ASAIO Journal, 2000, pp. 540-546.

(56) **References Cited**

OTHER PUBLICATIONS

Ward, et al., "A Wire-Based Dual-Analyte Sensor for Glucose and Lactate: In Vitro and In Vivo Evaluation," Diabetes Technology and Therapeutics, 2004, vol. 6 (3), pp. 389-401.
Wientjes K.J.C., "Development of a Glucose Sensor for Diabetic Patients," (Ph.D. Thesis), 2000, 212 pages.
Wikipedia., "Intravenous Therapy," http://en.wikipedia.org/wiki/Intravenous_therapy, Aug. 15, 2006, 6 pages.
Wilkins E., et al., "Glucose Monitoring: State of the Art and Future Possibilities," Med. Eng. Phys., vol. 18 (4), 1996, pp. 273-288.
Wilkins E., et al., "Integrated Implantable Device for Long-Term Glucose Monitoring," Biosensors & Bioelectronics, vol. 10, 1995, pp. 485-494.
Wilkins E.S., et al., "The Coated Wire Electrode Glucose Sensor," Horm Metab Res Suppl., vol. 20, 1988, pp. 50-55.
Wilson G.S., et al., "Enzyme-Based Biosensors for In Vivo Measurements," Chem. Rev., vol. 100, 2000, pp. 2693-2704.
Wilson G.S., et al., "Progress Toward the Development of an Implantable Sensor for Glucose," Clinical Chemistry, vol. 38 (9), 1992, pp. 1613-1617.
Wolpert H., "Establishing a Continuous Glucose Monitoring Program," Journal of Diabetes Science and Technology, Mar. 2008, vol. 2 (2), pp. 307-310.
Wood W D., et al., "Hermetic Sealing with Epoxy," Pave Technology-Mechanical Engineering, Mar. 1990, 3 pages.
Woodward S.C., "How Fibroblasts and Giant Cells Encapsulate Implants: Considerations in Design of Glucose Sensor," Diabetes Care, vol. 5 (3) May-Jun. 1982, pp. 278-281.
Worsley G.J et al., "Measurement of Glucose in Blood with a Phenylboronic Acid Optical Sensor," Journal of Diabetes Science and Technology, vol. 2 (2), Mar. 2008, pp. 213-220.
Wright M., et al., "Bioelectrochemical Dehalogenations Via Direct Electrochemistry of Poly(ethylene oxide)-Modified Myoglobin," Electrochemistry Communications, vol. 1, 1999, pp. 609-613.
Written Opinion for Application No. PCT/US2002/023903 dated Nov. 15, 2004, 5 pages.
Wu H., et al., "In Situ Electrochemical Oxygen Generation with an Immunoisolation Device," Annals New York Academy of Sciences, vol. 875, 1999, pp. 105-125.
Yamasaki Y., "The Development of a Needle-Type Glucose Sensor for Wearable Artificial Endocrine Pancreas," Medical Journal of Osaka University, vol. 35 (1-2), Sep. 1984, pp. 25-34.
Yamasaki Y., et al., "Direct Measurement of Whole Blood Glucose by a Needle-Type Sensor," Clinica Chimica Acta. 93, 1989, pp. 93-98.
Yang C., et al., "A Comparison of Physical Properties and Fuel Cell Performance of Nafion and Zirconium Phosphate/Nation Composite Membranes," Journal of Membrane Science, vol. 237, 2004, pp. 145-161.
Yang Q., et al., "Development of Needle-Type Glucose Sensor with High Selectivity," Science and Actuators B, vol. 46, 1998, pp. 249-256.
Yang S., et al., "A Glucose Biosensor Based on an Oxygen Electrode: In-Vitro Performances in a Model Buffer Solution and in Blood Plasma," Biomedical Instrumentation & Technology, vol. 30 (1), 1996, pp. 55-61.
Yang S., et al., "Glucose Biosensors with Enzyme Entrapped in Polymer Coating," Biomedical Instrument and Technology, Mar./Apr. 1995, vol. 29 (2), pp. 125-133.
Yang X., et al., "Polyelectrolyte and Molecular Host Ion Self-Assembly to Multilayer Thin Films: An Approach to Thin Film Chemical Sensors," Sensors and Actuators B, vol. 45, 1997, pp. 87-92.
Ye L., et al., "High Current Density Wired' Quinoprotein Glucose Dehydrogenase Electrode," Analytical Chemistry, vol. 65, 1993, pp. 238-241.
Zamzow K.L., et al., "Development and Evaluation of a Wearable Blood Glucose Monitor," ASAIO Transactions, vol. 36 (3), 1990, pp. M588-M591.
Zavalkoff S.R., et al., "Evaluation of Conventional Blood Glucose Monitoring as an Indicator of Integrated Glucose Values Using a Continuous Subcutaneous Sensor," Diabetes Care, vol. 25(9), 2002, pp. 1603-1606.
Zethelius B., et al., "Use of Multiple Biomarkers to Improve the Prediction of Death From Cardiovascular Causes," N. Engl. J. Med., vol. 358, May 2008, pp. 2107-2116.
Zhang Y., et al., "Electrochemical Oxidation of H2O2 on Pt and Pt + Ir Electrodes in Physiological Buffer and its Applicability to H2O2-Based Biosensors," J. Electro Analytical Chemistry, vol. 345, 1993, pp. 253-271.
Zhang Y., et al., "In Vitro and In Vivo Evaluation of Oxygen Effects on a Glucose Oxidase Based Implantable Glucose Sensor," Analytica Chimica Acta, vol. 281, 1993, pp. 513-520.
Zhang, et al., "Elimination of the Acetaminophen Interference in an Implantable Glucose Sensor," Analytical Chemistry, 1994, vol. 66 (7), pp. 1183-1188.
Zhu, et al., "Fabrication and Characterization of Glucose Sensors Based on a Microarray H2O2 Electrode," Biosensors & Bioelectronics, 1994, vol. 9, pp. 295-300.
Zhu, et al., "Planar Amperometric Glucose Sensor Based on Glucose Oxidase Immobilized by Chitosan Film on Prussian blue Layer," Sensors, 2002, vol. 2, pp. 127-136.
Ziaie, et al., "A Single-Channel Implantable Microstimulator for Functional Neuromuscular Stimulation," IEEE Transactions on Biomedical Engineering, 1997, vol. 44(10), pp. 909-920.

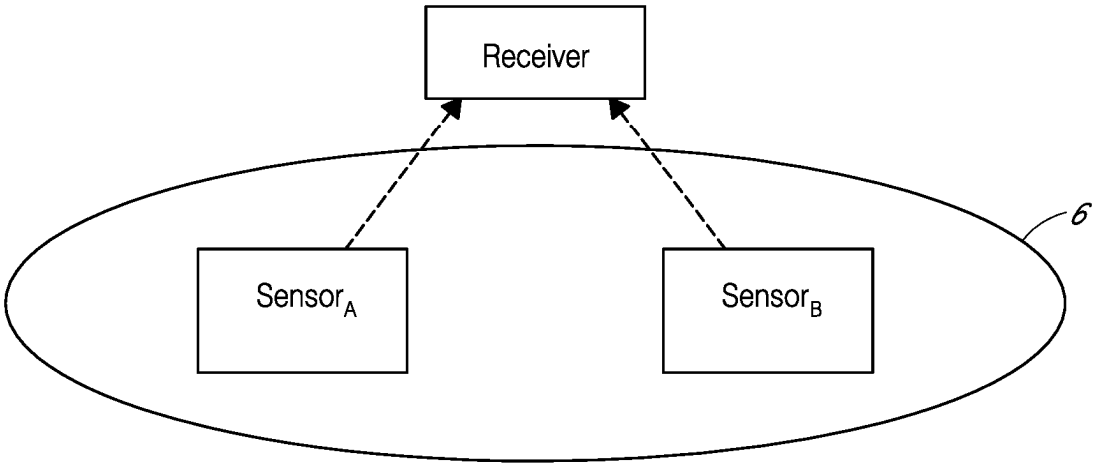

Receiver
Sensor$_A$
Sensor$_B$
6

FIG. 1A

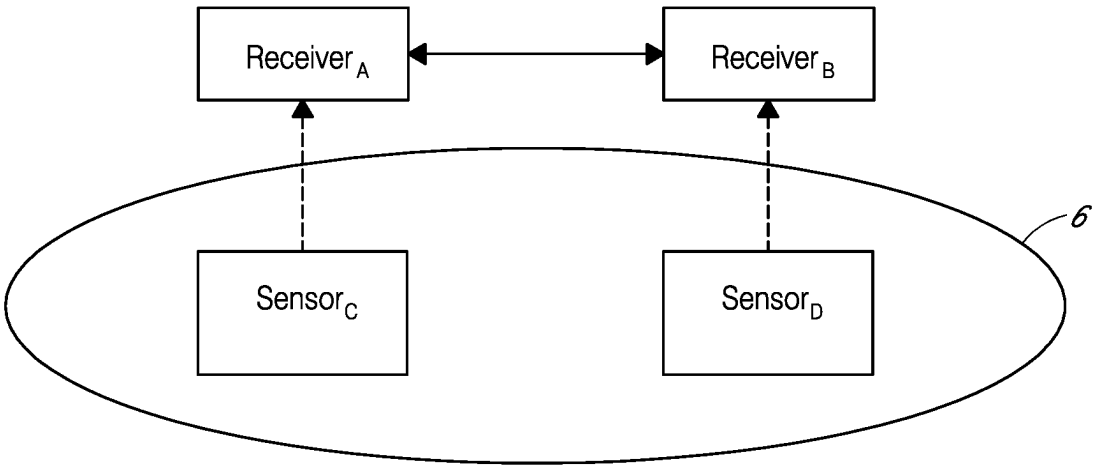

Receiver$_A$
Receiver$_B$
Sensor$_C$
Sensor$_D$
6

FIG. 1B

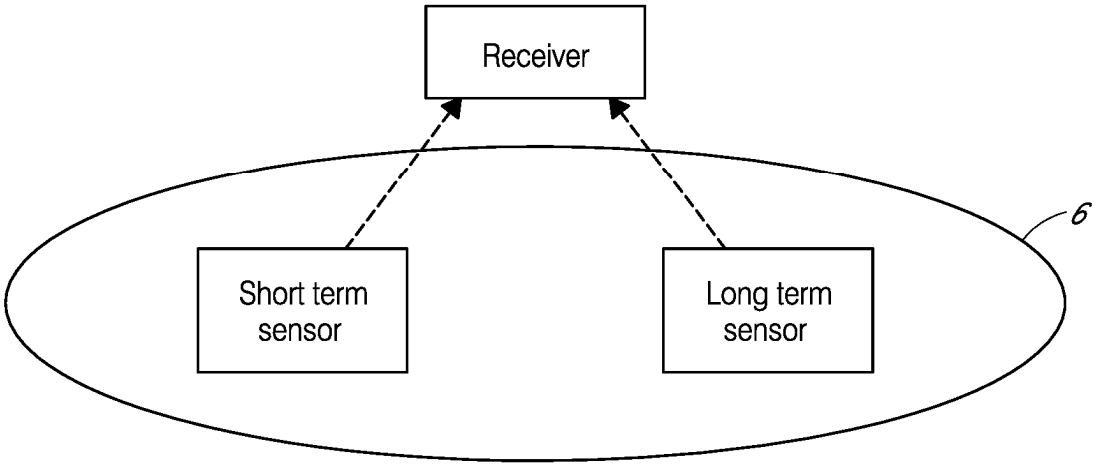

Receiver
Short term sensor
Long term sensor
6

FIG. 1C

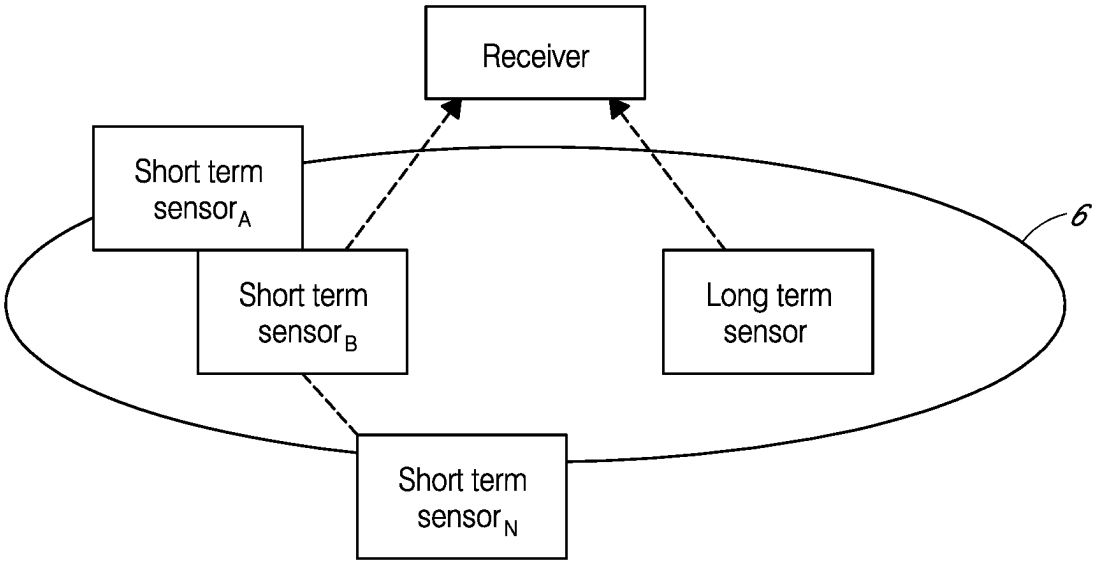

Receiver
Short term sensor$_A$
Short term sensor$_B$
Long term sensor
Short term sensor$_N$
6

FIG. 1D

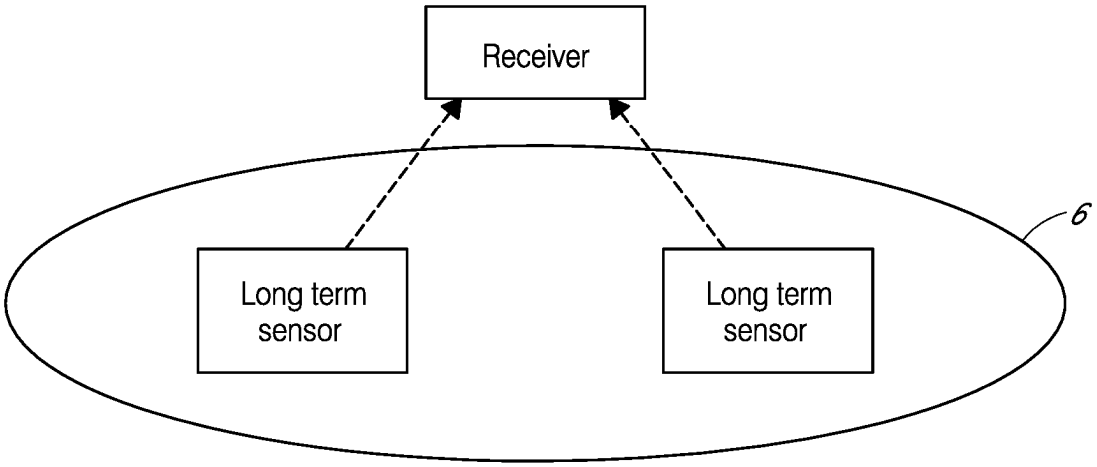

Receiver
Long term sensor
Long term sensor
6

FIG. 1E

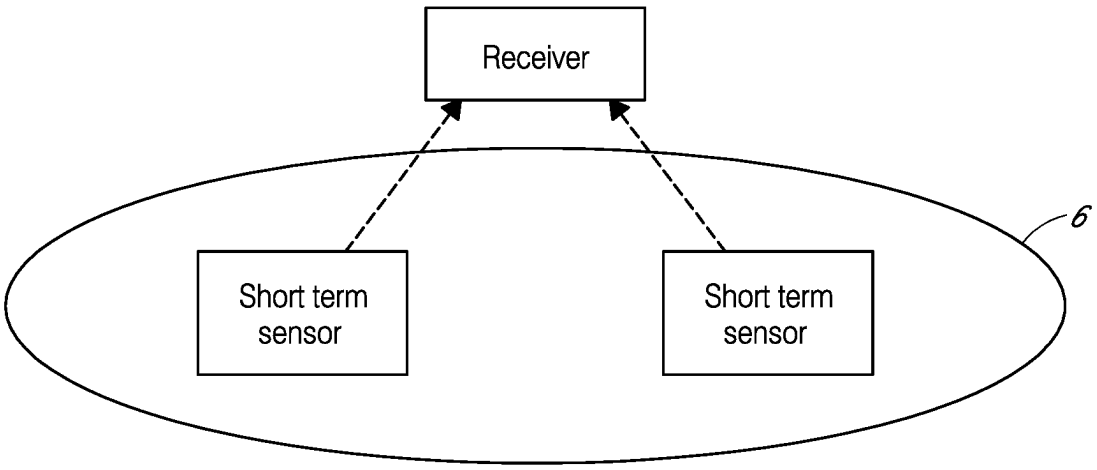

Receiver
Short term sensor
Short term sensor
6

FIG. 1F

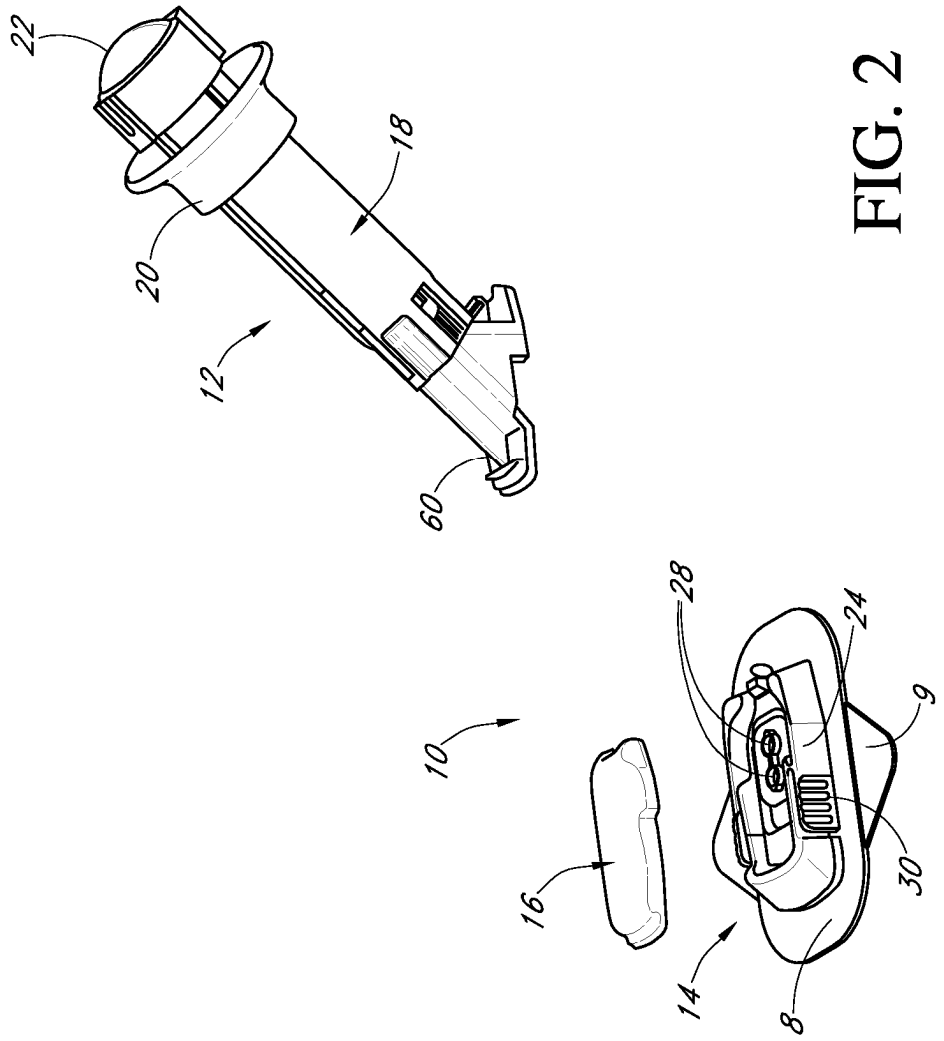


FIG. 2

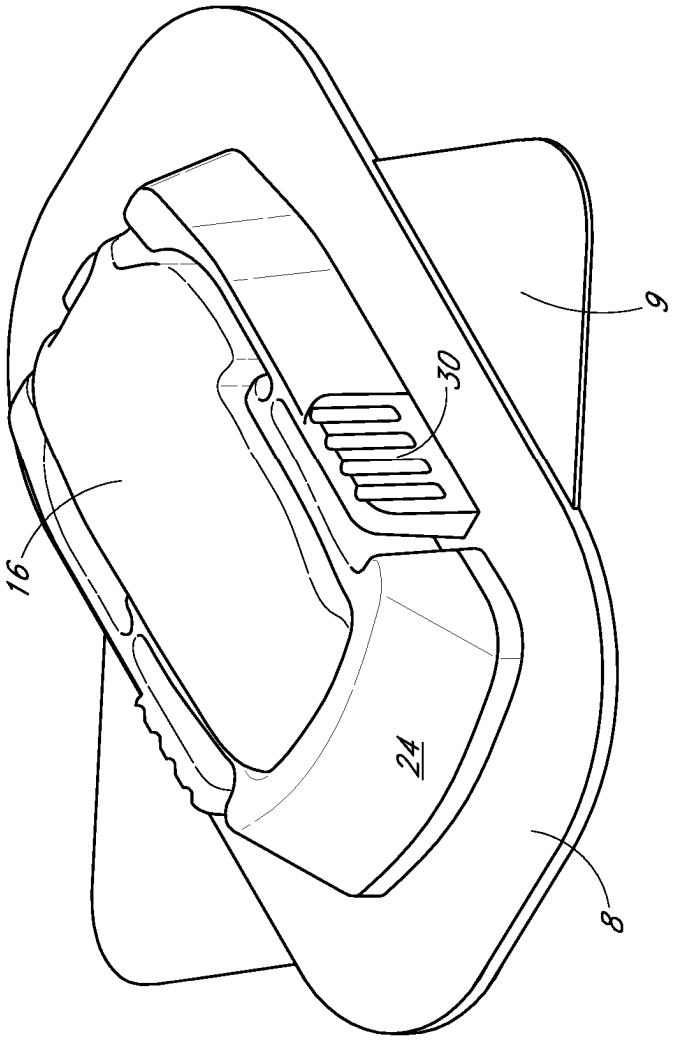


FIG. 3

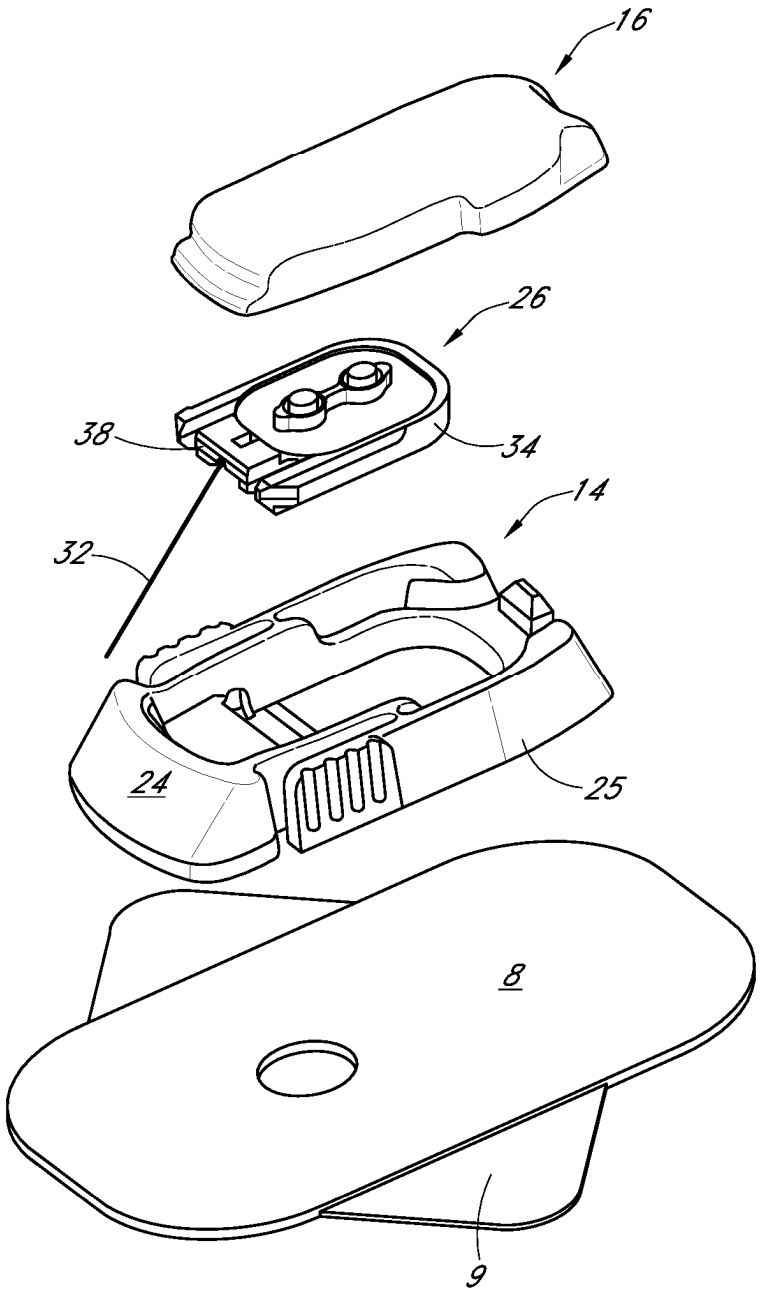


FIG. 4

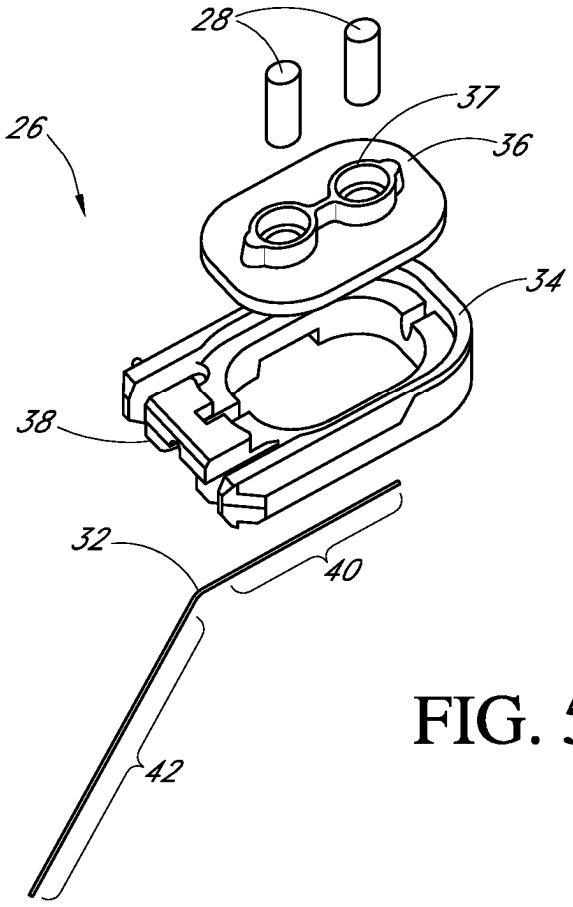


FIG. 5A

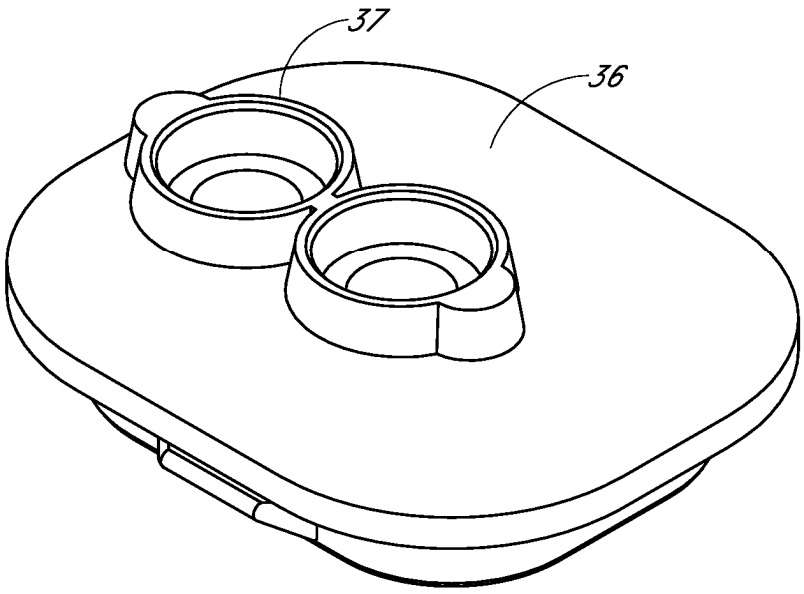


FIG. 5B

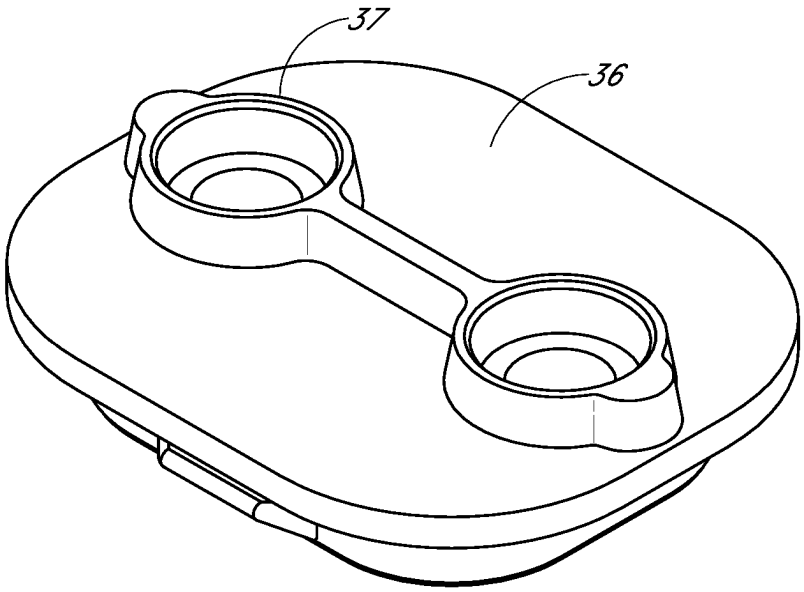


FIG. 5C

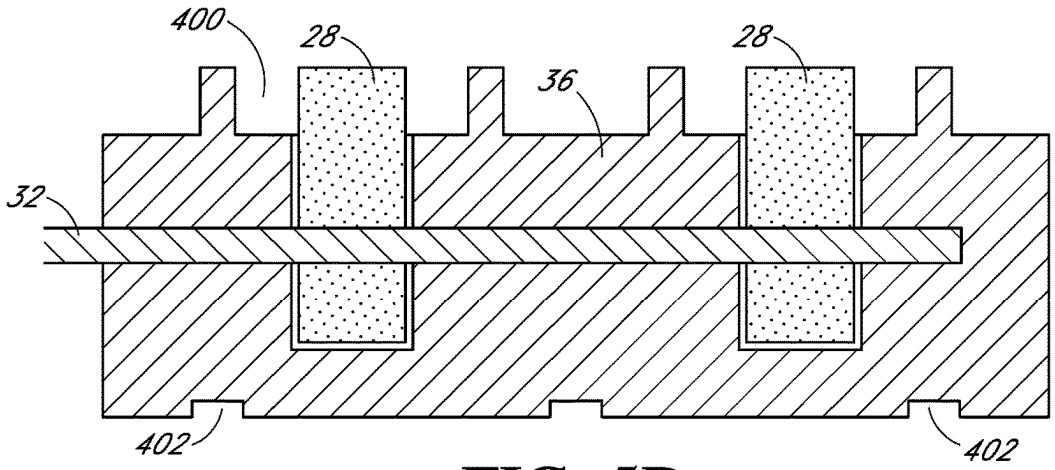


FIG. 5D

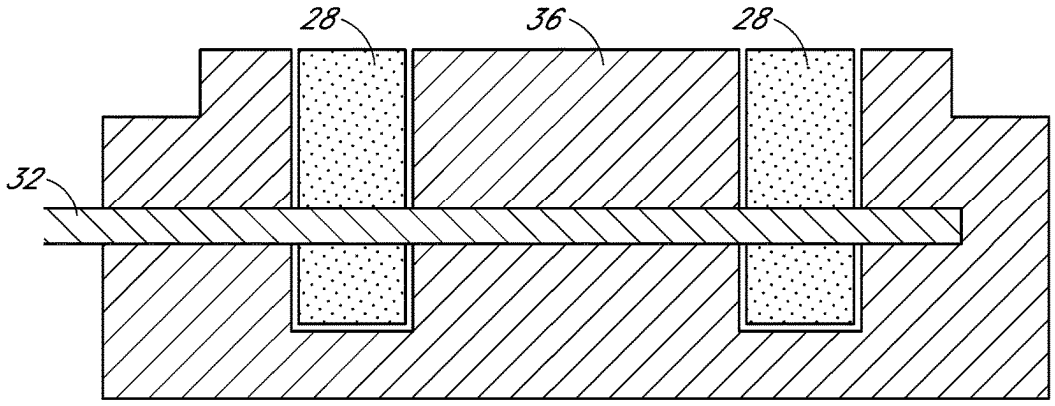


FIG. 5E

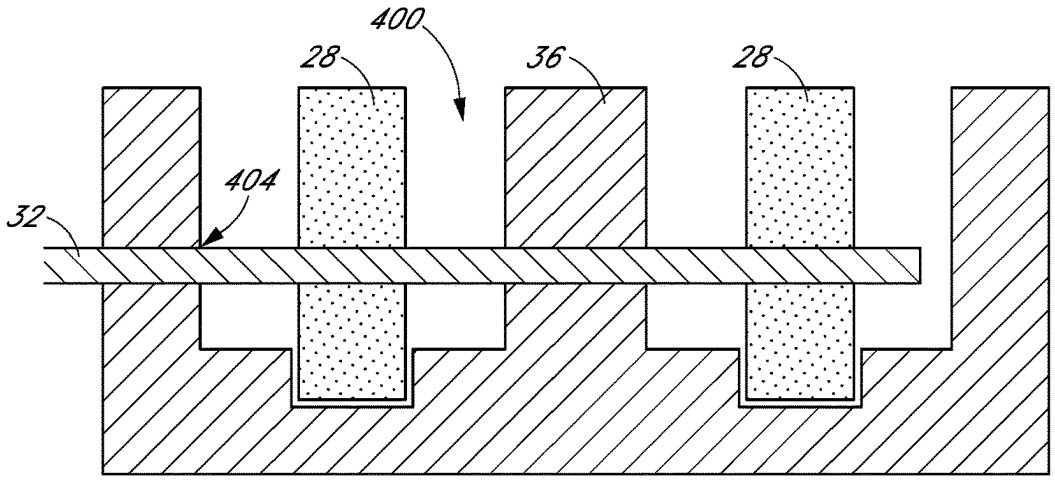


FIG. 5F

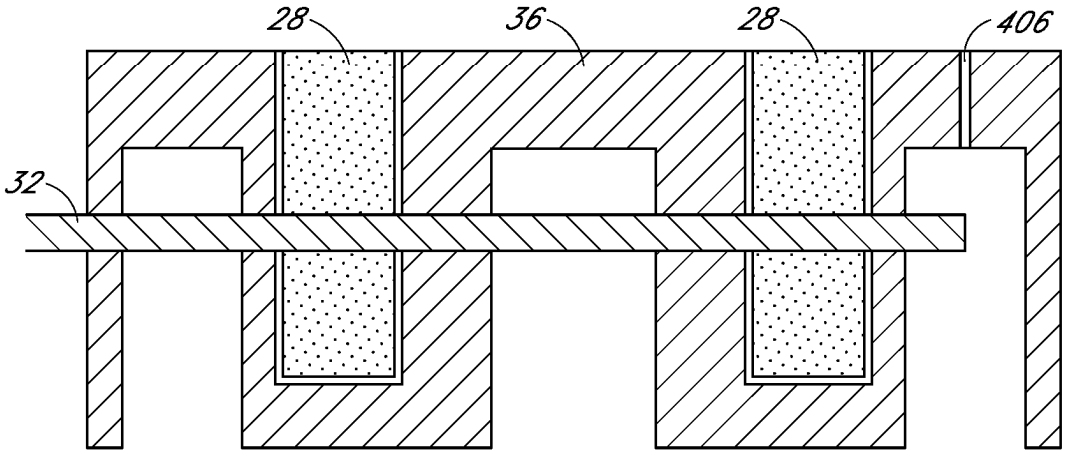


FIG. 5G

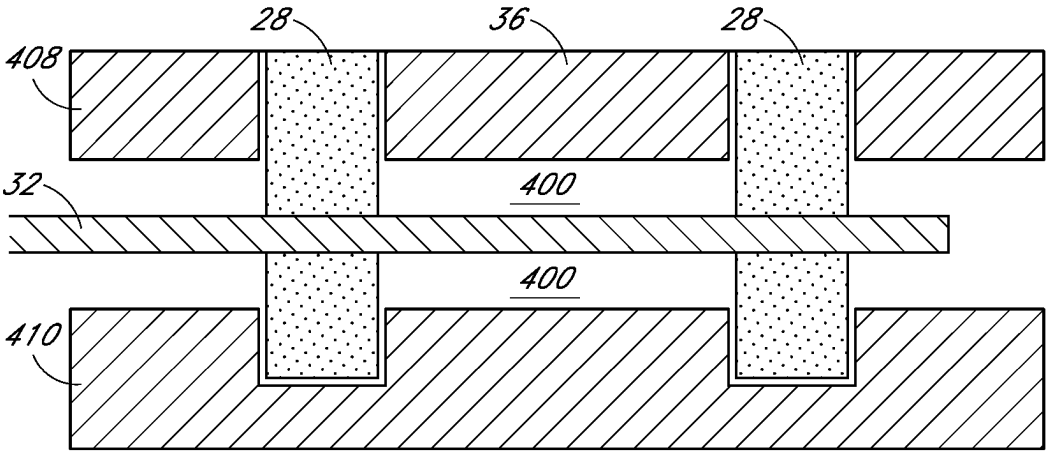


FIG. 5H

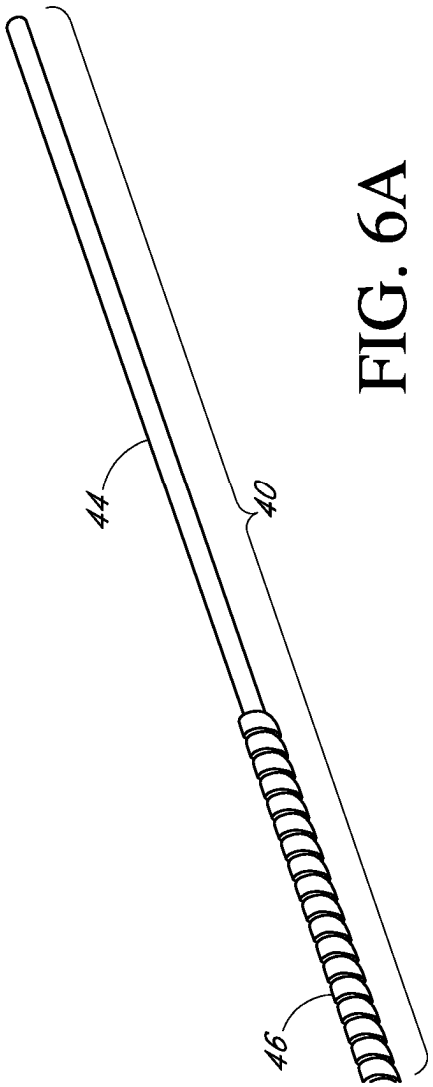


FIG. 6A

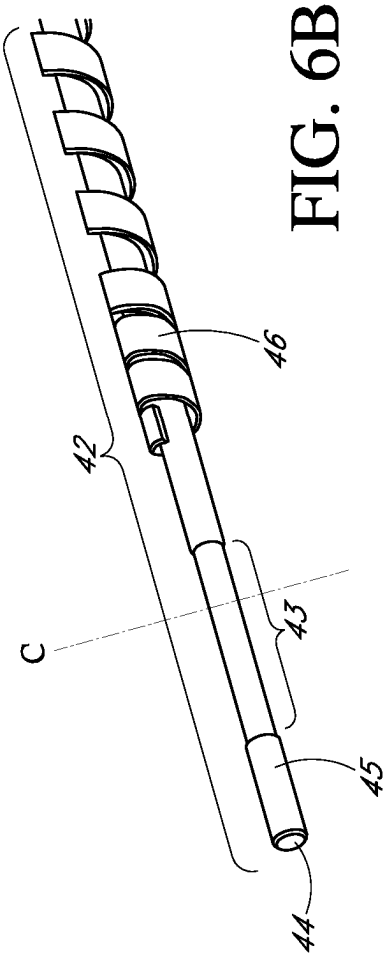


FIG. 6B

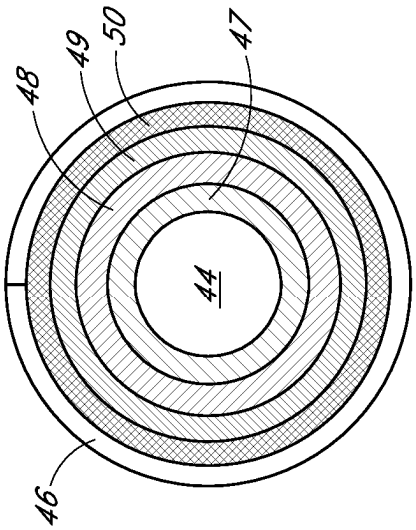


FIG. 6C

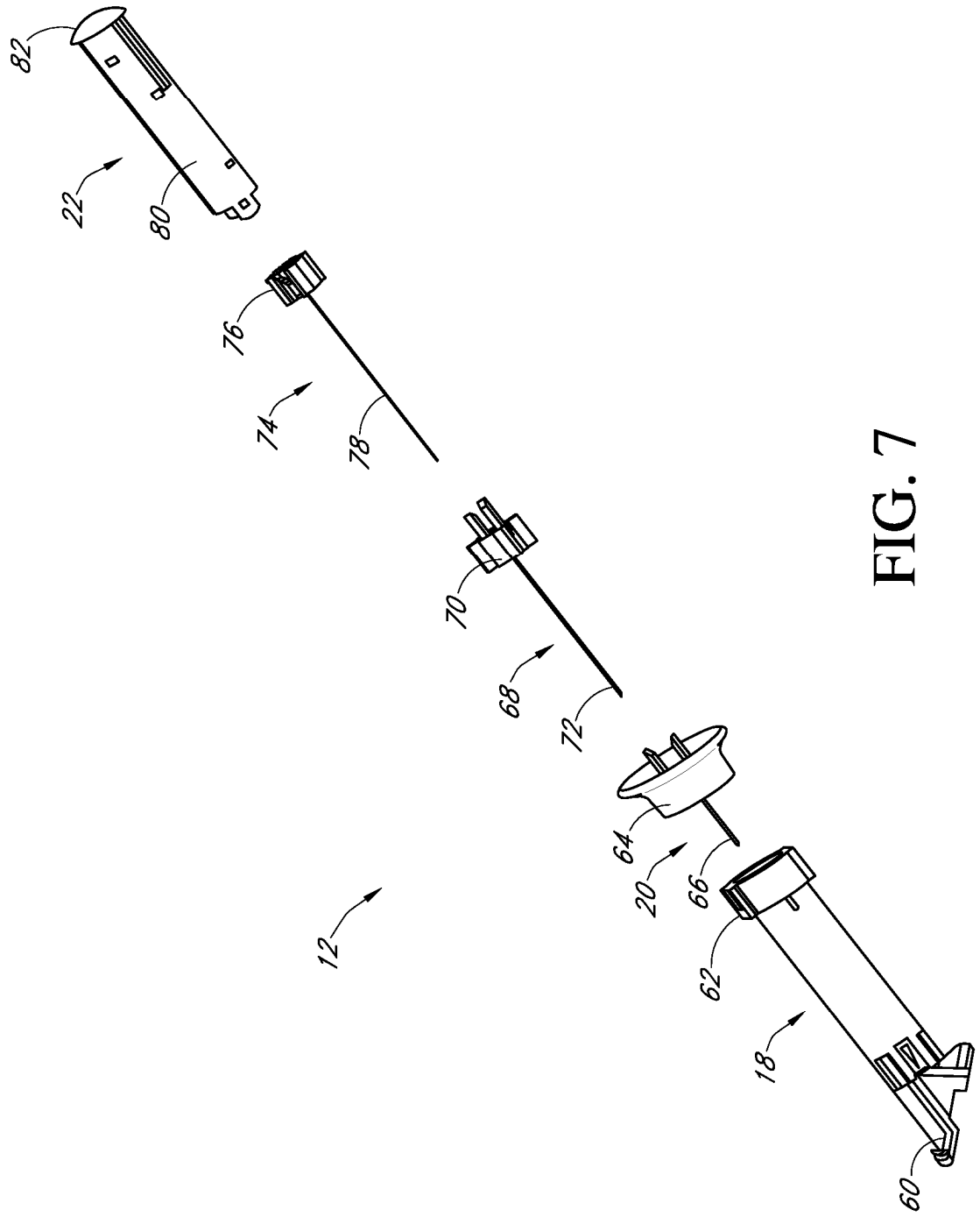


FIG. 7

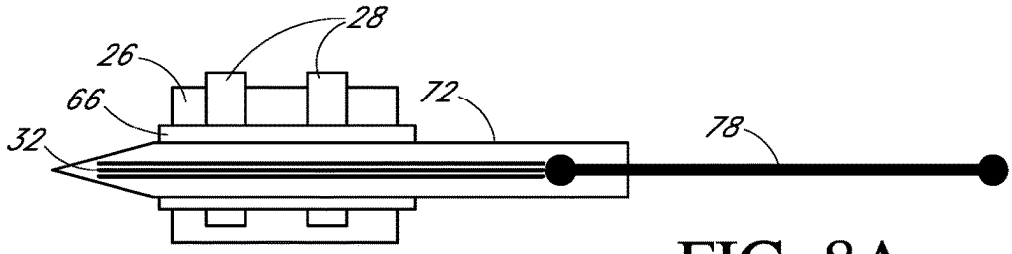


FIG. 8A

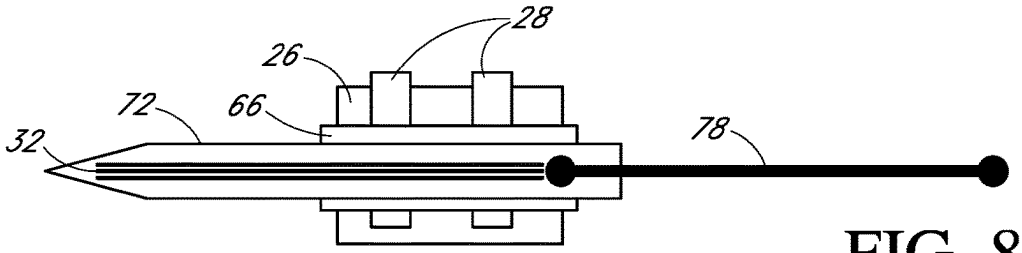


FIG. 8B

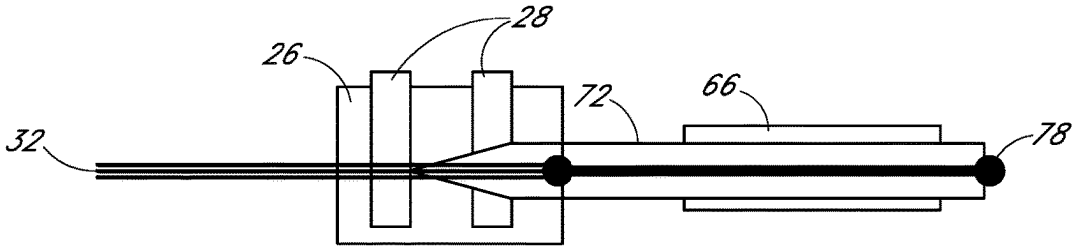


FIG. 8C

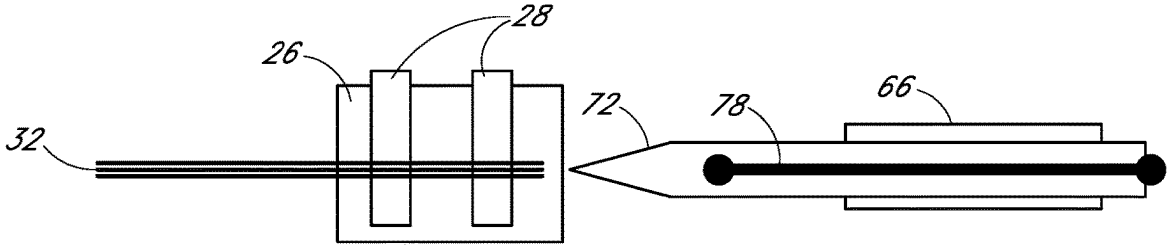


FIG. 8D

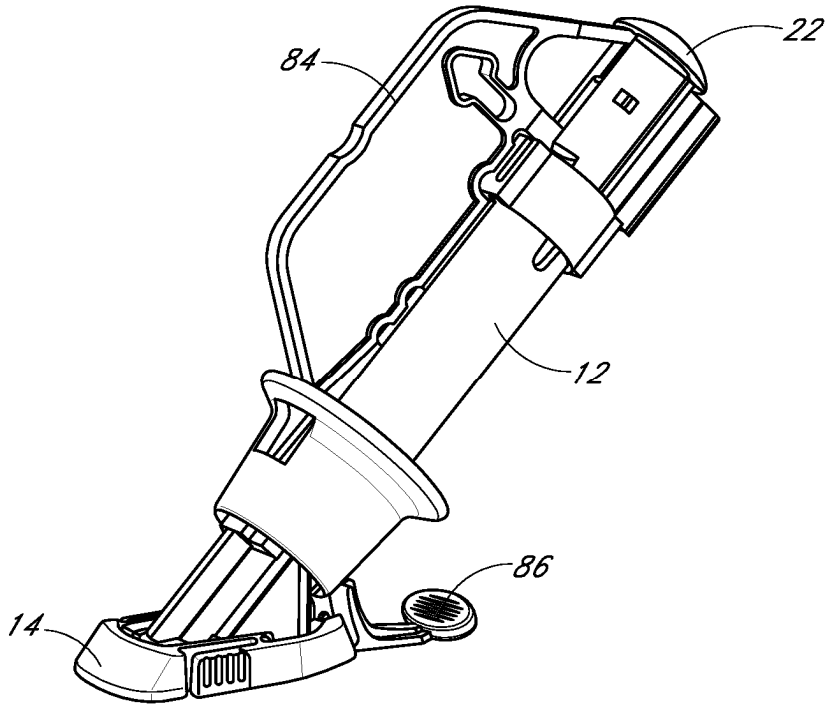


FIG. 9A

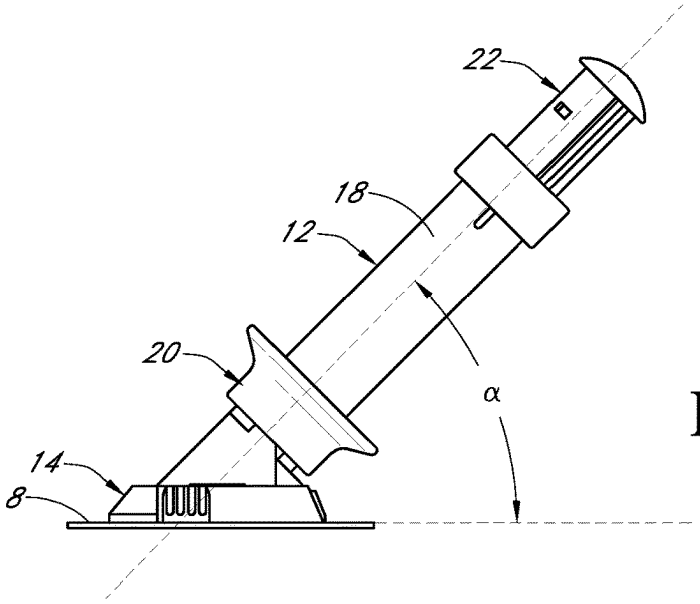


FIG. 9B

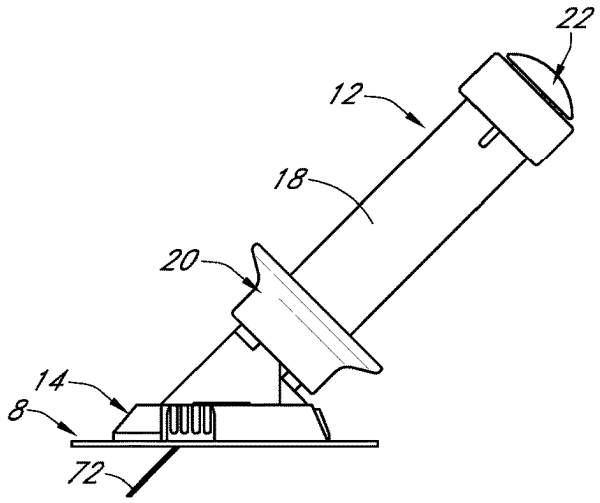


FIG. 9C

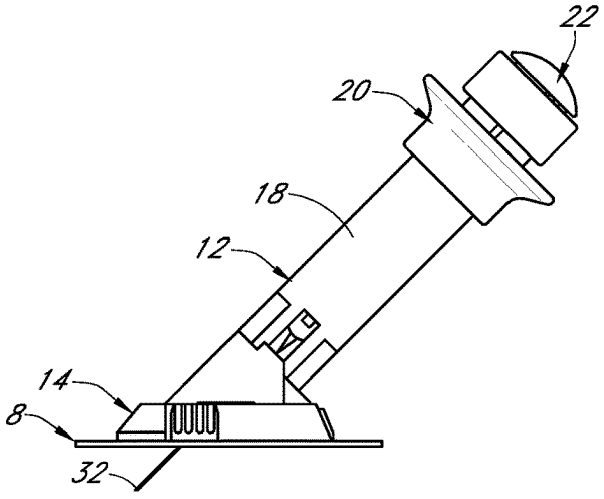


FIG. 9D

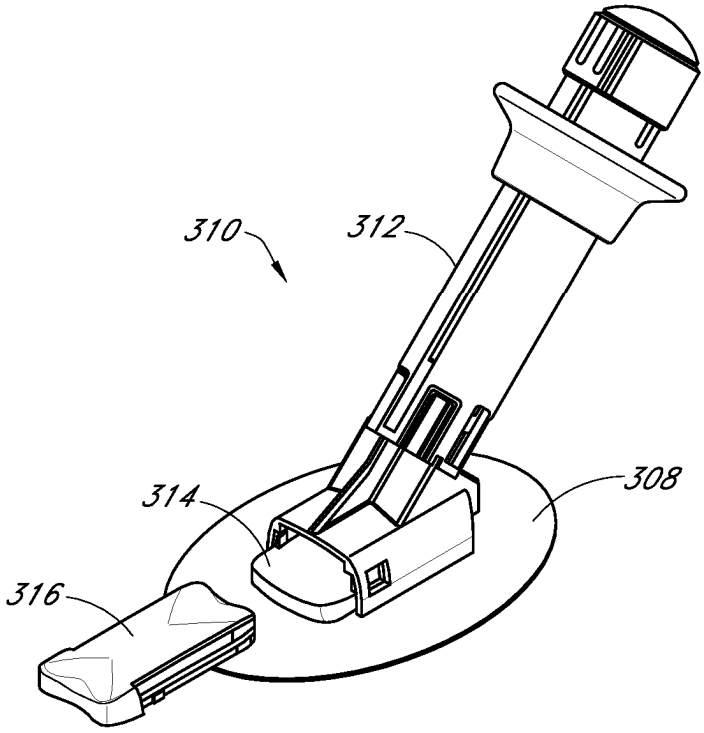


FIG. 9E

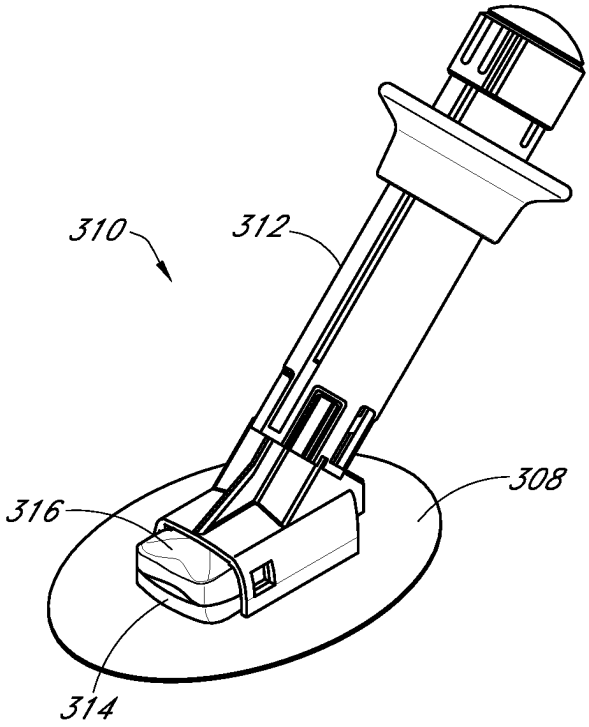


FIG. 9F

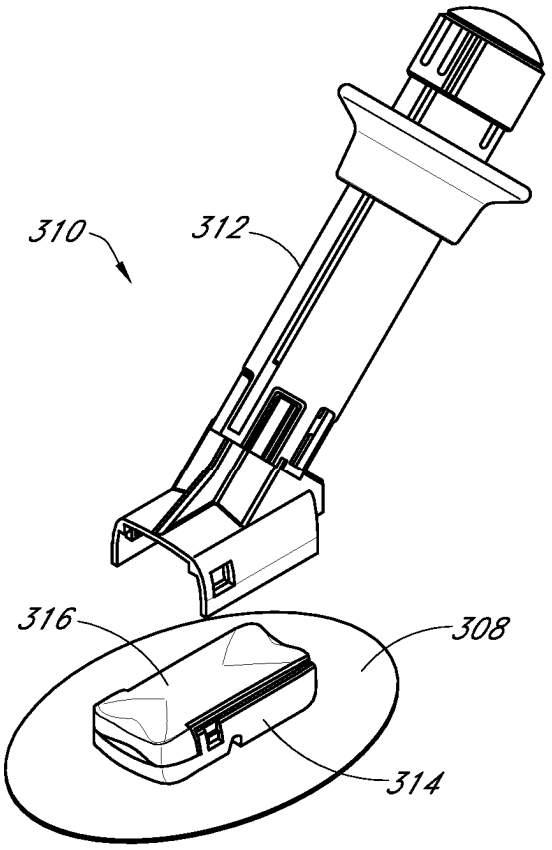


FIG. 9G

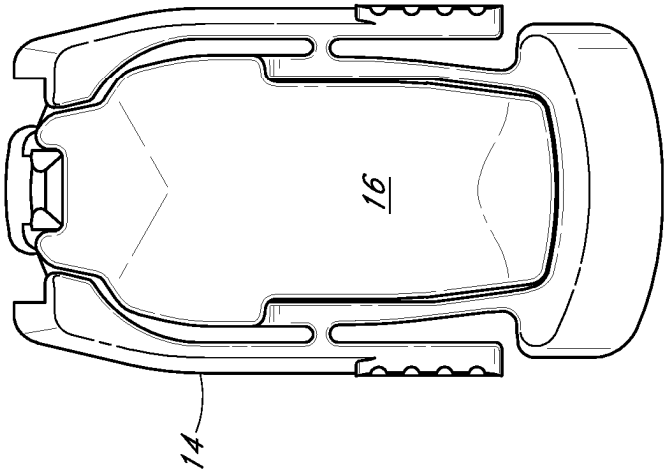


FIG. 9I

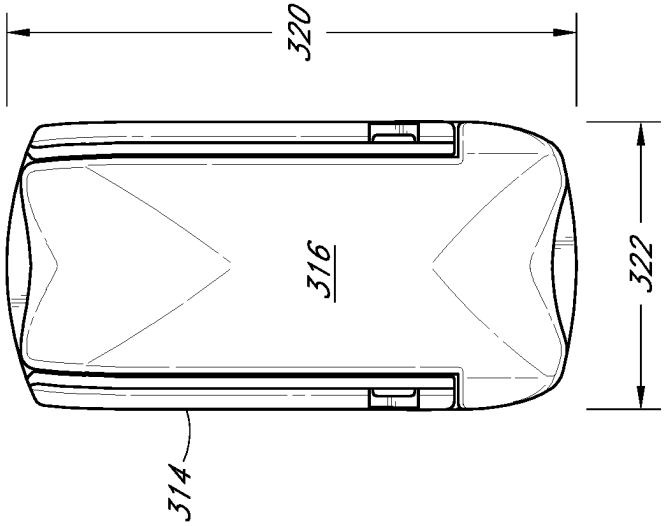


FIG. 9H

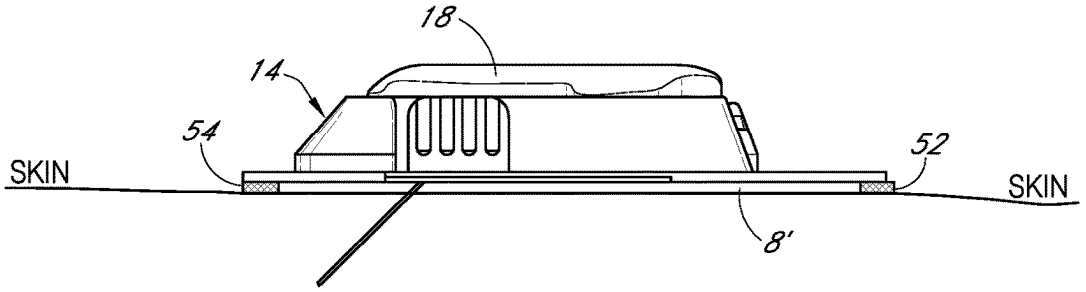


FIG. 10A

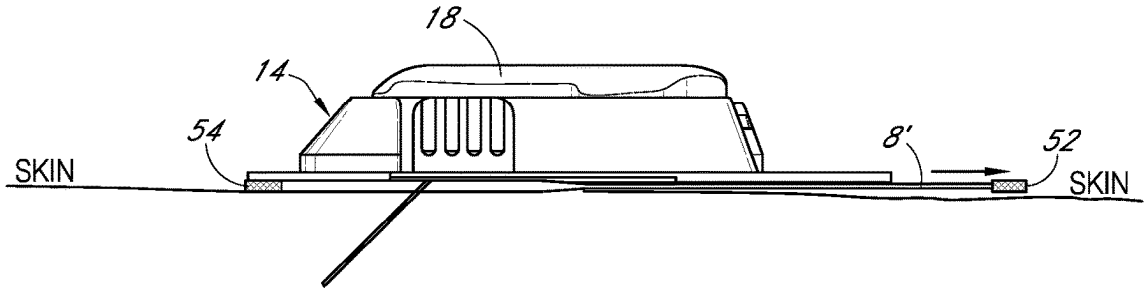


FIG. 10B

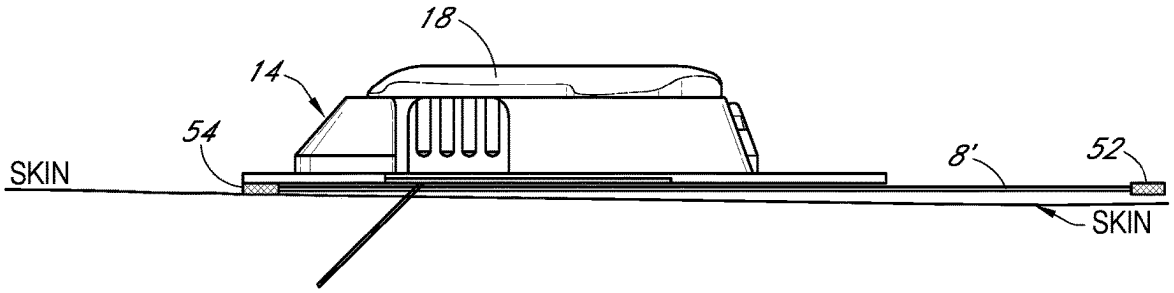


FIG. 10C

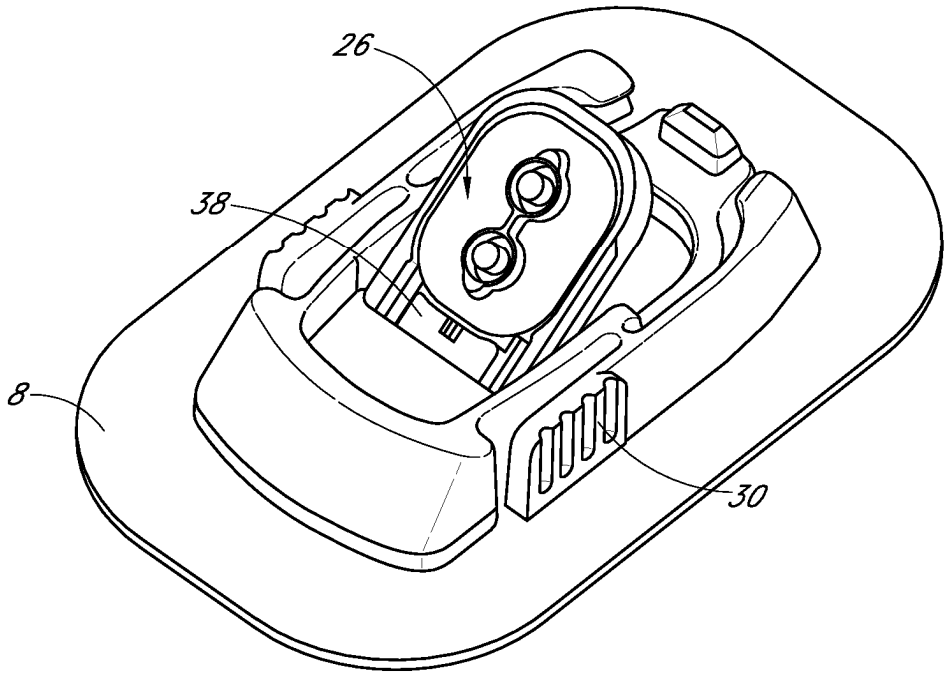


FIG. 11A

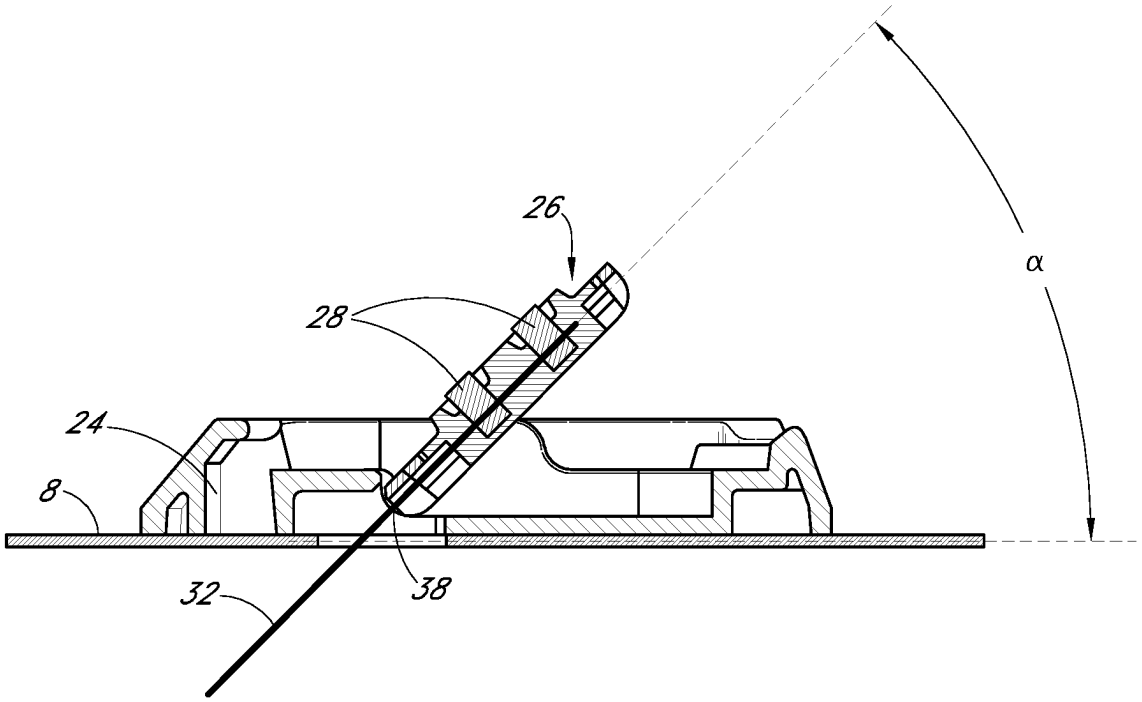


FIG. 11B

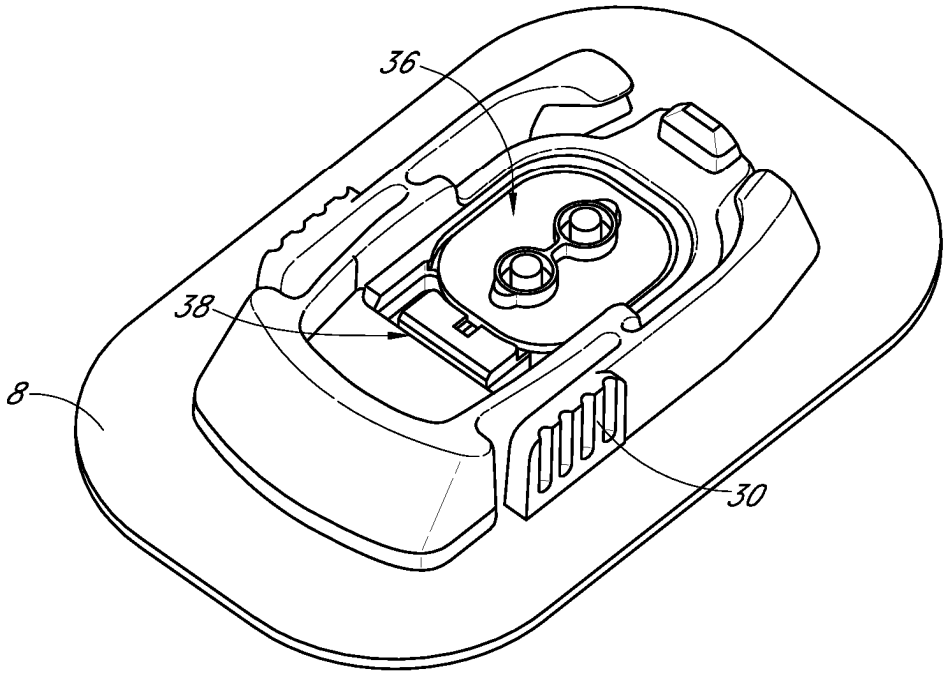


FIG. 12A

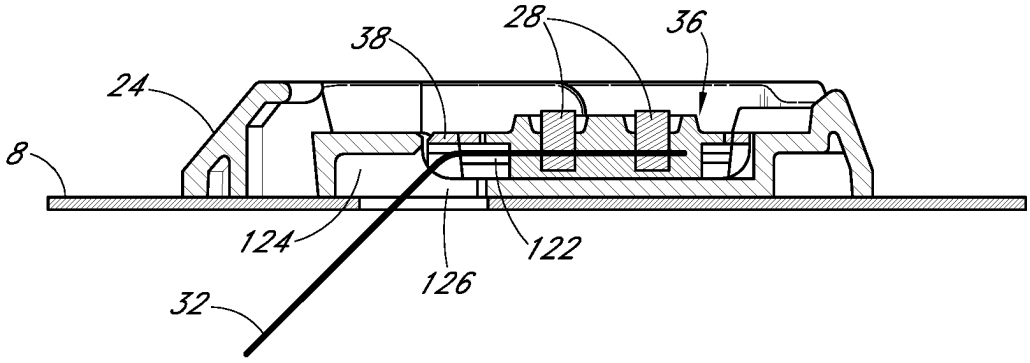


FIG. 12B

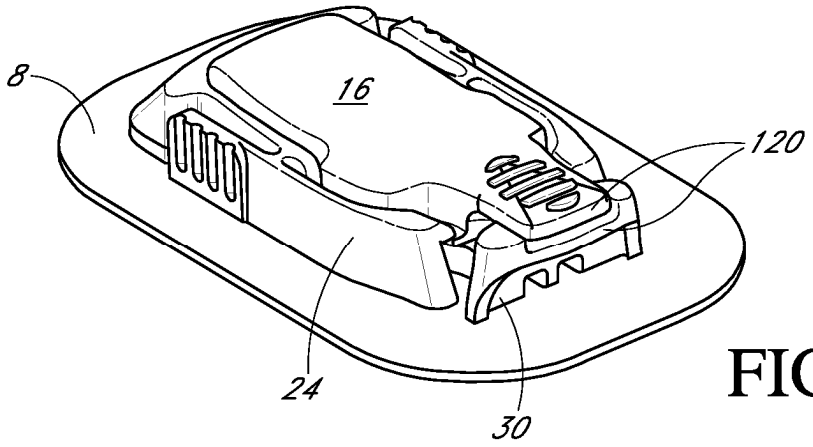


FIG. 13A

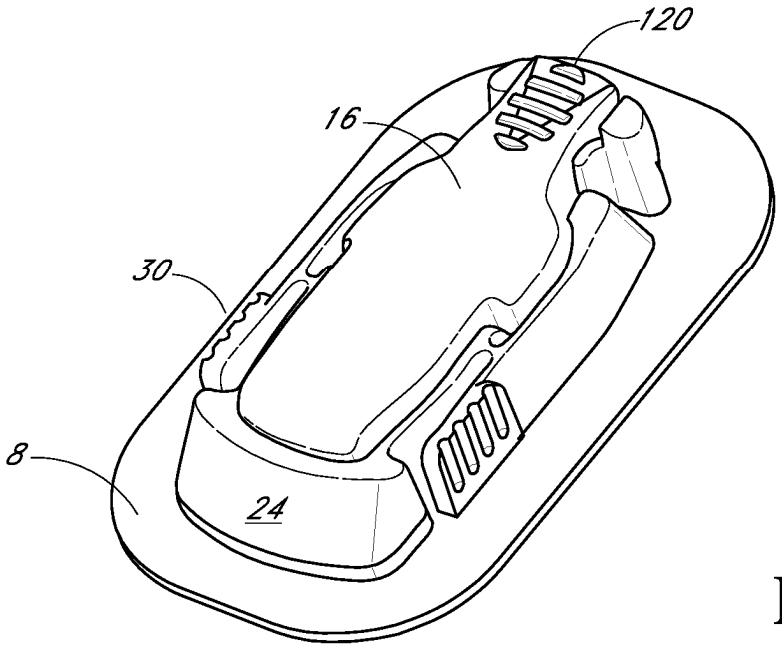


FIG. 13B

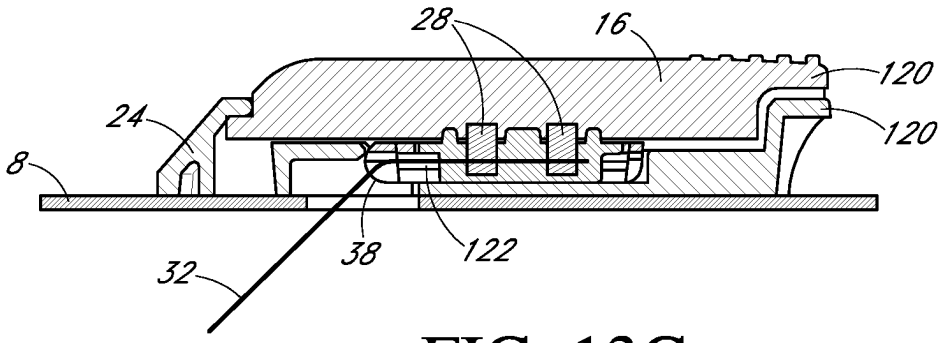


FIG. 13C

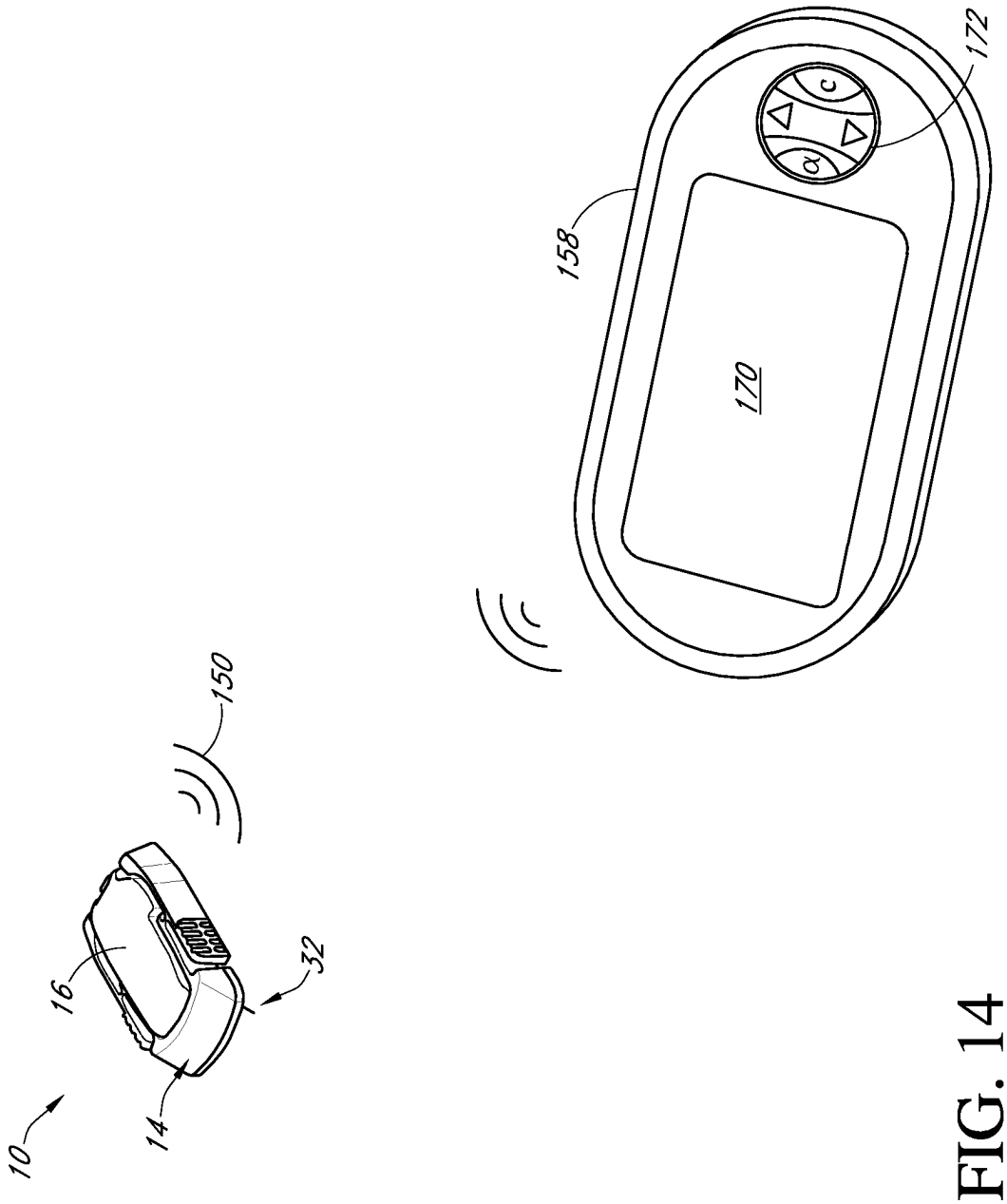


FIG. 14

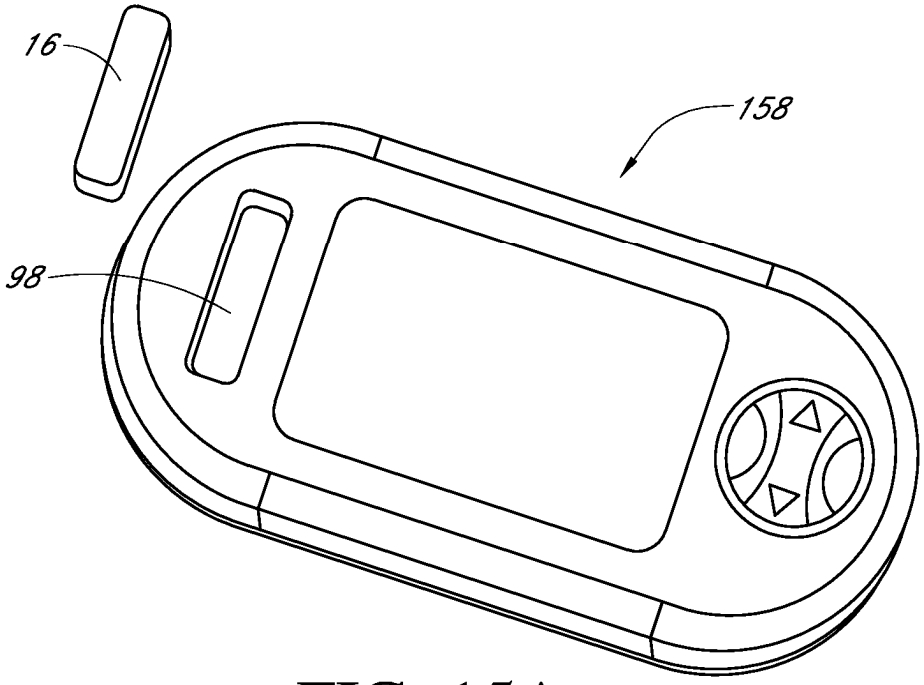


FIG. 15A

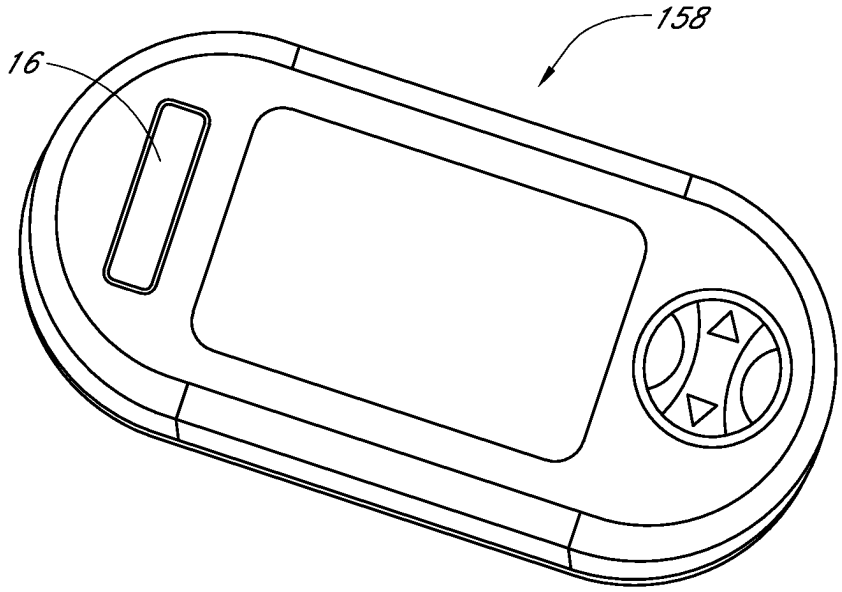


FIG. 15B

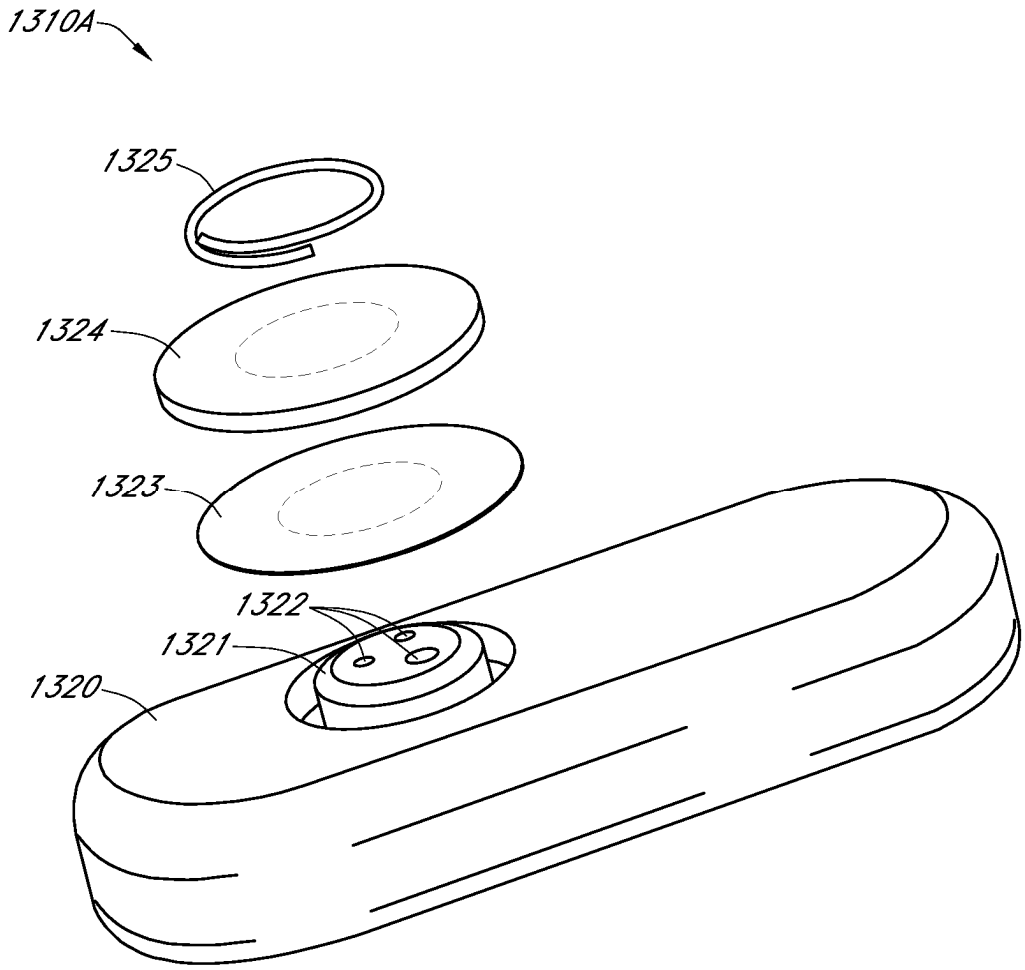


FIG. 16

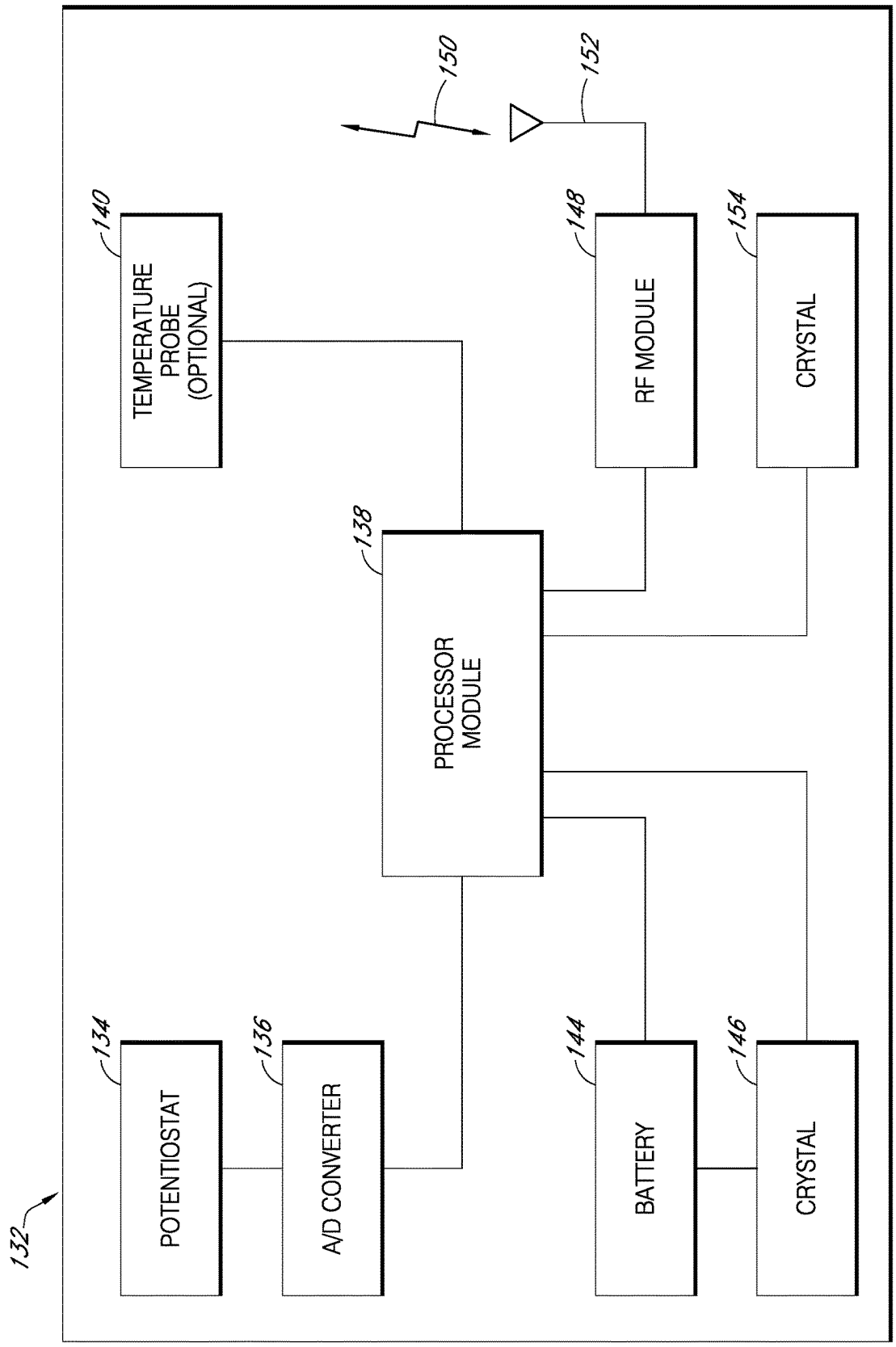

150
152
140
TEMPERATURE PROBE (OPTIONAL)
148
RF MODULE
154
CRYSTAL
138
PROCESSOR MODULE
134
POTENTIOSTAT
136
A/D CONVERTER
144
BATTERY
146
CRYSTAL
132

FIG. 17

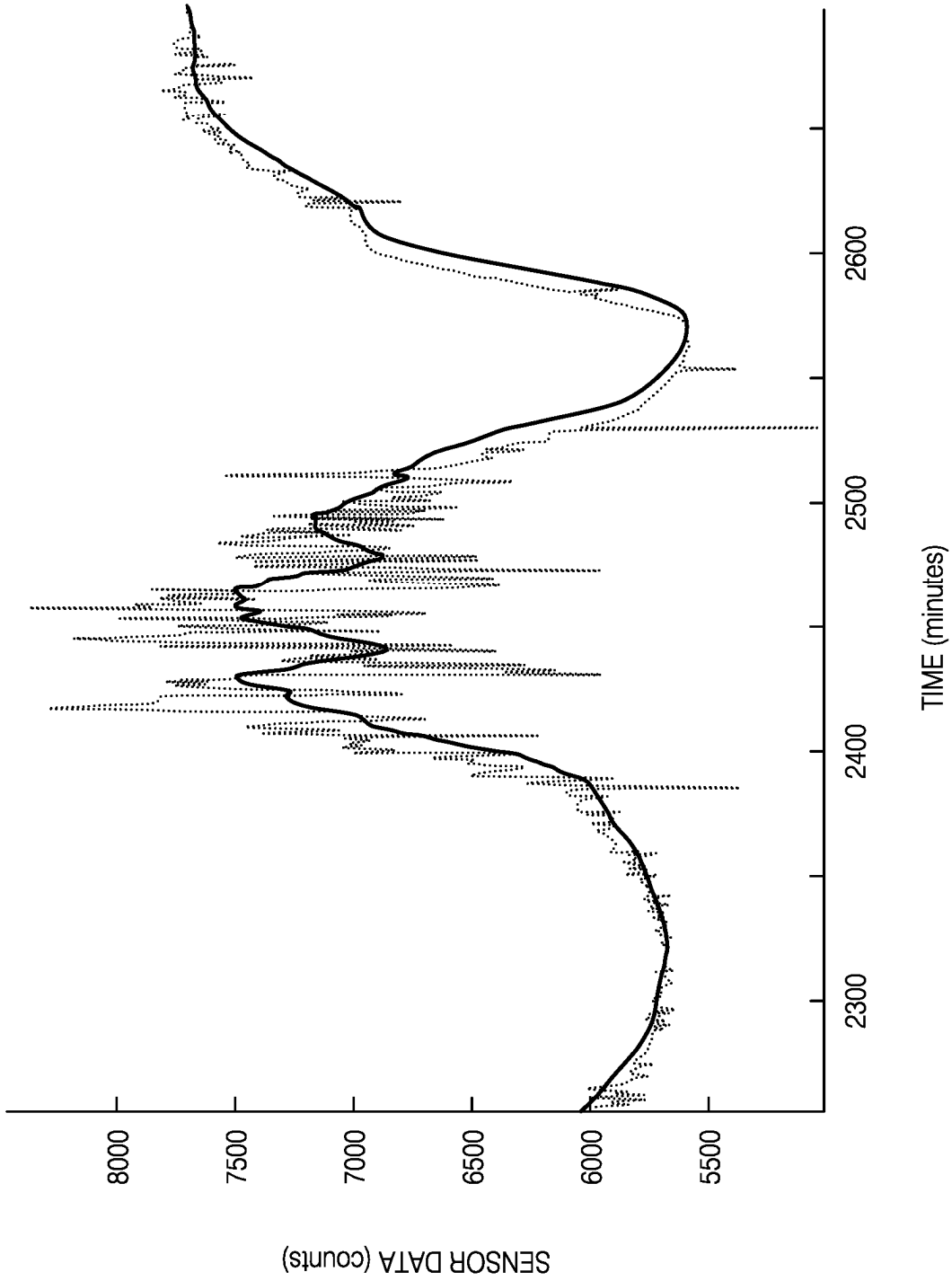

SENSOR DATA (counts)
8000
7500
7000
6500
6000
5500
TIME (minutes)
2300
2400
2500
2600

FIG. 18

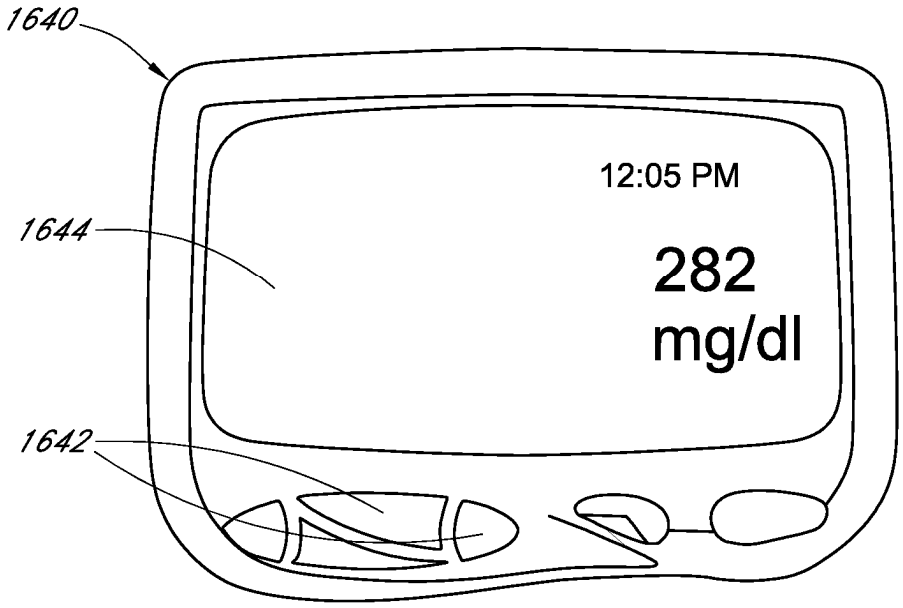


FIG. 19A

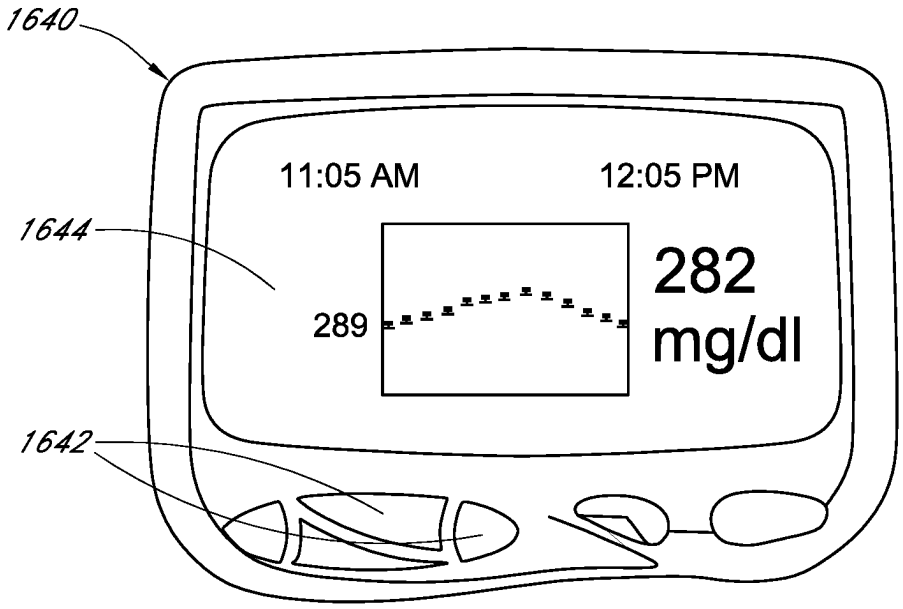


FIG. 19B

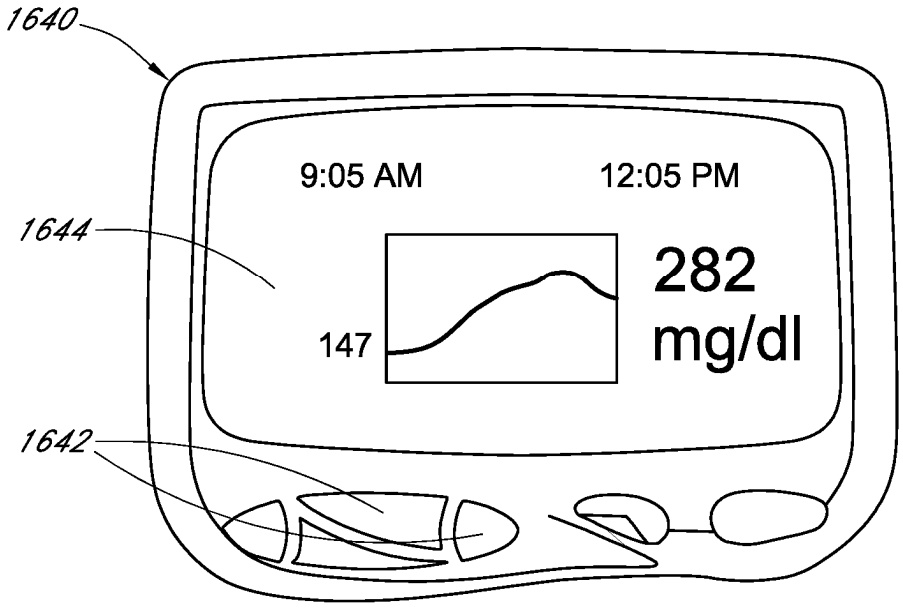


FIG. 19C

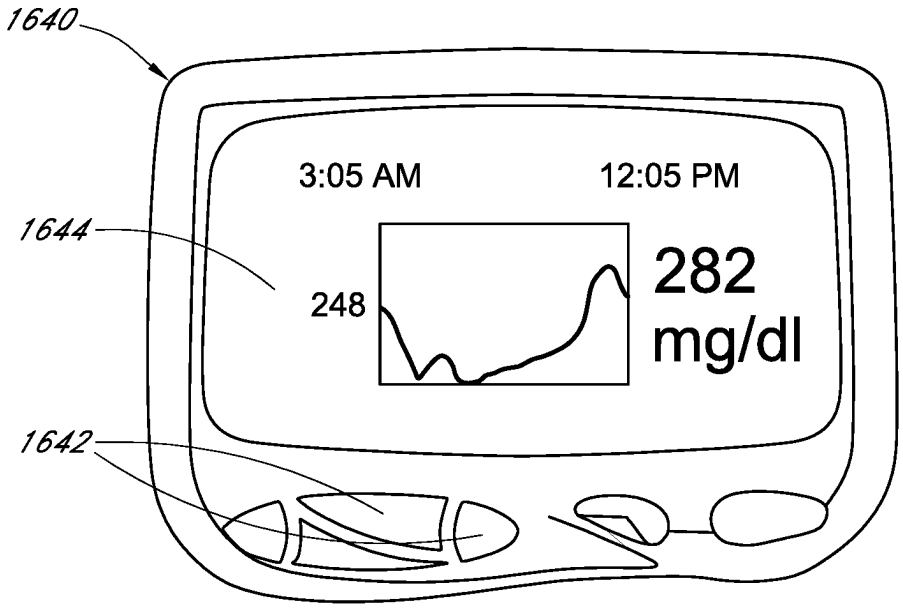


FIG. 19D

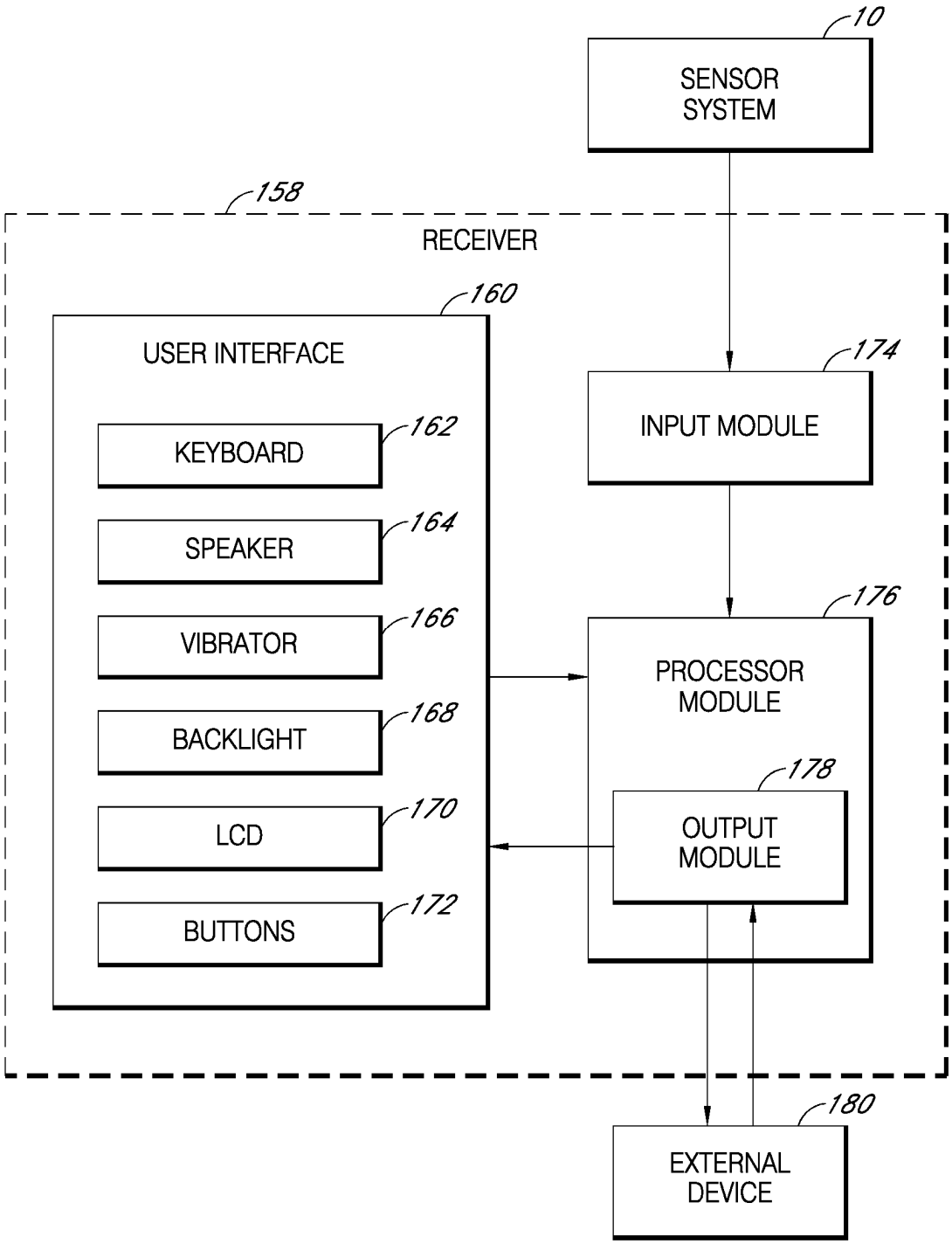

10
SENSOR SYSTEM
158
RECEIVER
160
USER INTERFACE
162
KEYBOARD
164
SPEAKER
166
VIBRATOR
168
BACKLIGHT
170
LCD
172
BUTTONS
174
INPUT MODULE
176
PROCESSOR MODULE
178
OUTPUT MODULE
180
EXTERNAL DEVICE

FIG. 20A

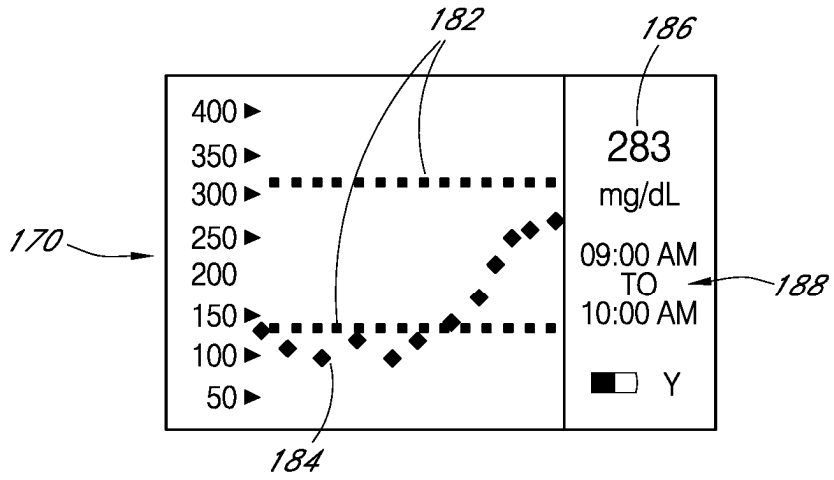


FIG. 20B

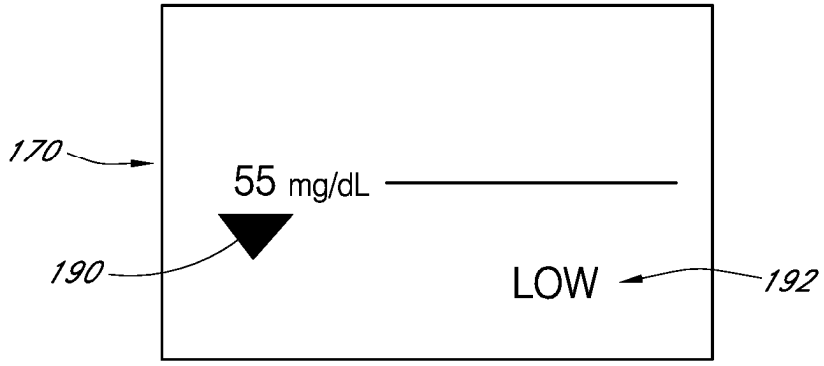


FIG. 20C

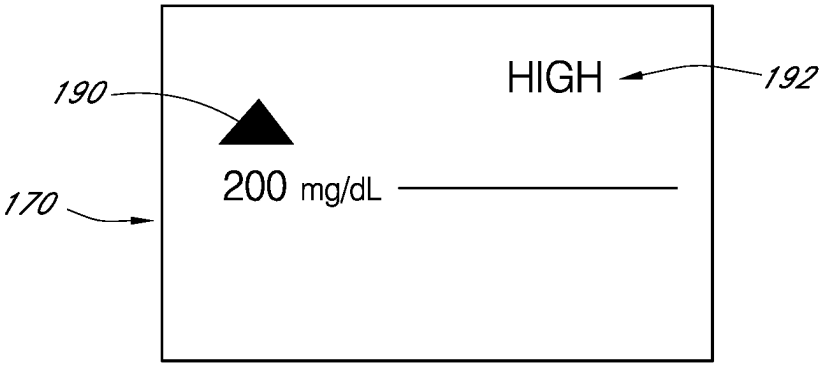


FIG. 20D

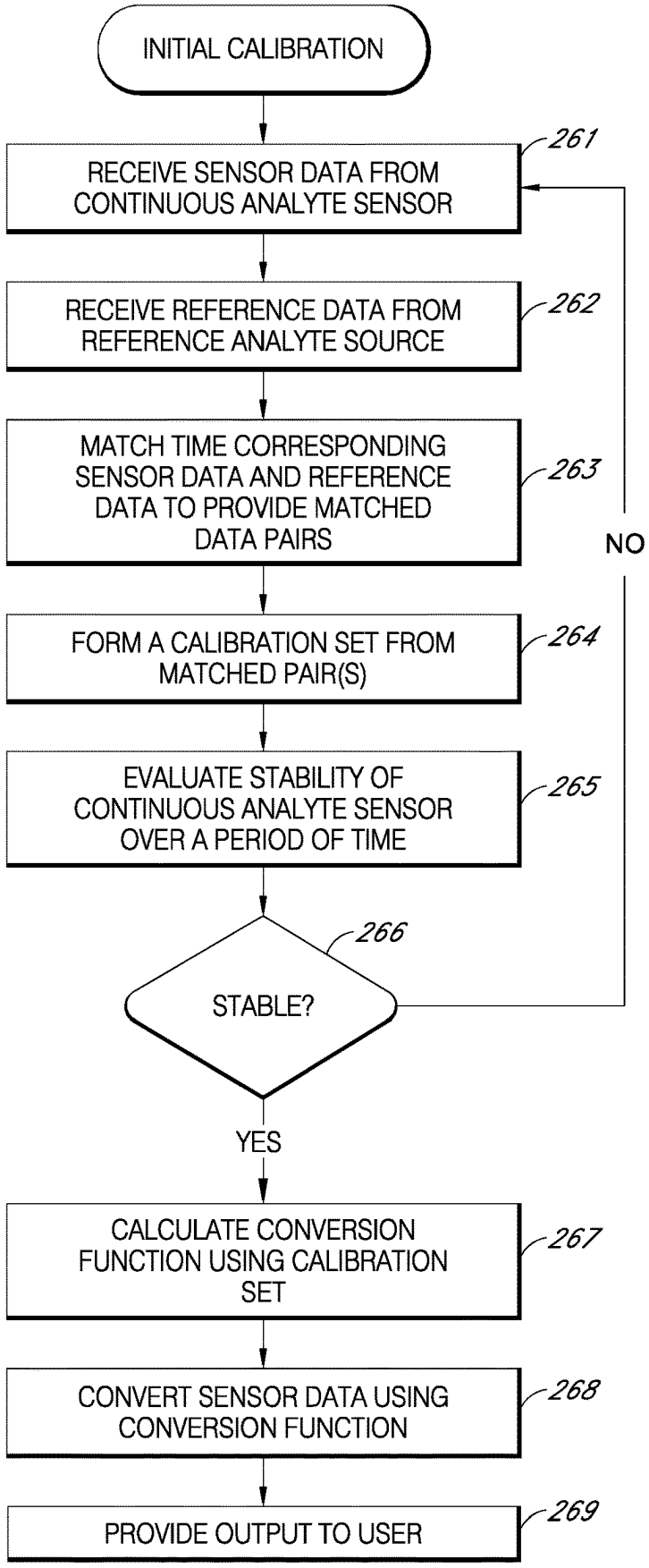

INITIAL CALIBRATION
RECEIVE SENSOR DATA FROM CONTINUOUS ANALYTE SENSOR
261
RECEIVE REFERENCE DATA FROM REFERENCE ANALYTE SOURCE
262
MATCH TIME CORRESPONDING SENSOR DATA AND REFERENCE DATA TO PROVIDE MATCHED DATA PAIRS
263
FORM A CALIBRATION SET FROM MATCHED PAIR(S)
264
EVALUATE STABILITY OF CONTINUOUS ANALYTE SENSOR OVER A PERIOD OF TIME
265
STABLE?
266
NO
YES
CALCULATE CONVERSION FUNCTION USING CALIBRATION SET
267
CONVERT SENSOR DATA USING CONVERSION FUNCTION
268
PROVIDE OUTPUT TO USER
269

FIG. 21

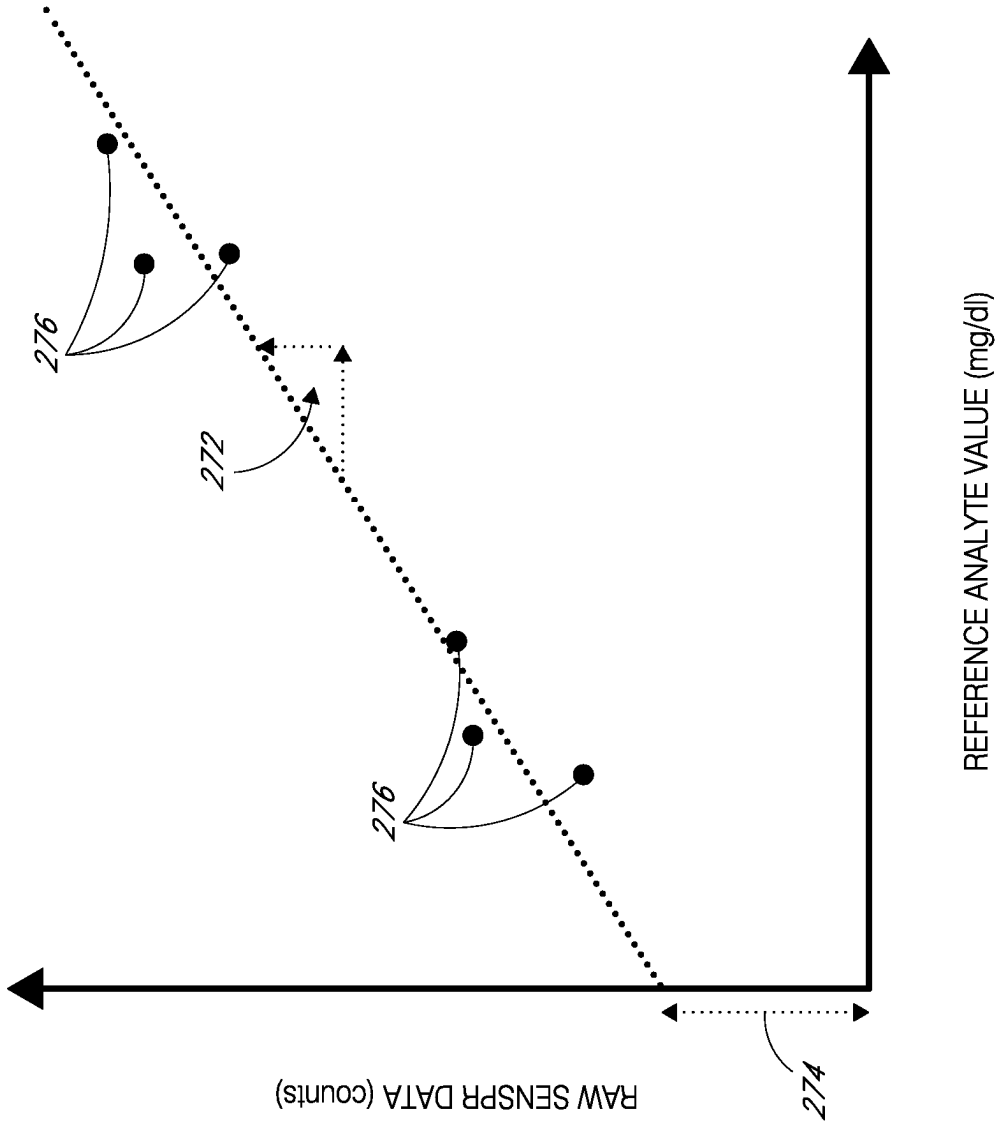

276
272
276
274
RAW SENSPR DATA (counts)
REFERENCE ANALYTE VALUE (mg/dl)

FIG. 22A

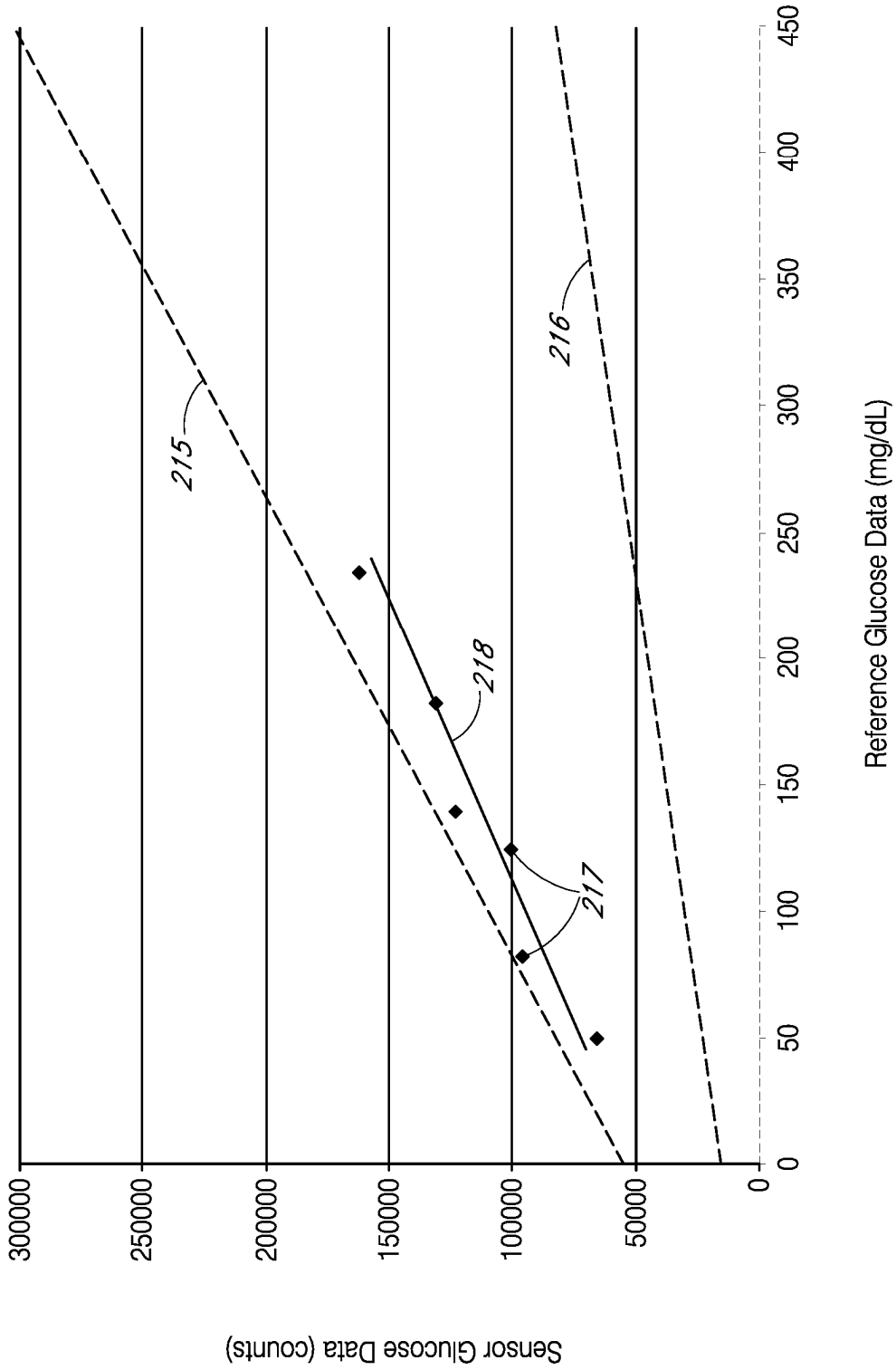

Sensor Glucose Data (counts)
300000
250000
200000
150000
100000
50000
0
Reference Glucose Data (mg/dL)
0
50
100
150
200
250
300
350
400
450
215
216
217
218

FIG. 22B

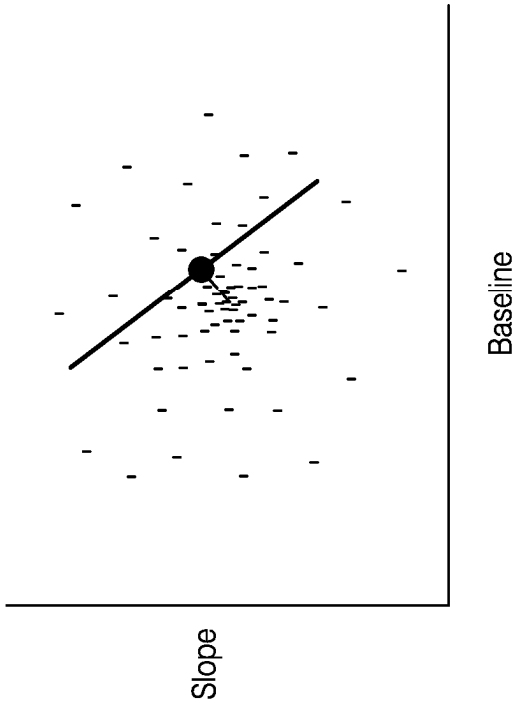


FIG. 22C

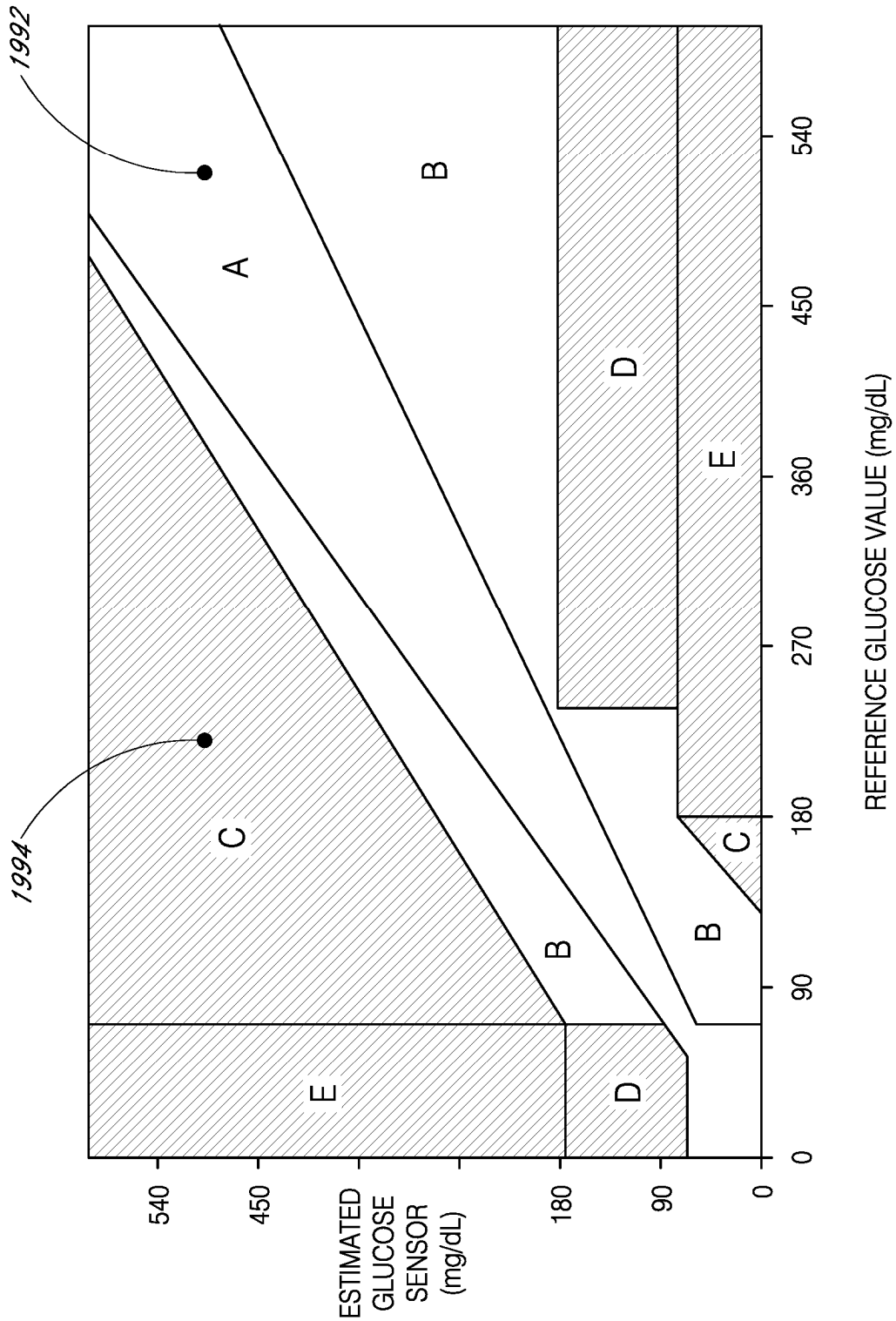

1992
1994
A
B
C
D
E
ESTIMATED GLUCOSE SENSOR (mg/dL)
REFERENCE GLUCOSE VALUE (mg/dL)
0
90
180
270
360
450
540

FIG. 23

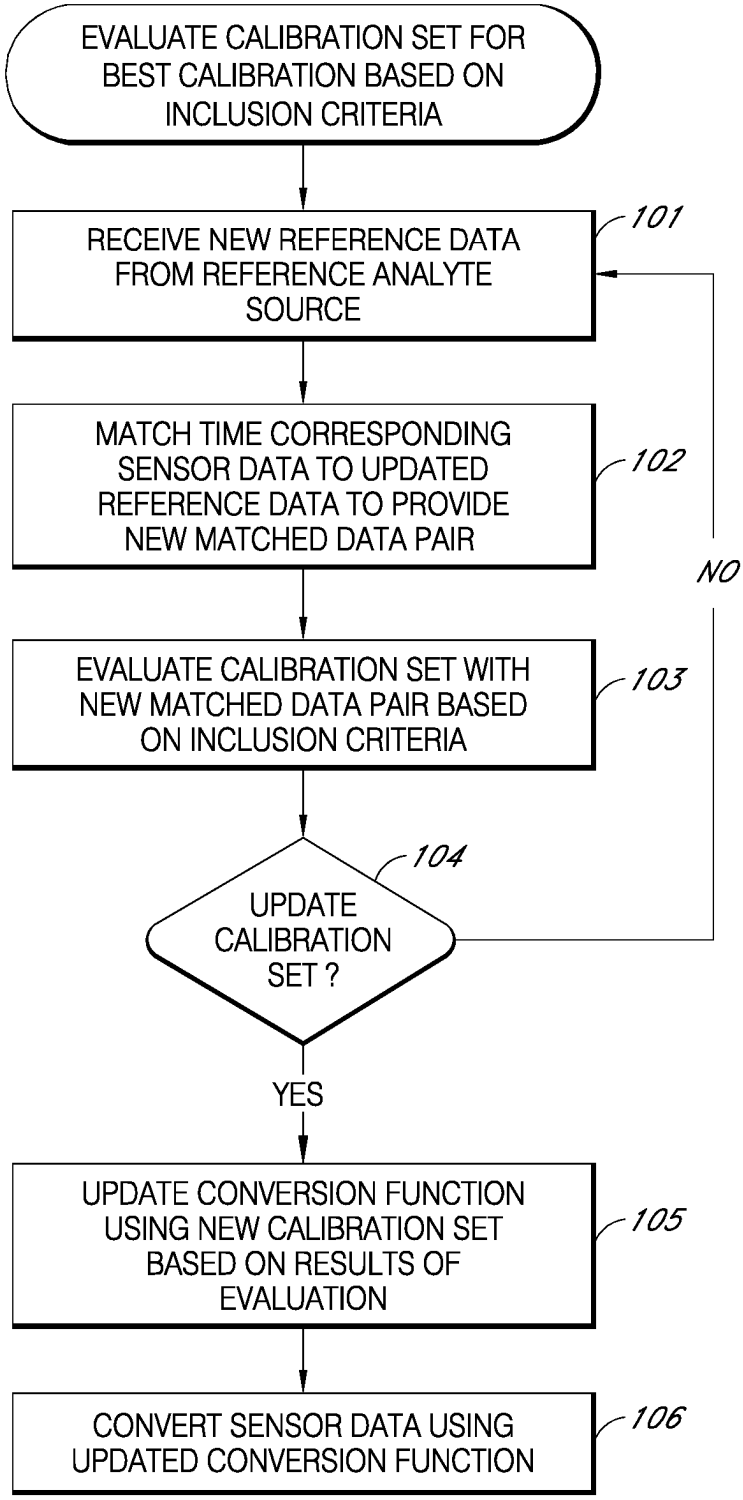

EVALUATE CALIBRATION SET FOR BEST CALIBRATION BASED ON INCLUSION CRITERIA
RECEIVE NEW REFERENCE DATA FROM REFERENCE ANALYTE SOURCE
101
MATCH TIME CORRESPONDING SENSOR DATA TO UPDATED REFERENCE DATA TO PROVIDE NEW MATCHED DATA PAIR
102
EVALUATE CALIBRATION SET WITH NEW MATCHED DATA PAIR BASED ON INCLUSION CRITERIA
103
104
UPDATE CALIBRATION SET ?
NO
YES
UPDATE CONVERSION FUNCTION USING NEW CALIBRATION SET BASED ON RESULTS OF EVALUATION
105
CONVERT SENSOR DATA USING UPDATED CONVERSION FUNCTION
106

FIG. 24

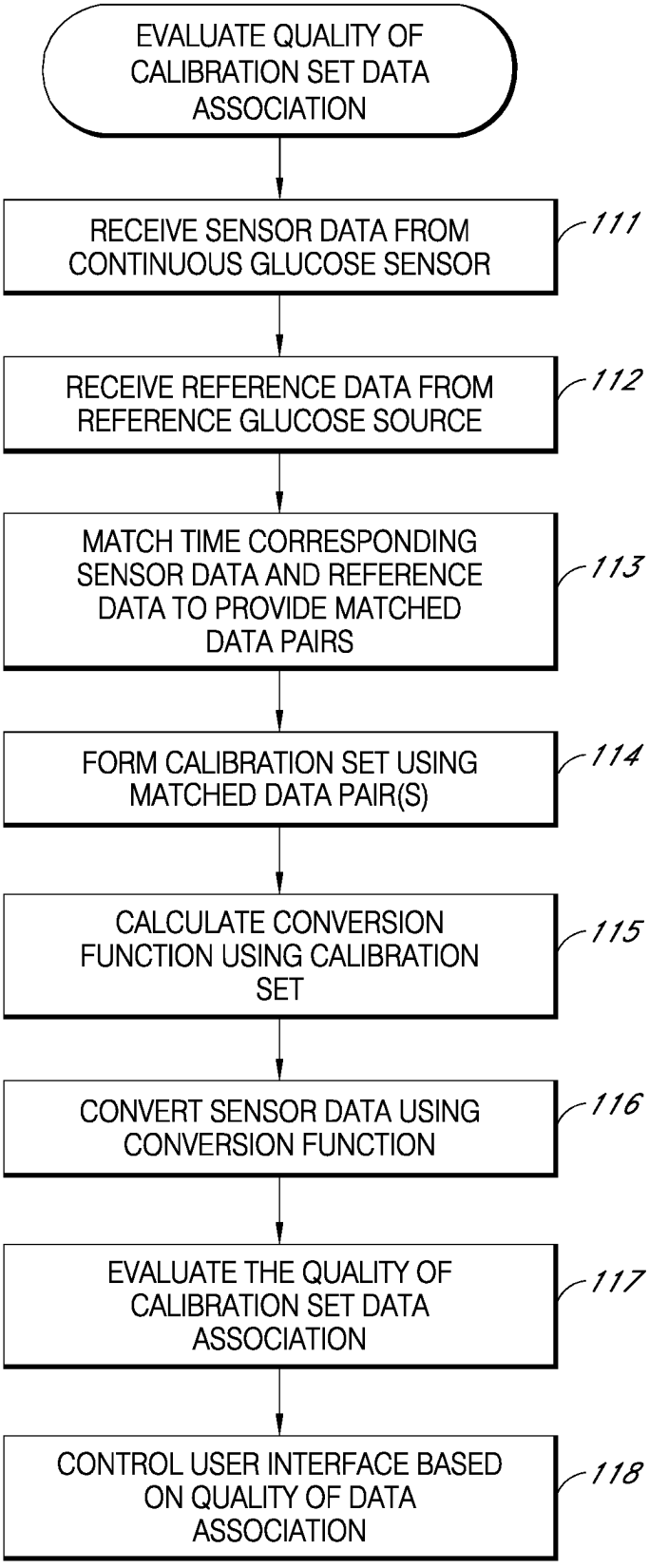

EVALUATE QUALITY OF CALIBRATION SET DATA ASSOCIATION
RECEIVE SENSOR DATA FROM CONTINUOUS GLUCOSE SENSOR
111
RECEIVE REFERENCE DATA FROM REFERENCE GLUCOSE SOURCE
112
MATCH TIME CORRESPONDING SENSOR DATA AND REFERENCE DATA TO PROVIDE MATCHED DATA PAIRS
113
FORM CALIBRATION SET USING MATCHED DATA PAIR(S)
114
CALCULATE CONVERSION FUNCTION USING CALIBRATION SET
115
CONVERT SENSOR DATA USING CONVERSION FUNCTION
116
EVALUATE THE QUALITY OF CALIBRATION SET DATA ASSOCIATION
117
CONTROL USER INTERFACE BASED ON QUALITY OF DATA ASSOCIATION
118

FIG. 25

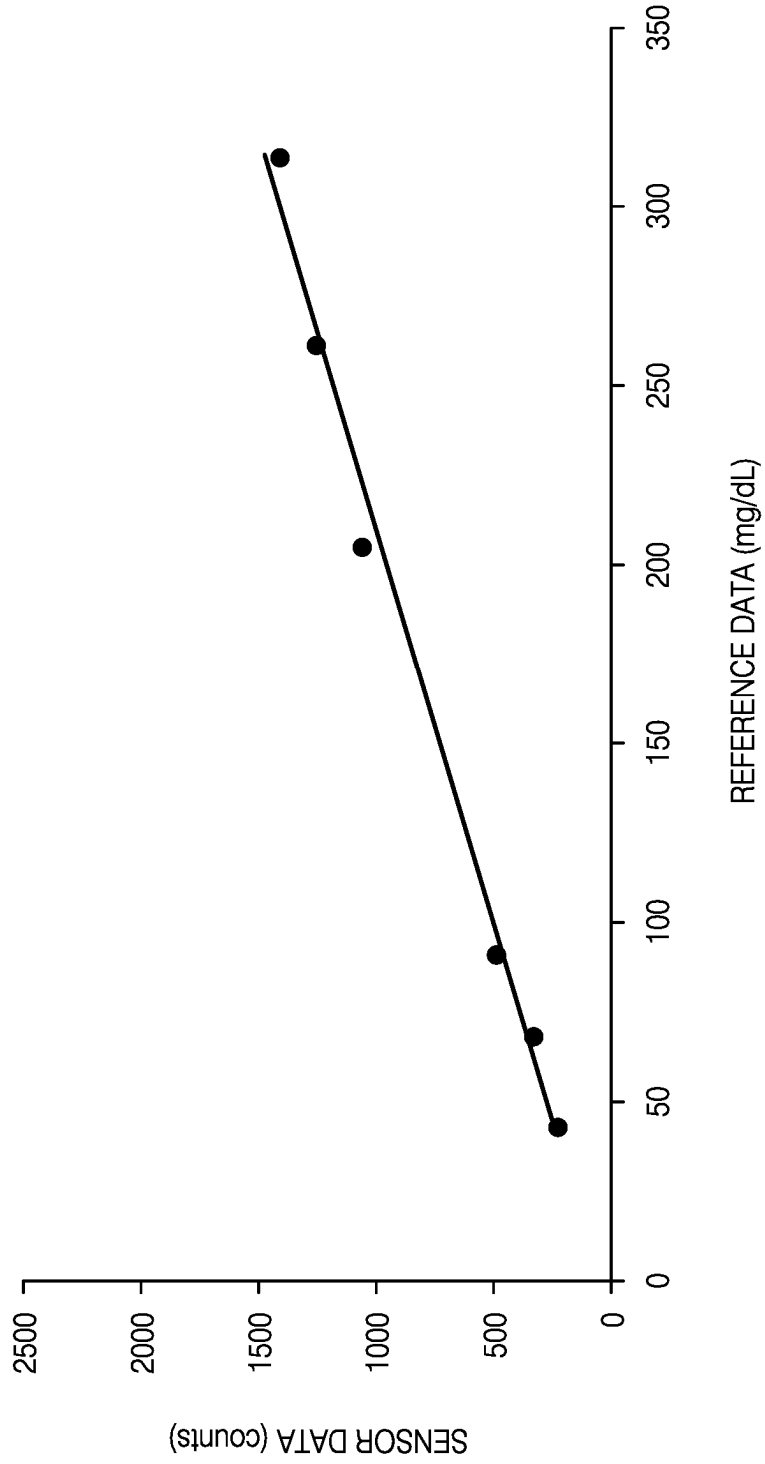

SENSOR DATA (counts)
2500
2000
1500
1000
500
0
0
50
100
150
200
250
300
350
REFERENCE DATA (mg/dL)

FIG. 26A

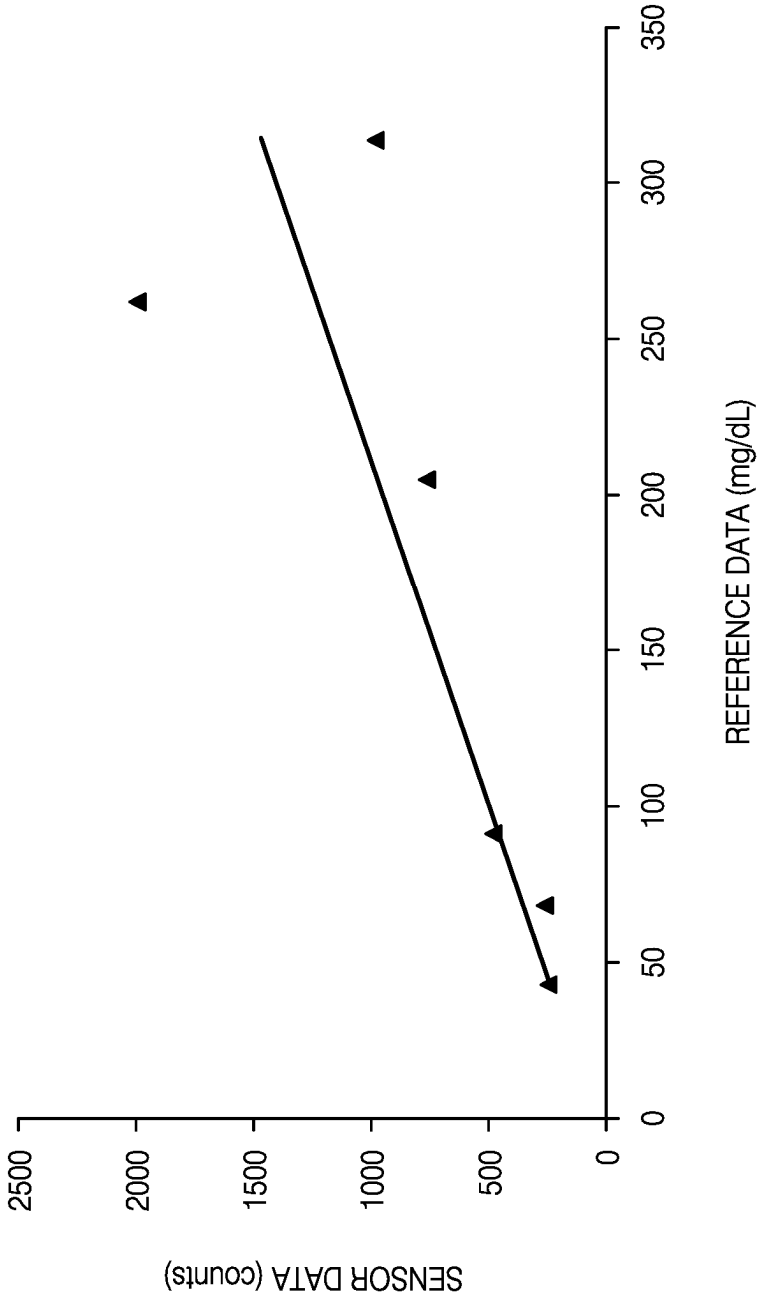
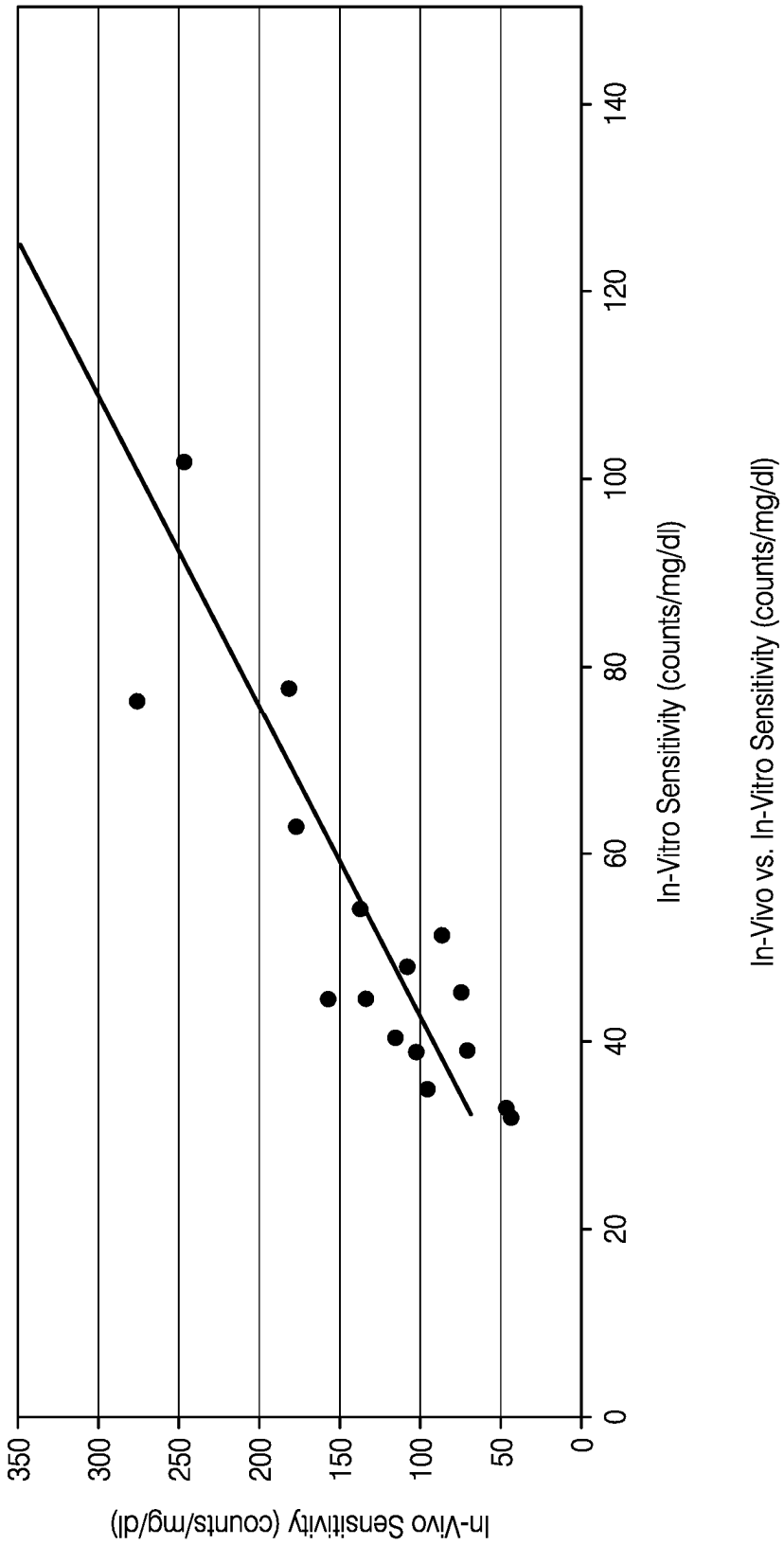

SENSOR DATA (counts)
2500
2000
1500
1000
500
0
0
50
100
150
200
250
300
350
REFERENCE DATA (mg/dL)

FIG. 26B



In-Vivo vs. In-Vitro Sensitivity (counts/mg/dl)

FIG. 27

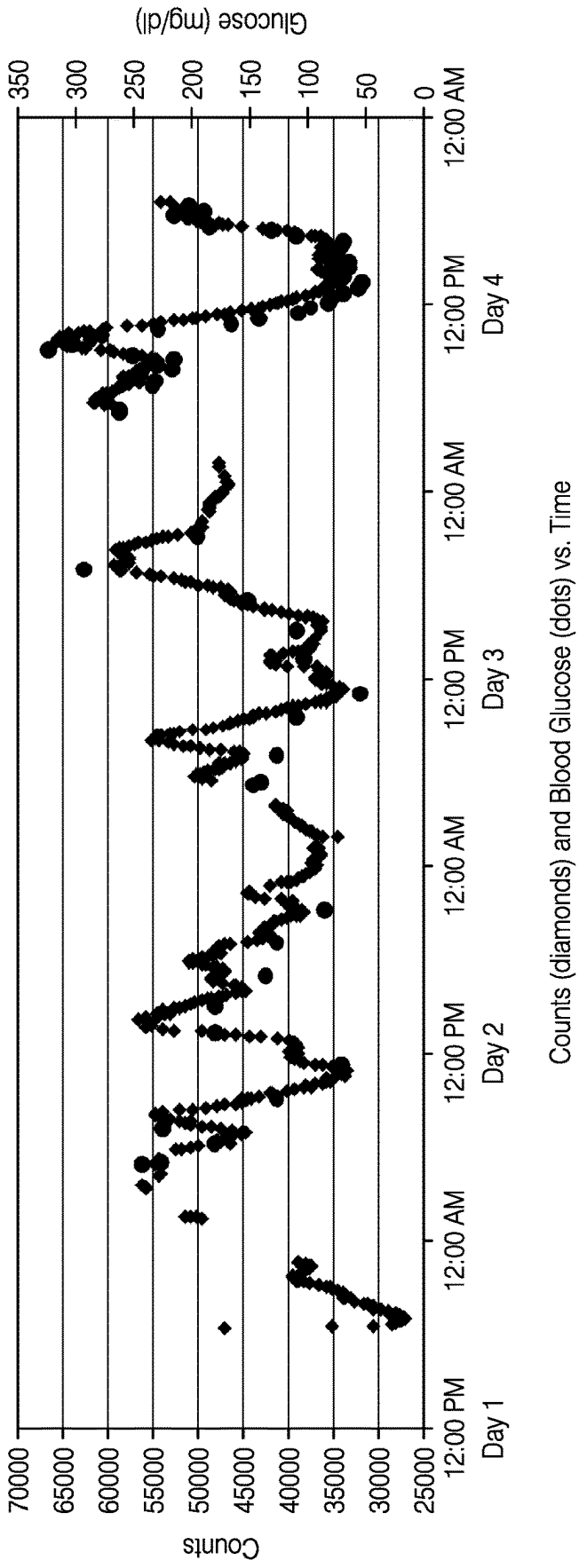
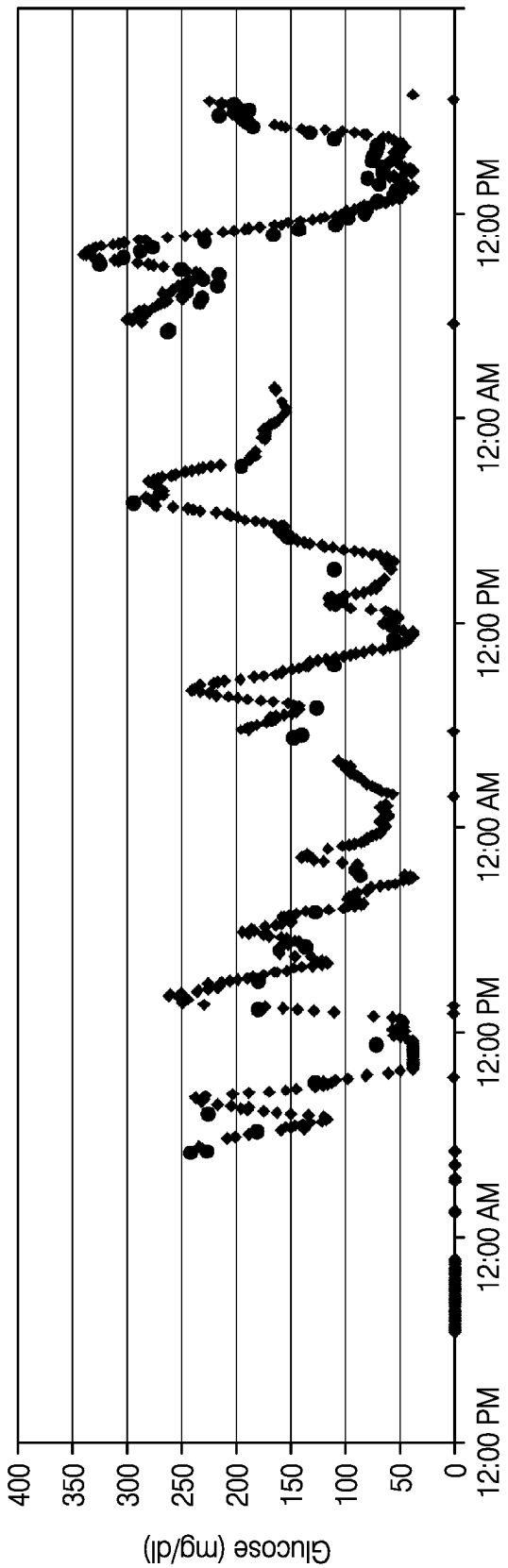

Glucose (mg/dl)
350
300
250
200
150
100
50
0
Counts
70000
65000
60000
55000
50000
45000
40000
35000
30000
25000
12:00 PM
Day 1
12:00 AM
12:00 PM
Day 2
12:00 AM
12:00 PM
Day 3
12:00 AM
12:00 PM
Day 4
12:00 AM
Counts (diamonds) and Blood Glucose (dots) vs. Time

FIG. 28



Sensor (diamonds) and Blood (dots) Glucose vs. Time. Calibration points shown by black dotx. Predicted in-vivo sensitivity used for calibration during day 1.

FIG. 29

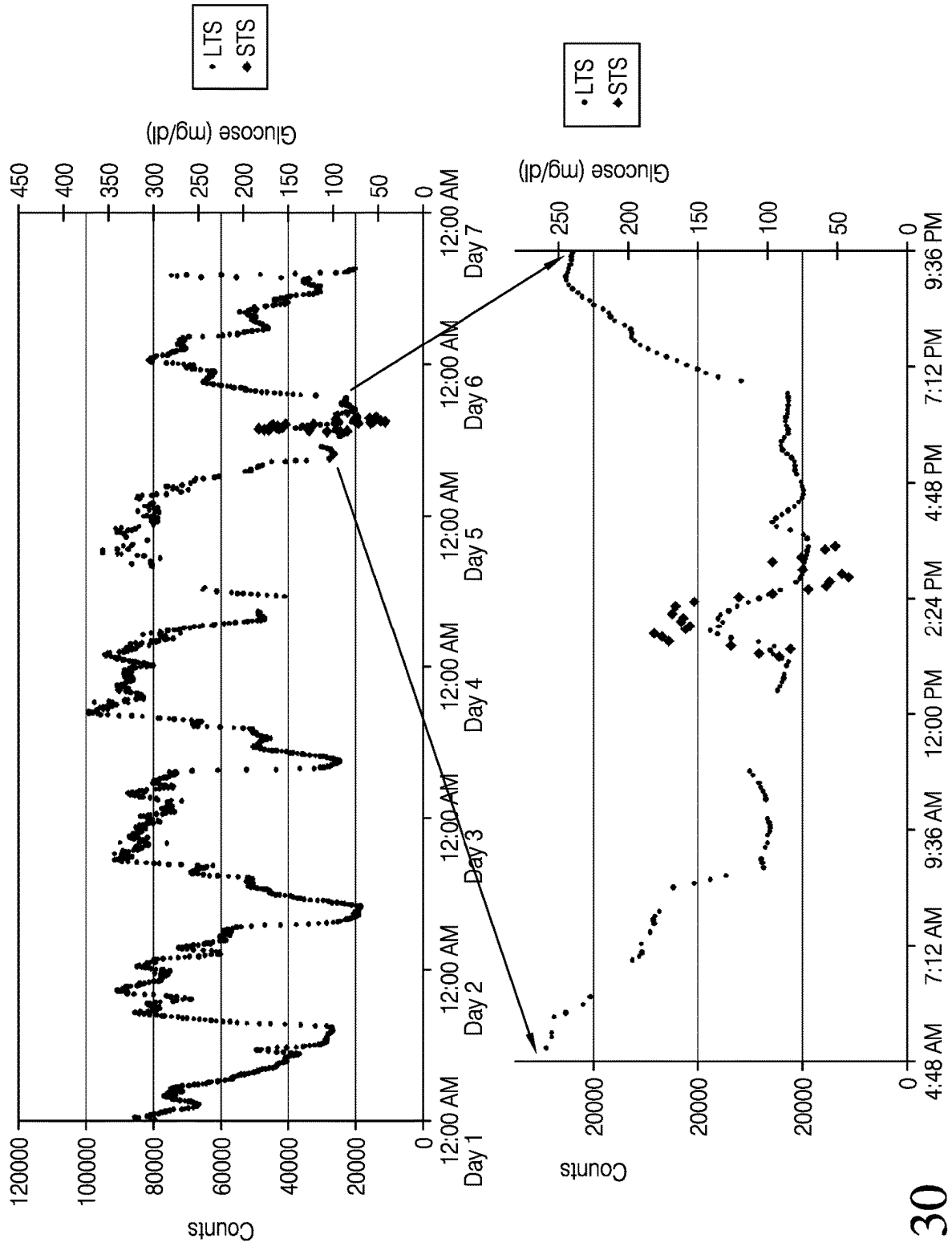

LTS
STS
Glucose (mg/dl)
Counts
12:00 AM Day 1
12:00 AM Day 2
12:00 AM Day 3
12:00 AM Day 4
12:00 AM Day 5
12:00 AM Day 6
12:00 AM Day 7
4:48 AM
7:12 AM
9:36 AM
12:00 PM
2:24 PM
4:48 PM
7:12 PM
9:36 PM

FIG. 30

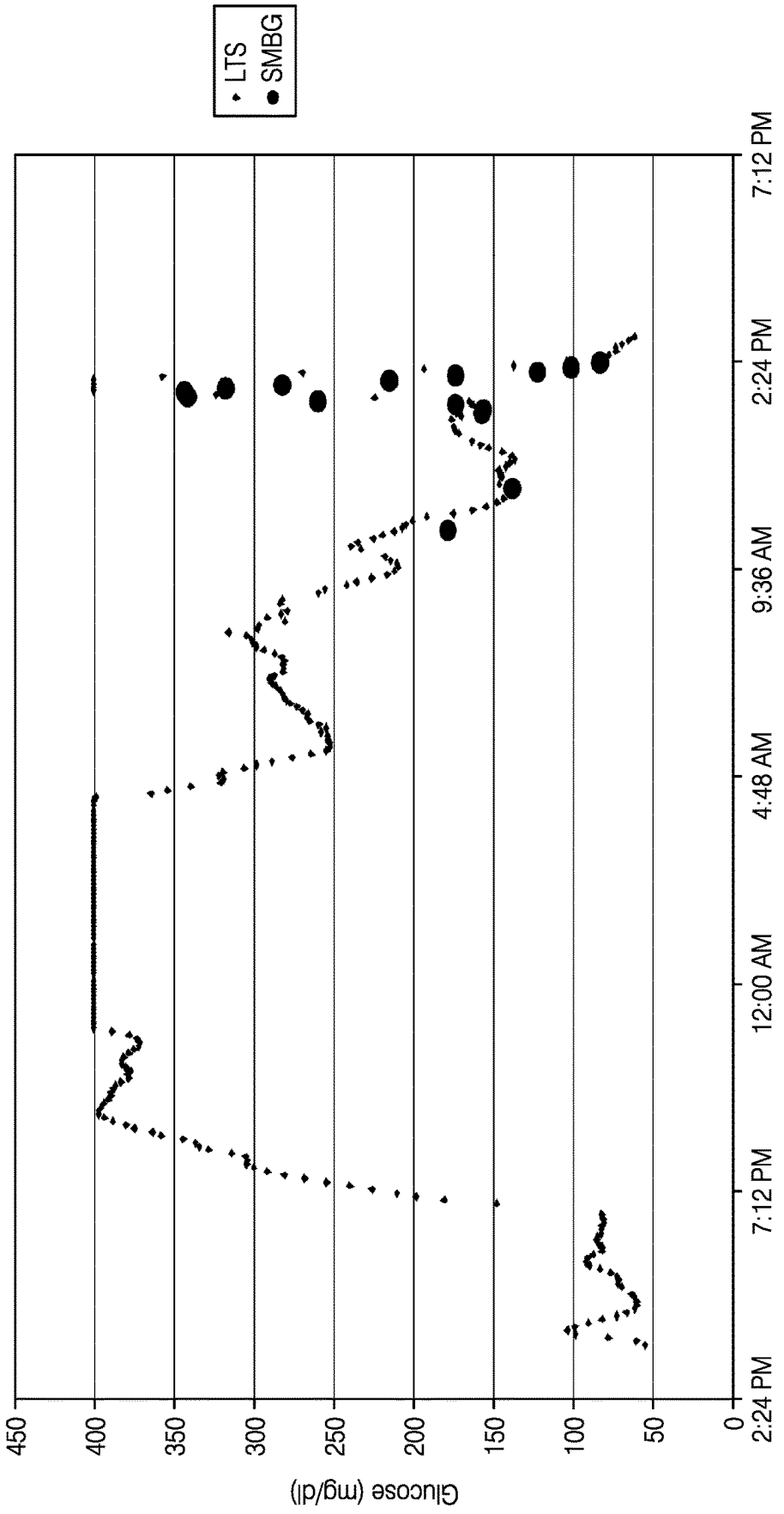

LTS
SMBG
Glucose (mg/dl)
450
400
350
300
250
200
150
100
50
0
2:24 PM
7:12 PM
12:00 AM
4:48 AM
9:36 AM
2:24 PM
7:12 PM

FIG. 31

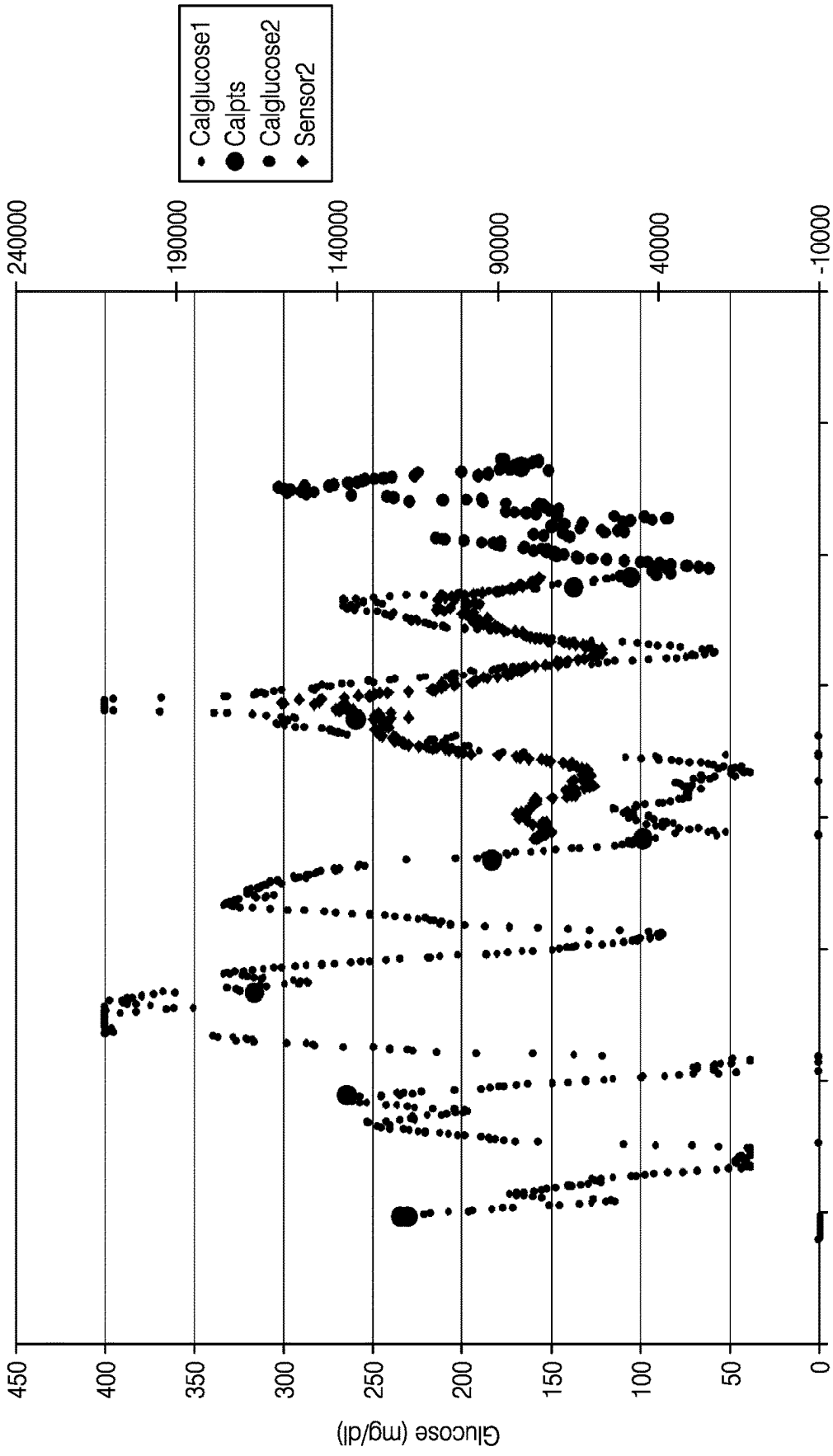

Calglucose1
Calpts
Calglucose2
Sensor2
240000
190000
140000
90000
40000
-10000
450
400
350
300
250
200
150
100
50
0
Glucose (mg/dl)
12:00 AM
12:00 PM
12:00 AM
12:00 PM
12:00 AM
12:00 PM
12:00 AM
12:00 PM
Day 1
Day 2
Day 3
Day 4

FIG. 32

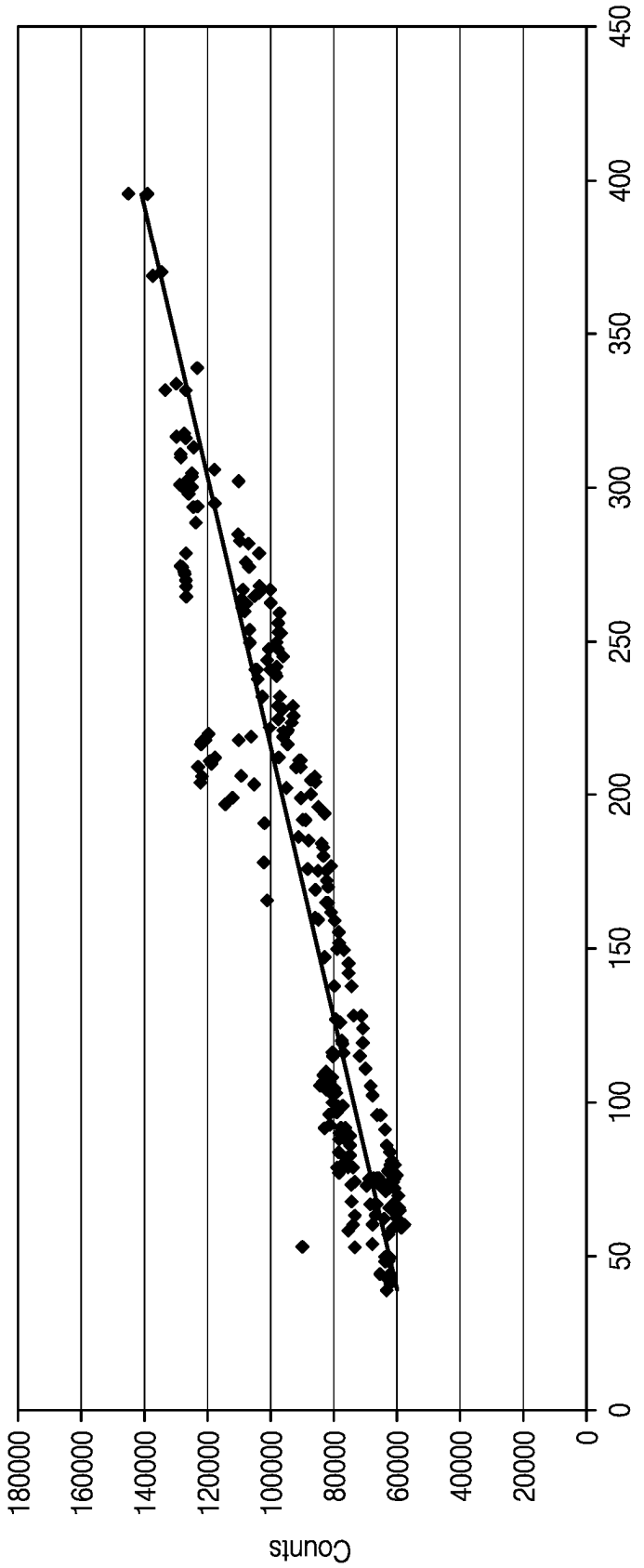

Counts
180000
160000
140000
120000
100000
80000
60000
40000
20000
0
mg/dl
0
50
100
150
200
250
300
350
400
450

FIG. 33

SYSTEM AND METHODS FOR PROCESSING ANALYTE SENSOR DATA FOR SENSOR CALIBRATION

INCORPORATION BY REFERENCE TO RELATED APPLICATIONS

Any and all priority claims identified in the Application Data Sheet, or any correction thereto, are hereby incorporated by reference under 37 CFR 1.57. This application is a continuation of U.S. application Ser. No. 16/457,628, filed Jun. 28, 2019, which is a continuation of U.S. application Ser. No. 15/787,595, filed Oct. 18, 2017, which is a continuation of U.S. application Ser. No. 15/065,623, filed Mar. 9, 2016, now U.S. Pat. No. 9,918,668, which is a continuation of U.S. application Ser. No. 13/607,162, filed Sep. 7, 2012, now U.S. Pat. No. 9,314,196, which is a continuation of U.S. application Ser. No. 12/683,755, filed on Jan. 7, 2010, now U.S. Pat. No. 8,611,978, which is a continuation of U.S. application Ser. No. 11/373,628, filed on Mar. 9, 2006, now U.S. Pat. No. 7,920,906, which claims the benefit of U.S. Provisional Application No. 60/660,743, filed on Mar. 10, 2005. Each of the aforementioned applications is incorporated by reference herein in its entirety, and each is hereby expressly made a part of this specification.

FIELD OF THE INVENTION

The present invention relates generally to systems and methods for analyte sensor data processing. Particularly, the present invention relates calibration of sensors.

BACKGROUND OF THE INVENTION

Diabetes mellitus is a disorder in which the pancreas cannot create sufficient insulin (Type I or insulin dependent) and/or in which insulin is not effective (Type 2 or non-insulin dependent). In the diabetic state, the victim suffers from high blood sugar, which can cause an array of physiological derangements (e.g., kidney failure, skin ulcers, or bleeding into the vitreous of the eye) associated with the deterioration of small blood vessels. A hypoglycemic reaction (low blood sugar) can be induced by an inadvertent overdose of insulin, or after a normal dose of insulin or glucose-lowering agent accompanied by extraordinary exercise or insufficient food intake.

Conventionally, a diabetic person carries a self-monitoring blood glucose (SMBG) monitor, which typically comprises uncomfortable finger pricking methods. Due to the lack of comfort and convenience, a diabetic will normally only measure his or her glucose level two to four times per day. Unfortunately, these time intervals are so far spread apart that the diabetic will likely find out too late, sometimes incurring dangerous side effects, of a hyper- or hypoglycemic condition. Alternatively, a long term sensor implanted in a diabetic person can provide substantially continuous blood glucose measurements to a receiver carried by the diabetic and obviate the finger pricking method. Due to its biological interface, after it is implanted a long term sensor typically requires a waiting period after which its sensor data must be calibrated using the finger prick method or the like.

SUMMARY OF THE INVENTION

Systems and methods for providing blood glucose measurements are needed that can shorten the calibration process of a long term sensor, avoid or reduce dependence on using the finger prick method during calibration, or overcome other problems known in the art. Systems and methods are disclosed that provide calibration of a long term sensor data using sensor data from another sensor. In one embodiment, the invention includes a method for calibrating an analyte sensor, the method including receiving sensor data from a first analyte sensor, receiving sensor data from a second analyte sensor, and calibrating the sensor data from the first analyte sensor using the sensor data from the second analyte sensor.

In one aspect of the first embodiment, the first analyte sensor is a long term substantially continuous analyte sensor and the second analyte sensor is a short term substantially continuous sensor. One embodiment of this aspect further includes receiving sensor data from a set of substantially continuous analyte sensors, the set of sensors comprising one or more short term sensors, and the set of sensors being employed in a host in series such that the overall time period during which sensor data from the set of sensors is received is greater than the useful lifespan of one of the short term sensors and calibrating the sensor data from the first analyte sensor using the sensor data from the second analyte sensor and the set of sensors. In another embodiment of this aspect calibrating the long term sensor data further includes using sensor data from the short term analyte sensor to update the calibration of the long term analyte sensor.

In a second aspect of the first embodiment, the first and second analyte sensors are glucose sensors.

In a third aspect of the first embodiment, the sensor data from the first analyte sensor includes at least one sensor data point, wherein the sensor data from the second analyte sensor includes at least one data point, and wherein calibrating the sensor data further comprises forming one or more matched data pairs by matching at least one sensor data point from the first analyte sensor to at least one sensor data point from the second analyte sensor and forming a calibration set comprising at least one matched data pair.

In a fourth aspect of the first embodiment, the sensor data from the first analyte sensor includes at least six sensor data points, wherein the sensor data from the second analyte sensor includes at least six sensor data points, and wherein calibrating the sensor data further comprises forming at least six matched data pairs by matching each sensor data point from the first analyte sensor to a corresponding sensor data point from the second analyte sensor and forming a calibration set comprising at least six matched data pairs.

In a fifth aspect of the first embodiment, the sensor data from the first analyte sensor includes at least twenty sensor data points, wherein the sensor data from the second analyte sensor includes at least twenty sensor data points, and wherein calibrating the sensor data further comprises forming at least twenty matched data pairs by matching each sensor data point from the first analyte sensor to a corresponding sensor data point from the second analyte sensor and forming a calibration set comprising at least twenty matched data pairs.

In a sixth aspect of the first embodiment, the method further includes constructing a first curve from the sensor data from the first analyte sensor and constructing a second curve from the sensor data from the second analyte sensor, wherein calibrating the sensor data from the first analyte sensor comprises matching the first curve with the second curve.

In a seventh aspect of the first embodiment, the first analyte sensor is a short term substantially continuous

analyte sensor and the second analyte sensor is a short term substantially continuous analyte sensor.

In a eighth aspect of the first embodiment, the first analyte sensor is a long term substantially continuous analyte sensor and the second analyte sensor is a long term substantially continuous analyte sensor.

In a ninth aspect of the first embodiment, the first analyte sensor is a short term substantially continuous analyte sensor and the second analyte sensor is a long term substantially continuous analyte sensor.

In a tenth aspect of the first embodiment, the method further includes receiving data from a non-continuous reference source, and wherein said calibrating further includes using data from the non-continuous reference source to calibrate the sensor data from the second analyte sensor.

In an eleventh aspect of the first embodiment, the non-continuous reference source is a blood glucose monitor.

In a twelfth aspect of the first embodiment, the non-continuous reference source is an in-vitro calibration.

In a thirteenth aspect of the first embodiment, the non-continuous reference source is an optical sensor.

In a fourteenth aspect of the first embodiment, calibrating the long term sensor data further includes using sensor data from the short term analyte sensor to update the calibration of the long term analyte sensor.

In a second embodiment, the invention includes a method of processing data from a substantially continuous analyte sensor, the method including testing an analyte sensor prior to insertion into a host to determine at least one sensor data characteristic, employing the analyte sensor in the host, receiving sensor data from the analyte sensor, and calibrating the sensor data using the sensor data characteristic.

In one aspect of the second embodiment, the substantially continuous analyte sensor is a short term sensor.

In a second aspect of the second embodiment, the substantially continuous analyte sensor is an implantable long term sensor.

In a third aspect of the second embodiment, the method further includes receiving reference data from a reference analyte monitor, the reference data comprising at least one reference data characteristic and calibrating the sensor data using the reference data characteristic.

In a fourth embodiment, the invention includes a system for calibrating a substantially continuous analyte sensor, the system including a first substantially continuous analyte sensor, a second substantially continuous analyte sensor, a first sensor data receiving module operably linked to said first sensor and configured to receive at least one sensor data point from said first sensor, a second sensor data receiving module operably linked to said second sensor and configured to receive at least one sensor data point from said second sensor, and a processor module in data communication with the first sensor data receiving module linked to said first sensor and further in data communication with the second sensor data receiving module linked to said second sensor, said processor module configured to calibrate the sensor data from the first sensor using sensor data from the second sensor.

In one aspect of the fourth embodiment, the processor module is further configured to match at least one time-matched data point from said first analyte sensor and said second analyte sensor to form at least one calibration set for calibrating the first analyte sensor including at least one matched data pair.

In a second aspect of the fourth embodiment, the first analyte sensor is a long term substantially continuous analyte sensor and the second analyte sensor is a short term substantially continuous analyte sensor.

In a third aspect of the fourth embodiment, the first analyte sensor is a short term substantially continuous analyte sensor and the second analyte sensor is a short term substantially continuous analyte sensor.

In a fourth aspect of the fourth embodiment, the first analyte sensor is a long term substantially continuous analyte sensor and the second analyte sensor is a long term substantially continuous analyte sensor.

In a fifth aspect of the fourth embodiment, the first analyte sensor is a short term substantially continuous analyte sensor and the second analyte sensor is a long term substantially continuous analyte sensor.

In a sixth aspect of the fourth embodiment, the first substantially continuous analyte sensor and said second substantially continuous analyte sensor are each glucose sensors.

In a fifth embodiment, the invention includes a method for simultaneously utilizing at least two analyte sensors, the method including receiving sensor data from a first analyte sensor, receiving sensor data from a second analyte sensor, and processing the sensor data from the first analyte sensor using the sensor data from the second analyte sensor.

In one aspect of the fifth embodiment, the sensor data from the second analyte sensor comprises time delay information and processing comprises modifying the sensor data from a first analyte sensor responsive to the time delay information.

In a second aspect of the fifth embodiment, processing comprises utilizing the sensor data from the second analyte sensor to assess performance of the first analyte sensor.

In a third aspect of the fifth embodiment, the first analyte sensor is a transcutaneous sensor.

In a fourth aspect of the fifth embodiment, the first analyte sensor is a wholly implantable sensor. In one embodiment of this aspect, the second analyte sensor is a transcutaneous sensor.

In a fifth aspect of the fifth embodiment, the second analyte sensor is a transcutaneous sensor.

In a sixth aspect of the fifth embodiment, the second analyte sensor is a wholly implantable sensor.

In a seventh aspect of the fifth embodiment, the processing comprises calibrating the first analyte sensor.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. **1**A is a schematic of a system with one receiver where one sensor is used to calibrate another sensor.

FIG. **1**B is a schematic of a system with more than one receiver where one sensor is used to calibrate another sensor.

FIG. **1**C is a schematic of a system that uses sensor data from a short term sensor to calibrate sensor data from a long term sensor.

FIG. **1**D is a schematic of a system that uses two or more short term sensors to calibrate short term sensors and/or long term sensors.

FIG. **1**E is a schematic of a system that uses sensor data from a long term sensor to calibrate sensor data from another long term sensor.

FIG. **1**F is a schematic of a system that uses sensor data from a short term sensor to calibrate sensor data from another short term sensor.

FIG. **2** is a perspective view of a transcutaneous analyte sensor system, including an applicator, a mounting unit, and an electronics unit.

FIG. **3** is a perspective view of a mounting unit, including the electronics unit in its functional position.

FIG. **4** is an exploded perspective view of a mounting unit, showing its individual components.

FIG. **5**A is an exploded perspective view of a contact subassembly, showing its individual components.

FIG. **5**B is a perspective view of an alternative contact configuration.

FIG. **5**C is a perspective view of another alternative contact configuration.

FIGS. **5**D to **5**H are schematic cross-sectional views of a portion of the contact subassembly; namely, a variety of embodiments illustrating alternative sealing member configurations.

FIG. **6**A is an expanded cutaway view of a proximal portion of a sensor.

FIG. **6**B is an expanded cutaway view of a distal portion of a sensor.

FIG. **6**C is a cross-sectional view through the sensor of FIG. **5**B on line C-C, showing an exposed electroactive surface of a working electrode surrounded by a membrane system.

FIG. **7** is an exploded side view of an applicator, showing the components that facilitate sensor insertion and subsequent needle retraction.

FIGS. **8**A to **8**D are schematic side cross-sectional views that illustrate applicator components and their cooperating relationships.

FIG. **9**A is a perspective view of an applicator and mounting unit in one embodiment including a safety latch mechanism.

FIG. **9**B is a side view of an applicator matingly engaged to a mounting unit in one embodiment, prior to sensor insertion.

FIG. **9**C is a side view of a mounting unit and applicator depicted in the embodiment of FIG. **9**B, after the plunger subassembly has been pushed, extending the needle and sensor from the mounting unit.

FIG. **9**D is a side view of a mounting unit and applicator depicted in the embodiment of FIG. **9**B, after the guide tube subassembly has been retracted, retracting the needle back into the applicator.

FIG. **9**E is a perspective view of an applicator, in an alternative embodiment, matingly engaged to the mounting unit after to sensor insertion.

FIG. **9**F is a perspective view of the mounting unit and applicator, as depicted in the alternative embodiment of FIG. **9**E, matingly engaged while the electronics unit is slidingly inserted into the mounting unit.

FIG. **9**G is a perspective view of the electronics unit, as depicted in the alternative embodiment of FIG. **9**E, matingly engaged to the mounting unit after the applicator has been released.

FIGS. **9**H and **9**I are comparative top views of the sensor system shown in the alternative embodiment illustrated in FIGS. **9**E to **9**G as compared to the embodiments illustrated in FIGS. **9**B to **9**D.

FIGS. **10**A to **10**C are side views of a sensor system adhered with an extensible adhesive pad in one embodiment. The figures illustrate the system prior to and during initial and continued release of the mounting unit from the host's skin.

FIGS. **11**A and **11**B are perspective and side cross-sectional views, respectively, of a sensor system showing the mounting unit immediately following sensor insertion and release of the applicator from the mounting unit.

FIGS. **12**A and **12**B are perspective and side cross-sectional views, respectively, of a sensor system showing the mounting unit after pivoting the contact subassembly to its functional position.

FIGS. **13**A to **13**C are perspective and side views, respectively, of the sensor system showing the sensor, mounting unit, and electronics unit in their functional positions.

FIG. **14** is a perspective view of a sensor system wirelessly communicating with a receiver.

FIGS. **15**A and **15**B are perspective views of a receiver in one preferred embodiment, wherein the receiver is provided with a docking station for receiving and holding the electronics unit (from the sensor assembly) when not in use.

FIG. **16** is an exploded perspective view of one exemplary embodiment of a continuous glucose sensor

FIG. **17** is a block diagram that illustrates electronics associated with a sensor system.

FIG. **18** is a graph that illustrates data smoothing of a raw data signal in one embodiment.

FIG. **19**A illustrates a first embodiment wherein the receiver shows a numeric representation of the estimated analyte value on its user interface, which is described in more detail elsewhere herein.

FIG. **19**B illustrates a second embodiment wherein the receiver shows an estimated glucose value and one hour of historical trend data on its user interface, which is described in more detail elsewhere herein.

FIG. **19**C illustrates a third embodiment wherein the receiver shows an estimated glucose value and three hours of historical trend data on its user interface, which is described in more detail elsewhere herein.

FIG. **19**D illustrates a fourth embodiment wherein the receiver shows an estimated glucose value and nine hours of historical trend data on its user interface, which is described in more detail elsewhere herein.

FIG. **20**A is a block diagram that illustrates a configuration of a medical device including a continuous analyte sensor, a receiver, and an external device.

FIGS. **20**B to **20**D are illustrations of receiver liquid crystal displays showing embodiments of screen displays.

FIG. **21** is a flow chart that illustrates the initial calibration and data output of the sensor data in one embodiment.

FIG. **22**A is a graph that illustrates a regression performed on a calibration set to obtain a conversion function in one exemplary embodiment.

FIG. **22**B is a graph that illustrates one example of using prior information for slope and baseline.

FIG. **22**C is a slope-baseline graph illustrating one example of using prior distribution information for determining a calibration slope and baseline.

FIG. **23** is a graph of two data pairs on a Clarke Error Grid to illustrate the evaluation of clinical acceptability in one exemplary embodiment.

FIG. **24** is a flow chart that illustrates the process of evaluation of calibration data for best calibration based on inclusion criteria of matched data pairs in one embodiment.

FIG. **25** is a flow chart that illustrates the process of evaluating the quality of the calibration in one embodiment.

FIG. **26**A and FIG. **26**B are graphs that illustrate an evaluation of the quality of calibration based on data association in one exemplary embodiment using a correlation coefficient.

FIG. **27** is a graph that illustrates an exemplary relationship between in-vitro and in-vivo sensitivity.

FIG. **28** is a graphical representation showing sensor data resulting from applying the in-vitro/in-vivo relationship of

FIG. **27** to a substantially continuous analyte sensor and showing blood glucose readings over a period of time, from an exemplary application.

FIG. **29** is a graphical representation of glucose data from an analyte sensor and blood glucose readings over a time period from an exemplary application.

FIG. **30** is a graphical representation of sensor data from an un-calibrated long term sensor and a calibrated short term sensor employed on the same host.

FIG. **31** is a graphical representation of sensor data from a long term sensor, that was calibrated using the short term sensor data shown in FIG. **30**, prospectively applied and compared to reference glucose measurements.

FIG. **32** is a graphical representation of sensor data from a short term sensor calibrated by sensor data from another short term sensor.

FIG. **33** is a graphical representation of a regression used to calibrate sensor data shown in FIG. **32**.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENT

The following description and examples illustrate a preferred embodiment of the present invention in detail. Those of skill in the art will recognize that there are numerous variations and modifications of this invention that are encompassed by its scope. Accordingly, the description of a preferred embodiment should not be deemed to limit the scope of the present invention.

Definitions

In order to facilitate an understanding of the disclosed invention, a number of terms are defined below.

The term "A/D Converter" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to hardware that converts analog signals into digital signals.

The term "analyte" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a substance or chemical constituent in a biological fluid (for example, blood, interstitial fluid, cerebral spinal fluid, lymph fluid or urine) that can be analyzed. Analytes can include naturally occurring substances, artificial substances, metabolites, and/or reaction products. In some embodiments, the analyte for measurement by the sensing regions, devices, and methods is glucose. However, other analytes are contemplated as well, including but not limited to acarboxyprothrombin; acylcarnitine; adenine phosphoribosyl transferase; adenosine deaminase; albumin; alpha-fetoprotein; amino acid profiles (arginine (Krebs cycle), histidine/urocanic acid, homocysteine, phenylalanine/tyrosine, tryptophan); andrenostenedione; antipyrine; arabinitol enantiomers; arginase; benzoylecgonine (cocaine); biotinidase; biopterin; c-reactive protein; carnitine; carnosinase; CD4; ceruloplasmin; chenodeoxycholic acid; chloroquine; cholesterol; cholinesterase; conjugated 1-ß hydroxy-cholic acid; cortisol; creatine kinase; creatine kinase MM isoenzyme; cyclosporin A; d-penicillamine; de-ethylchloroquine; dehydroepiandrosterone sulfate; DNA (acetylator polymorphism, alcohol dehydrogenase, alpha 1-antitrypsin, cystic fibrosis, Duchenne/Becker muscular dystrophy, glucose-6-phosphate dehydrogenase, hemoglobin A, hemoglobin S, hemoglobin C, hemoglobin D, hemoglobin E, hemoglobin F, D-Punjab, beta-thalassemia, hepatitis B virus, HCMV, HTLV-1, Leber hereditary optic neuropathy, MCAD, RNA, PKU, *Plasmodium vivax*, sexual differentiation, 21-deoxycorti sol); desbutylhalofantrine; dihydropteridine reductase; diptheria/tetanus antitoxin; erythrocyte arginase; erythrocyte protoporphyrin; esterase D; fatty acids/acylglycines; free ß-human chorionic gonadotropin; free erythrocyte porphyrin; free thyroxine (FT4); free tri-iodothyronine (FT3); fumarylacetoacetase; galactose/gal-1-phosphate; galactose-1-phosphate uridyltransferase; gentamicin; glucose-6-phosphate dehydrogenase; glutathione; glutathione perioxidase; glycocholic acid; glycosylated hemoglobin; halofantrine; hemoglobin variants; hexosaminidase A; human erythrocyte carbonic anhydrase I; 17-alpha-hydroxyprogesterone; hypoxanthine phosphoribosyl transferase; immunoreactive trypsin; lactate; lead; lipoproteins ((a), B/A-1, ß); lysozyme; mefloquine; netilmicin; phenobarbitone; phenytoin; phytanic/pristanic acid; progesterone; prolactin; prolidase; purine nucleoside phosphorylase; quinine; reverse tri-iodothyronine (rT3); selenium; serum pancreatic lipase; sissomicin; somatomedin C; specific antibodies (adenovirus, anti-nuclear antibody, anti-zeta antibody, arbovirus, Aujeszky's disease virus, dengue virus, *Dracunculus medinensis, Echinococcus granulosus, Entamoeba histolytica*, enterovirus, *Giardia duodenalisa, Helicobacter pylori*, hepatitis B virus, herpes virus, HIV-1, IgE (atopic disease), influenza virus, *Leishmania donovani*, leptospira, measles/mumps/rubella, *Mycobacterium leprae, Mycoplasma pneumoniae*, Myoglobin, *Onchocerca volvulus*, parainfluenza virus, *Plasmodium falciparum*, poliovirus, *Pseudomonas aeruginosa*, respiratory syncytial virus, rickettsia (scrub typhus), *Schistosoma mansoni, Toxoplasma gondii, Trepenoma pallidium, Trypanosoma cruzi/rangeli*, vesicular stomatis virus, *Wuchereria bancrofti*, yellow fever virus); specific antigens (hepatitis B virus, HIV-1); succinylacetone; sulfadoxine; theophylline; thyrotropin (TSH); thyroxine (T4); thyroxine-binding globulin; trace elements; transferrin; UDP-galactose-4-epimerase; urea; uroporphyrinogen I synthase; vitamin A; white blood cells; and zinc protoporphyrin. Salts, sugar, protein, fat, vitamins and hormones naturally occurring in blood or interstitial fluids can also constitute analytes in certain embodiments. The analyte can be naturally present in the biological fluid, for example, a metabolic product, a hormone, an antigen, an antibody, and the like. Alternatively, the analyte can be introduced into the body, for example, a contrast agent for imaging, a radioisotope, a chemical agent, a fluorocarbon-based synthetic blood, or a drug or pharmaceutical composition, including but not limited to insulin; ethanol; cannabis (marijuana, tetrahydrocannabinol, hashish); inhalants (nitrous oxide, amyl nitrite, butyl nitrite, chlorohydrocarbons, hydrocarbons); cocaine (crack cocaine); stimulants (amphetamines, methamphetamines, Ritalin, Cylert, Preludin, Didrex, PreState, Voranil, Sandrex, Plegine); depressants (barbituates, methaqualone, tranquilizers such as Valium, Librium, Miltown, Serax, Equanil, Tranxene); hallucinogens (phencyclidine, lysergic acid, mescaline, peyote, psilocybin); narcotics (heroin, codeine, morphine, opium, meperidine, Percocet, Percodan, Tussionex, Fentanyl, Darvon, Talwin, Lomotil); designer drugs (analogs of fentanyl, meperidine, amphetamines, methamphetamines, and phencyclidine, for example, Ecstasy); anabolic steroids; and nicotine. The metabolic products of drugs and pharmaceutical compositions are also contemplated analytes. Analytes such as neurochemicals and other chemicals generated within the body can also be analyzed, such as, for example, ascorbic acid, uric acid, dopamine, noradrena-

line, 3-methoxytyramine (3MT), 3,4-dihydroxyphenylacetic acid (DOPAC), homovanillic acid (HVA), 5-hydroxytryptamine (5HT), and 5-hydroxyindoleacetic acid (FHIAA).

The term "analyte sensor" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to any mechanism (e.g., enzymatic or non-enzymatic) by which analyte can be quantified. For example, some embodiments utilize a membrane that contains glucose oxidase that catalyzes the conversion of oxygen and glucose to hydrogen peroxide and gluconate:

$$\text{Glucose}+O_2 \rightarrow \text{Gluconate}+H_2O_2$$

Because for each glucose molecule metabolized, there is a proportional change in the co-reactant O_2 and the product H_2O_2, one can use an electrode to monitor the current change in either the co-reactant or the product to determine glucose concentration.

The term "biointerface membrane" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a permeable membrane that can be comprised of two or more domains and constructed of materials of a few microns thickness or more, which can be placed over the sensor body to keep host cells (e.g., macrophages) from gaining proximity to, and thereby damaging, the sensing membrane or forming a barrier cell layer and interfering with the transport of analyte across the tissue-device interface. The term "exit-site" as used herein is a broad term and is used in its ordinary sense, including, without limitation, the area where a medical device (for example, a sensor and/or needle) exits from the host's body.

The term "Clarke Error Grid" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to an error grid analysis, which evaluates the clinical significance of the difference between a reference glucose value and a sensor generated glucose value, taking into account (1) the value of the reference glucose measurement; (2) the value of the sensor glucose measurement; (3) the relative difference between the two values; and (4) the clinical significance of this difference. See Clarke et al., "Evaluating Clinical Accuracy of Systems for Self-Monitoring of Blood Glucose", Diabetes Care, Volume 10, Number 5, September-October 1987.

The term "clinical acceptability" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to determination of the risk of inaccuracies to a patient. Clinical acceptability considers a deviation between time corresponding glucose measurements (e.g., data from a glucose sensor and data from a reference glucose monitor) and the risk (e.g., to the decision making of a diabetic patient) associated with that deviation based on the glucose value indicated by the sensor and/or reference data. One example of clinical acceptability can be 85% of a given set of measured analyte values within the "A" and "B" region of a standard Clarke Error Grid when the sensor measurements are compared to a standard reference measurement.

The term "concordant" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to being in agreement or harmony, and/or free from discord.

The term "congruence" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the quality or state of agreeing, coinciding, or being concordant. In one example, congruence can be determined using rank correlation.

The term "Consensus Error Grid" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to an error grid analysis that assigns a specific level of clinical risk to any possible error between two time corresponding glucose measurements. The Consensus Error Grid is divided into zones signifying the degree of risk posed by the deviation. See Parkes et al., "A New Consensus Error Grid to Evaluate the Clinical Significance of Inaccuracies in the Measurement of Blood Glucose", Diabetes Care, Volume 23, Number 8, August 2000.

The phrase "continuous" as it relates to analyte sensing or an analyte sensor and as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to continuously, continually, and/or intermittently (regularly or irregularly) monitoring an analyte concentration, performed, for example, about every 1 second to 20 minutes.

The term "counts" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a unit of measurement of a digital signal. In one example, a raw data signal measured in counts is directly related to a voltage (converted by an A/D converter), which is directly related to current.

The terms "data association" and "data association function" as used herein are broad terms, and are to be given their ordinary and customary meaning to a person of ordinary skill in the art (and are not to be limited to a special or customized meaning), and refer without limitation to a statistical analysis of data and particularly its correlation to, or deviation from, from a particular curve. A data association function is used to show data association. For example, the data that forms that calibration set as described herein can be analyzed mathematically to determine its correlation to, or deviation from, a curve (e.g., line or set of lines) that defines the conversion function; this correlation or deviation is the data association. A data association function is used to determine data association. Examples of data association functions include, but are not limited to, linear regression, non-linear mapping/regression, rank (e.g., non-parametric) correlation, least mean square fit, mean absolute deviation (MAD), mean absolute relative difference. In one such example, the correlation coefficient of linear regression is indicative of the amount of data association of the calibration set that forms the conversion function, and thus the quality of the calibration.

The term "domain" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to regions of the biointerface membrane that can be

layers, uniform or non-uniform gradients (for example, anisotropic) or provided as portions of the membrane.

The term "EEPROM" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to electrically erasable programmable read-only memory, which is user-modifiable read-only memory (ROM) that can be erased and reprogrammed (e.g., written to) repeatedly through the application of higher than normal electrical voltage.

The term "electrochemically reactive surface" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the surface of an electrode where an electrochemical reaction takes place. In one example, a working electrode measures hydrogen peroxide produced by the enzyme catalyzed reaction of the analyte being detected reacts creating an electric current (for example, detection of glucose analyte utilizing glucose oxidase produces H_2O_2 as a by product, H_2O_2 reacts with the surface of the working electrode producing two protons ($2H^+$), two electrons ($2e^-$) and one molecule of oxygen (O_2) which produces the electronic current being detected). In the case of the counter electrode, a reducible species, for example, O_2 is reduced at the electrode surface in order to balance the current being generated by the working electrode.

The term "electronic connection" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to any electronic connection known to those in the art that can be utilized to interface the sensor head electrodes with the electronic circuitry of a device such as mechanical (e.g., pin and socket) or soldered.

The term "electronic connection" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to any electronic connection known to those in the art that can be utilized to interface the sensing region electrodes with the electronic circuitry of a device such as mechanical (for example, pin and socket) or soldered.

The term "ex vivo portion" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the portion of the device (for example, sensor) adapted to remain and/or exist outside the living body of a host.

The term "host" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to mammals, particularly humans.

The term "in vivo portion" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the portion of the device (for example, sensor) adapted for insertion and/or existence within the living body of a host.

The term "jitter" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to uncertainty or variability of waveform timing, which can be cause by ubiquitous noise caused by a circuit and/or environmental effects; jitter can be seen in amplitude, phase timing, or the width of the signal pulse.

The term "long term" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the lifespan of an analyte sensor. For example, the term "long term analyte sensor" is used herein relative to the term "short term analyte sensor" and designates an analyte sensor with a lifespan that is more than the lifespan of the short term analyte sensor. For example, a short term analyte sensor can have a lifespan of about less than about an hour to about three weeks, and a long term analyte sensor has a corresponding lifespan of longer than three weeks.

The term "matched data pairs" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to reference data (e.g., one or more reference analyte data points) matched with substantially time corresponding sensor data (e.g., one or more sensor data points).

The term "microprocessor" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a computer system or processor designed to perform arithmetic and logic operations using logic circuitry that responds to and processes the basic instructions that drive a computer.

The phrase "noncontinuous" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to single point analyte monitoring of an analyte concentration, for example, performed using blood glucose meters (e.g., using finger stick blood samples) and optical measuring techniques (e.g., near infrared spectroscopy, infrared spectroscopy, raman spectroscopy, photoacoustic spectroscopy, scatter and polarization changes), etc.

The terms "operable connection," "operably connected," and "operably linked" as used herein are broad terms, and are to be given their ordinary and customary meaning to a person of ordinary skill in the art (and are not to be limited to a special or customized meaning), and refer without limitation to one or more components being linked to another component(s) in a manner that allows transmission of signals between the components, e.g., wired or wirelessly. For example, one or more electrodes can be used to detect the amount of analyte in a sample and convert that information into a signal; the signal can then be transmitted to an electronic circuit means. In this case, the electrode is "operably linked" to the electronic circuitry.

The term "oxygen antenna domain" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a domain composed of a material that has higher oxygen solubility than aqueous media so that it concentrates oxygen from the biological fluid surrounding the biointerface membrane. The domain can then act as an oxygen reservoir during times of minimal oxygen need and has the capacity to provide on demand a higher oxygen gradient to facilitate oxygen transport across the membrane. This enhances function in the enzyme reaction domain and

at the counter electrode surface when glucose conversion to hydrogen peroxide in the enzyme domain consumes oxygen from the surrounding domains. Thus, this ability of the oxygen antenna domain to apply a higher flux of oxygen to critical domains when needed improves overall sensor function.

The term "quality of calibration" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the statistical association of matched data pairs in the calibration set used to create the conversion function. For example, an R-value can be calculated for a calibration set to determine its statistical data association, wherein an R-value greater than 0.79 determines a statistically acceptable calibration quality, while an R-value less than 0.79 determines statistically unacceptable calibration quality.

The term "raw data signal" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to an analog or digital signal directly related to the measured analyte from the analyte sensor. In one example, the raw data signal is digital data in "counts" converted by an A/D converter from an analog signal (e.g., voltage or amps) representative of an analyte concentration.

The term "RF transceiver" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a radio frequency transmitter and/or receiver for transmitting and/or receiving signals.

The term "R-value" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to one conventional way of summarizing the correlation of data; that is, a statement of what residuals (e.g., root mean square deviations) are to be expected if the data are fitted to a straight line by the a regression.

The term "sensing membrane" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to a permeable or semi-permeable membrane that can be comprised of two or more domains and constructed of materials of a few microns thickness or more, which are permeable to oxygen and can or can not be permeable to an analyte of interest. In one example, the sensing membrane comprises an immobilized glucose oxidase enzyme, which enables an electrochemical reaction to occur to measure a concentration of glucose.

The term "sensing region" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the region of a monitoring device responsible for the detection of a particular analyte. The sensing region generally comprises a non-conductive body, a working electrode (anode), a reference electrode (optional), and/or a counter electrode (cathode) passing through and secured within the body forming electrochemically reactive surfaces on the body and an electronic connective means at another location on the body, and a multi-domain membrane affixed to the body and covering the electrochemically reactive surface.

The term "sensor head" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the region of a monitoring device responsible for the detection of a particular analyte. In one example, a sensor head comprises a non-conductive body, a working electrode (anode), a reference electrode and a counter electrode (cathode) passing through and secured within the body forming an electrochemically reactive surface at one location on the body and an electronic connective means at another location on the body, and a sensing membrane affixed to the body and covering the electrochemically reactive surface. The counter electrode has a greater electrochemically reactive surface area than the working electrode. During general operation of the sensor a biological sample (e.g., blood or interstitial fluid) or a portion thereof contacts (directly or after passage through one or more membranes or domains) an enzyme (e.g., glucose oxidase); the reaction of the biological sample (or portion thereof) results in the formation of reaction products that allow a determination of the analyte (e.g., glucose) level in the biological sample. In some embodiments, the sensing membrane further comprises an enzyme domain (e.g., and enzyme layer), and an electrolyte phase (e.g., a free-flowing liquid phase comprising an electrolyte-containing fluid described further below).

The term "short term" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to the lifespan of an analyte sensor. For example, the term "short term analyte sensor" is used herein relative to the term "long term analyte sensor" and designates an analyte sensor with a lifespan less than the lifespan of a long term analyte sensor. For example, a short term analyte sensor can have a lifespan of less than an hour to about three weeks, and a long term analyte sensor has a corresponding lifespan of longer than three weeks.

The term "SRAM" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to static random access memory (RAM) that retains data bits in its memory as long as power is being supplied.

The term "substantially" as used herein is a broad term, and is to be given its ordinary and customary meaning to a person of ordinary skill in the art (and is not to be limited to a special or customized meaning), and refers without limitation to being largely but not necessarily wholly that which is specified.

In the disclosure which follows, the following abbreviations apply: Eq and Eqs (equivalents); mEq (milliequivalents); M (molar); mM (millimolar) μM (micromolar); N (Normal); mol (moles); mmol (millimoles); μmol (micromoles); nmol (nanomoles); g (grams); mg (milligrams); μg (micrograms); Kg (kilograms); L (liters); mL (milliliters); dL (deciliters); μL (microliters); cm (centimeters); mm (millimeters); μm (micrometers); nm (nanometers); h and hr (hours); min. (minutes); s and sec. (seconds); ° C. (degrees Centigrade).

Overview

Some of the aspects of the invention relate to calibrating a substantially continuous analyte sensor (e.g., a subcutaneous, transdermal (e.g., transcutaneous), or intravascular

device) using sensor data from another substantially continuous analyte sensor. The analyte sensor can be any type of sensor that measures a concentration of an analyte of interest or a substance indicative of the concentration or presence of the analyte. The analyte sensors can be short or long term sensors. For example, in some embodiments, the sensor data from a short term substantially continuous analyte sensor is used to calibrate the sensor data from another short term substantially continuous analyte sensor. In another embodiment, the sensor data from a short term substantially continuous analyte sensor is used to calibrate the sensor data from a long term substantially continuous analyte sensor. In one embodiment, the sensor data from a long term substantially continuous analyte sensor is used to calibrate the sensor data from another long term substantially continuous analyte sensor. In other embodiments, the sensor data from a long term substantially continuous analyte sensor is used to calibrate the sensor data from a short term substantially continuous analyte sensor. The analyte sensors can use any method of analyte-sensing, including enzymatic, chemical, physical, electrochemical, spectrophotometric, polarimetric, calorimetric, radiometric, or the like.

FIG. **1**A is a schematic illustrating one embodiment of a system where data from one is used to calibrate another sensor. The system includes two substantially continuous analyte sensors, $sensor_A$ and $sensor_B$, configured to provide sensor data to a receiver. $Sensor_A$ and $sensor_B$ can be any substantially continuous analyte sensors capable of determining the level of an analyte in the body, for example glucose, oxygen, lactase, insulin, hormones, cholesterol, medicaments, viruses, or the like. $Sensor_A$ and $sensor_B$ can be short term or long term sensors. $Sensor_A$ and $sensor_B$ detect information relating to an analyte in the host **6** and transmit the information (e.g., sensor data) to the receiver **8**. The transmission of the sensor data can be through a wire or wireless communication channel. In this embodiment, $sensor_A$ and $sensor_B$ measure the same analyte (e.g., glucose). If there is correlation and/or a predictive relationship between the sensor data from $sensor_A$ and $sensor_B$, the sensor data from one analyte sensor can be used to calibrate the sensor data from the other analyte sensor. If $sensor_A$ and $sensor_B$ measure two different analytes but there is correlation and/or predictive behavior between the sensor data for the two different analytes, the sensor data from one analyte sensor can be used to calibrate the sensor data from the other analyte sensor. In such an embodiment, the receiver is configured to receive the sensor data from both sensors, process the data, and calibrate one of the sensors, using e.g., data processing and calibration techniques as described herein and other suitable sensor calibration techniques.

FIG. **1**B is a schematic illustrating another embodiment of a system where data from one sensor is used to calibrate another sensor. This embodiment also includes two substantially continuous analyte sensors, $sensor_C$ and $sensor_D$, that detect information relating to an analyte in the host **6** and transmit the information to a receiver. Sensory and $sensor_B$ can be short term or long term sensors. In this embodiment, $sensor_C$ transmits sensor data to $receiver_A$ and $sensor_D$ transmits sensor data to $receiver_B$. $Receiver_A$ and $receiver_B$ are configured to communicate information with each other (e.g., analyte measurement information, corresponding timestamp of analyte measurement, etc.) so that the sensor data from one sensor can be used to calibrate the sensor data from the other. Here, for example, so long as the receivers can communicate, the communication protocol, content, and data format for one sensor and its receiver can be different than for the other sensor and its receiver. Some of the embodiments described and illustrated herein, e.g., FIGS. **1**C and **1**E, include one receiver that can receive data from one or more sensors. However, in such embodiments, additional receivers can be employed (e.g., one receiver for each sensor).

FIG. **1**C is a schematic illustrating an embodiment of a system that calibrates a long term sensor using sensor data from a short term sensor. A long term analyte sensor can have an initial instability time period during which it is unstable for environmental, physiological, or other reasons. For example, for a long term analyte sensor implanted subcutaneously, its stabilization can be dependent upon the maturity of the tissue ingrowth around and within the sensor (see, e.g., U.S. Publication No. US-2005-0112169-A1). Accordingly, determination of sensor stability can include waiting a time period (e.g., an implantable sensor is known to require a time period for tissue, and a transdermal (e.g., transcutaneous) sensor is known to require time to equilibrate the sensor with the user's tissue); in some embodiments, this waiting period is from about one minute to about three weeks. This waiting period can be predetermined by prior testing the sensor under similar conditions, and/or by analysis of the data from the sensor to determine that the sensor is stable. In this embodiment, the short term sensor measures the analyte and provides data to a receiver during the waiting period. Once the long term sensor is deemed to be stable, the data from the long term sensor requires calibration to provide an accurate value. Where the sensor data from the short term sensor has a correlative or predictive relationship with the sensor data from the long term sensor, the data from the short term sensor can be used to calibrate the long term sensor. Calibration of the sensor data from a long term sensor using sensor data provided by the short term sensor facilitates the use of the long term sensor data sooner and/or reduces or obviates the need to calibrate the long term sensor with single point calibration techniques, e.g., using finger stick blood samples, optical measuring techniques etc.

The sensor data from a long term sensor can change over time, due to, for example, a decrease in its sensitivity. In one embodiment, a long term sensor can be recalibrated by employing a short term sensor on the same host as the long term sensor and then using the sensor data from the short term sensor to calibrate the sensor data from the long term sensor. For example, on a host having a long term (e.g., wholly implantable) sensor, a substantially continuous short term (e.g., transcutaneous) analyte sensor can also be employed. A short term sensor taking a measurement every five minutes provides sensor data for 288 measurements in a day. The receiver translates the short sensor data to estimate analyte values, which are used to re-calibrate the long term sensor using e.g., data processing techniques described herein, including data processing techniques described in reference to FIGS. **16** and **17**. As another example, a short term sensor can be used to trouble shoot a long term sensor, for example, when the long term sensor is wholly implanted in the host and experiences malfunctioning. In such a situation, a short term sensor can be inserted into the host (e.g., a transcutaneous sensor) and the data compared to that of the long term sensor (e.g., wholly implantable sensor) to diagnose potential problems with the long term sensor, for example, signal noise due to oxygen deficiencies or enzyme deficiency. In alternative embodiments, short and/or long term sensors can be used to diagnose problems in other short and/or long term sensors.

FIG. **1**D is a schematic illustrating an embodiment of a system that calibrates a long term sensor using sensor data

from one or more short term sensors. In certain circumstances a single short term sensor is not able to provide sensor data for the entire waiting period of the long term sensor. For example, the lifespan of the short term sensor may be less than the requisite waiting period. Accordingly, the use of two or more short term sensors can be used to provide sensor data for the entire waiting period for a long term sensor if the waiting period is greater than three days.

In FIG. **1**D, two or more short term sensors are used to provide analyte measurements during the waiting period of a long term sensor. For example, a short term sensor A is used to provide analyte measurements to a receiver for its usable lifespan. Prior to the end of the lifespan of sensor A, short term sensor B is employed to also provide analyte measurements to a receiver for the host **6**, so that sensor A and sensor B are both providing sensor data during an overlapping time period. If the waiting period of the long term sensor is longer than the period that can be covered by sensor A and B, another short-term sensor can be employed prior to the end of the lifespan of sensor B. Substitution of short term sensors in an overlapping manner can be repeated until the expiration of the waiting period. Once the long term sensor is stable, it can be calibrated using sensor data from one or more of the short term sensors. Alternatively, one or more short-term sensor may be implanted near the time when the long term sensor is expected to start working rather than having the short term sensors implanted during the entire waiting period of the long term sensor.

In one embodiment, where two or more short term sensors are used on the same host, the sensor data from one short term sensor can be used to calibrate the other short term sensor. For example, in the above-described embodiment, the receiver can use the sensor data from sensor A to calibrate the sensor data from B, which can be used to calibrate each successive short term sensor employed on the host **6**.

In some embodiments, one or more of the short term sensors are calibrated in-vitro, e.g., by a benchtop calibration process that predetermines the information particular to the sensor. In such embodiments, the series of short term sensors need not overlap each other for calibration purposes. In one example, a benchtop calibrated short term sensor can be used to calibrate a long term implantable glucose sensor so that manual glucose measurements (SMBG) are not needed. According to one embodiment of a sensor, the sensing mechanism of an enzyme-electrode based glucose sensor depends on two phenomena that are linear with glucose concentration: (1) diffusion of glucose through a membrane system situated between interstitial fluid and the electrode surface, and (2) an enzymatic reaction within the membrane system. Because of this linearity, calibration/recalibration of the sensor consists of solving the line

$$y=mx+b$$

where y denotes the sensor signal (in units of A/D counts), x the estimated glucose concentration (mg/dl), m the sensor sensitivity to glucose (counts/mg/dl), and b the baseline signal (counts).

Typically, calibration requires at least two independent, paired measurements (x_1, y_1; x_2, y_2) to solve for m and b and thus allow glucose estimation when only the sensor signal, y is available. However, if prior information is available for m and/or b, then calibration can occur with fewer paired measurements. In one embodiment of a semi-benchtop calibration technique, prior information (e.g., benchtop tests) determine the sensitivity of the sensor and/or the baseline signal of the sensor by analyzing sensor data from measurements taken by the sensor of a controlled solution (e.g., prior to inserting the sensor). If there exists a predictive relationship between benchtop sensor parameters and in-vivo parameters, then this information can be used by the calibration procedure. For example, if a predictive relationship exists between benchtop sensitivity and in-vivo sensitivity, $m \approx f(m_{benchtop})$, then the predicted m can be used, along with a single matched pair, to solve for b ($b=y-mx$, semi-benchtop calibration). If in addition, b can be assumed to be 0, for example with a biointerface electrode configuration, then both m and b are known a priori, matched pairs are not needed for calibration, and the sensor can be completely benchtop calibrated.

Other methods for using information obtained prior to sensor insertion in sensor calibration are discussed in more detail elsewhere herein.

In some embodiments, the long term sensor can be benchtop calibrated and then once implanted, the calibration of the data from the long term sensor can be adjusted using additional reference data, for example, reference data from a short term sensor. Typically for a long term sensor, as its biointerface matures, its m and b change. This change (a 10% change in signal amplitude, for example) over a period of time (two weeks) can be solved for with periodic use of a short term sensor as follows: a user implanted with a long term sensor inserts a short term sensor, the short term sensor glucose values (if collected at 5 min intervals, there are 288 per day) are used as the reference values, x, to which the long term sensor values, y, can be calibrated. This procedure would allow for accurate estimates of m, b, and time-lag (assuming the time-lag of the short term sensor signal relative to blood glucose). Accordingly, reference data from a short term glucose sensor can be used in the initial calibration process described herein and shown in FIG. **21**, and the update calibration process described herein and shown in FIG. **24**.

In the embodiment illustrated in FIG. **1**E, the sensor data for one long term sensor is used to calibrate the sensor data from another long term sensor. Long term sensor A can be employed on the host **6** and calibrated using, e.g., data from a short term sensor, a long term sensor, or another calibration technique. Prior to the end of the lifespan of long term sensor A, long term sensor B can be implanted into the host **6**. Once long term sensor B is stable, sensor data from long term sensor A can be used to calibrate the sensor data from long term sensor B. This calibration process can be repeated for additional long term sensors employed on host **6**, so that a series of long term sensors can be employed, with the sensor data from each newly employed sensor being calibrated using sensor data from the previously employed long term sensor. By this means, a patient can have continuity of data while only undergoing one procedure for each long term implant, for example, once per year. In one example, Sensor A is implanted at time 0. After 11 months, Sensor B is implanted, and Sensor A is left in place. During the next month, Sensor B starts up and is calibrated using data from sensor A. At time 23 months, Sensor C is implanted and Sensor A is removed. Sensor B is used to calibrate Sensor C, and so forth for as long as the patient wishes to continue placing new sensors. Other time frames of sensor implantation and overlap are also possible.

FIG. **1**F is a schematic illustrating an embodiment of a system that calibrates a short term sensor using sensor data from a short term sensor. It is known that some transdermal (e.g., transcutaneous) sensors require time to equilibrate the sensor with the user's tissue (e.g., due to electrochemical break-in). For example, when a first short term sensor's

lifespan is nearing its end, a second short term sensor can be implanted into the user's tissue (e.g., at another location) to allow that sensor to equilibrate while the first short term sensor is still providing data. In addition to providing data during the equilibration time of the second short term sensor (thereby allowing data continuity), the first short term sensor can provide additional data, including calibration information, time lag information, drift information, and the like, to the second short term sensor to increase its intelligence and enhance its performance.

The sensors of the preferred embodiments can be employed sequentially and/or simultaneously (e.g., redundant sensors). For example, one or more short or long term sensors can be used simultaneously in order to compare, trouble-shoot, and/or provide increased data to the host. In one situation, by providing at least two redundant sensors, transient problems experienced by one sensor can be negated by the use of the other sensor.

An analyte sensor uses any known method, including invasive, minimally invasive, and non-invasive sensing techniques, to provide an output signal indicative of the concentration of the analyte of interest. The output signal is typically a raw signal that is used to provide a value that can be used for calibrating another analyte sensor or an analyte sensor used in another application. Appropriate smoothing, calibration, and evaluation methods can be applied to the raw signal and/or system as a whole to provide relevant and acceptable analyte data for calibrating another analyte sensor or the same analyte sensor used in another application.

The analyte sensor(s) useful with the preferred embodiments can be any device capable of measuring the concentration of an analyte of interest. One exemplary embodiment is described below, which calibrates a long term implantable glucose sensor based on data from a short term implantable glucose sensor. However, it can be understood that the devices and methods described herein can be applied to any device capable of detecting a concentration of analyte of and providing an output signal that represents the concentration of the analyte for use in calibrating another sensor.

Short Term Sensor

In one embodiment, the short term sensors for use as described herein include a transcutaneous analyte sensor system that includes an applicator for inserting the transcutaneous (transdermal) analyte sensor under a host's skin. The sensor system includes a sensor for sensing the analyte, wherein the sensor is associated with a mounting unit adapted for mounting on the skin of the host. The mounting unit houses the electronics unit associated with the sensor and is adapted for fastening to the host's skin. In certain embodiments, the system further includes a receiver for receiving and/or processing sensor data.

FIG. **2** is a perspective view of a transcutaneous analyte sensor system **10**. In the preferred embodiment of a system as depicted in FIG. **2**, the sensor includes an applicator **12**, a mounting unit **14**, and an electronics unit **16**. The system can further include a receiver **158**, such as is described in more detail with reference to FIG. **14**.

The mounting unit (housing) **14** includes a base **24** adapted for mounting on the skin of a host, a sensor adapted for transdermal (e.g., transcutaneous) insertion through the skin of a host (see FIG. **5**A), and one or more contacts **28** configured to provide secure electrical contact between the sensor and the electronics unit **16**. The mounting unit **14** is designed to maintain the integrity of the sensor in the host so as to reduce or eliminate translation of motion between the mounting unit, the host, and/or the sensor.

In one embodiment, an applicator **12** is provided for inserting the sensor **32** through the host's skin at the appropriate insertion angle with the aid of a needle (see FIGS. **7** through **9**), and for subsequent removal of the needle using a continuous push-pull action. Preferably, the applicator comprises an applicator body **18** that guides the applicator components (see FIGS. **7** through **9**) and includes an applicator body base **60** configured to mate with the mounting unit **14** during insertion of the sensor into the host. The mate between the applicator body base **60** and the mounting unit **14** can use any known mating configuration, for example, a snap-fit, a press-fit, an interference-fit, or the like, to discourage separation during use. One or more release latches **30** enable release of the applicator body base **60**, for example, when the applicator body base **60** is snap fit into the mounting unit **14**.

The electronics unit **16** includes hardware, firmware, and/or software that enable measurement of levels of the analyte via the sensor. For example, the electronics unit **16** can comprise a potentiostat, a power source for providing power to the sensor, other components useful for signal processing, and preferably an RF module for transmitting data from the electronics unit **16** to a receiver. Electronics can be affixed to a printed circuit board (PCB), or the like, and can take a variety of forms. For example, the electronics can take the form of an integrated circuit (IC), such as an Application-Specific Integrated Circuit (ASIC), a microcontroller, or a processor. Preferably, electronics unit **16** houses the sensor electronics, which comprise systems and methods for processing sensor analyte data. Examples of systems and methods for processing sensor analyte data are described in more detail in U.S. Publication No. US-2005-0027463-A1.

After insertion of the sensor using the applicator **12**, and subsequent release of the applicator **12** from the mounting unit **14** (see FIGS. **9**B to **9**D), the electronics unit **16** is configured to releasably mate with the mounting unit **14** in a manner similar to that described above with reference to the applicator body base **60**. The electronics unit **16** includes contacts on its backside (not shown) configured to electrically connect with the contacts **28**, such as are described in more detail with reference to FIGS. **3** through **5**. In one embodiment, the electronics unit **16** is configured with programming, for example initialization, calibration reset, failure testing, or the like, each time it is initially inserted into the mounting unit **14** and/or each time it initially communicates with the sensor **32**.

Mounting Unit

FIG. **3** is a perspective view of a sensor system of a preferred embodiment, shown in its functional position, including a mounting unit and an electronics unit matingly engaged therein. FIGS. **13**A to **13**C illustrate the sensor is its functional position for measurement of an analyte concentration in a host.

In preferred embodiments, the mounting unit **14**, also referred to as a housing, comprises a base **24** adapted for fastening to a host's skin. The base can be formed from a variety of hard or soft materials, and preferably comprises a low profile for minimizing protrusion of the device from the host during use. In some embodiments, the base **24** is formed at least partially from a flexible material, which is believed to provide numerous advantages over conventional transcutaneous sensors, which, unfortunately, can suffer from motion-related artifacts associated with the host's movement when the host is using the device. For example, when a transcutaneous analyte sensor is inserted into the host, various movements of the sensor (for example, relative movement between the in vivo portion and the ex vivo

portion, movement of the skin, and/or movement within the host (dermis or subcutaneous)) create stresses on the device and can produce noise in the sensor signal. It is believed that even small movements of the skin can translate to discomfort and/or motion-related artifact, which can be reduced or obviated by a flexible or articulated base. Thus, by providing flexibility and/or articulation of the device against the host's skin, better conformity of the sensor system **10** to the regular use and movements of the host can be achieved. Flexibility or articulation is believed to increase adhesion (with the use of an adhesive pad) of the mounting unit **14** onto the skin, thereby decreasing motion-related artifact that can otherwise translate from the host's movements and reduced sensor performance.

FIG. **4** is an exploded perspective view of a sensor system of a preferred embodiment, showing a mounting unit, an associated contact subassembly, and an electronics unit. In some embodiments, the contacts **28** are mounted on or in a subassembly hereinafter referred to as a contact subassembly **26** (see FIG. **5**A), which includes a contact holder **34** configured to fit within the base **24** of the mounting unit **14** and a hinge **38** that allows the contact subassembly **26** to pivot between a first position (for insertion) and a second position (for use) relative to the mounting unit **14**, which is described in more detail with reference to FIGS. **11** and **12**. The term "hinge" as used herein is a broad term and is used in its ordinary sense, including, without limitation, to refer to any of a variety of pivoting, articulating, and/or hinging mechanisms, such as an adhesive hinge, a sliding joint, and the like; the term hinge does not necessarily imply a fulcrum or fixed point about which the articulation occurs.

In certain embodiments, the mounting unit **14** is provided with an adhesive pad **8**, preferably disposed on the mounting unit's back surface and preferably including a releasable backing layer **9**. Thus, removing the backing layer **9** and pressing the base portion **24** of the mounting unit onto the host's skin adheres the mounting unit **14** to the host's skin. Additionally or alternatively, an adhesive pad can be placed over some or all of the sensor system after sensor insertion is complete to ensure adhesion, and optionally to ensure an airtight seal or watertight seal around the wound exit-site (or sensor insertion site) (not shown). Appropriate adhesive pads can be chosen and designed to stretch, elongate, conform to, and/or aerate the region (e.g., host's skin).

In preferred embodiments, the adhesive pad **8** is formed from spun-laced, open- or closed-cell foam, and/or non-woven fibers, and includes an adhesive disposed thereon, however a variety of adhesive pads appropriate for adhesion to the host's skin can be used, as is appreciated by one skilled in the art of medical adhesive pads. In some embodiments, a double-sided adhesive pad is used to adhere the mounting unit to the host's skin. In other embodiments, the adhesive pad includes a foam layer, for example, a layer wherein the foam is disposed between the adhesive pad's side edges and acts as a shock absorber.

In some embodiments, the surface area of the adhesive pad **8** is greater than the surface area of the mounting unit's back surface. Alternatively, the adhesive pad can be sized with substantially the same surface area as the back surface of the base portion. Preferably, the adhesive pad has a surface area on the side to be mounted on the host's skin that is greater than about 1, 1.25, 1.5, 1.75, 2, 2.25, or 2.5 times the surface area of the back surface **25** of the mounting unit base **24**. Such a greater surface area can increase adhesion between the mounting unit and the host's skin, minimize movement between the mounting unit and the host's skin, and/or protect the wound exit-site (sensor insertion site) from environmental and/or biological contamination. In some alternative embodiments, however, the adhesive pad can be smaller in surface area than the back surface assuming a sufficient adhesion can be accomplished.

In some embodiments, the adhesive pad **8** is substantially the same shape as the back surface **25** of the base **24**, although other shapes can also be advantageously employed, for example, butterfly-shaped, round, square, or rectangular. The adhesive pad backing can be designed for two-step release, for example, a primary release wherein only a portion of the adhesive pad is initially exposed to allow adjustable positioning of the device, and a secondary release wherein the remaining adhesive pad is later exposed to firmly and securely adhere the device to the host's skin once appropriately positioned. The adhesive pad is preferably waterproof. Preferably, a stretch-release adhesive pad is provided on the back surface of the base portion to enable easy release from the host's skin at the end of the useable life of the sensor, as is described in more detail with reference to FIGS. **10**A to **10**C.

In some circumstances, it has been found that a conventional bond between the adhesive pad and the mounting unit may not be sufficient, for example, due to humidity that can cause release of the adhesive pad from the mounting unit. Accordingly, in some embodiments, the adhesive pad can be bonded using a bonding agent activated by or accelerated by an ultraviolet, acoustic, radio frequency, or humidity cure. In some embodiments, a eutectic bond of first and second composite materials can form a strong adhesion. In some embodiments, the surface of the mounting unit can be pretreated utilizing ozone, plasma, chemicals, or the like, in order to enhance the bondability of the surface.

A bioactive agent is preferably applied locally at the insertion site (exit-site) prior to or during sensor insertion. Suitable bioactive agents include those which are known to discourage or prevent bacterial growth and infection, for example, anti-inflammatory agents, antimicrobials, antibiotics, or the like. It is believed that the diffusion or presence of a bioactive agent can aid in prevention or elimination of bacteria adjacent to the exit-site. Additionally or alternatively, the bioactive agent can be integral with or coated on the adhesive pad, or no bioactive agent at all is employed.

FIG. **5**A is an exploded perspective view of the contact subassembly **26** in one embodiment, showing its individual components. Preferably, a watertight (waterproof or water-resistant) sealing member **36**, also referred to as a sealing material or seal, fits within a contact holder **34** and provides a watertight seal configured to surround the electrical connection at the electrode terminals within the mounting unit in order to protect the electrodes (and the respective operable connection with the contacts of the electronics unit **16**) from damage due to moisture, humidity, dirt, and other external environmental factors. In one embodiment, the sealing member **36** is formed from an elastomeric material, such as silicone; however, a variety of other elastomeric or sealing materials can also be used, for example, silicone-polyurethane hybrids, polyurethanes, and polysulfides. Preferably, the sealing member is configured to compress within the contact subassembly when the electronics unit is mated to the mounting unit. In some embodiments, the sealing member **36** comprises a self-lubricating material, for example, self-lubricating silicone or other materials impregnated with or otherwise comprising a lubricant configured to be released during use. In some embodiments, the sealing member **36** includes a self-sealing material, for example, one that leaches out a sealant such as a silicone oil. In some embodiments, bumps, ridges, or other raised portions (not

shown), can be added to a component of the sensor system, such as to the contact subassembly **26** (e.g., housing adjacent to the sealing member), electronics unit **16** and/or sealing member **36** to provide additional compression and improve the seal formed around the contacts **28** and/or sensor **32** when the contacts **28** are mated to the sensor electronics.

Preferably, the sealing member is selected using a durometer. A durometer is an instrument used for measuring the indentation hardness of rubber, plastics, and other materials. Durometers are built to various standards from ASTM, DIN, JIS, and ISO. The hardness of plastics is most commonly measured by the Shore (Durometer) test or Rockwell hardness test. Both methods measure the resistance of plastics toward indentation and provide an empirical hardness value. Shore Hardness, using either the Shore A or Shore D scale, is the preferred method for rubbers/elastomers and is also commonly used for softer plastics such as polyolefins, fluoropolymers, and vinyls. The Shore A scale is used for softer rubbers while the Shore D scale is used for harder ones. In preferred embodiments, the Shore A scale is employed in connection with selection of a sealing member.

The Shore hardness is measured with a Durometer and sometimes referred to as "Durometer hardness." The hardness value is determined by the penetration of the Durometer indenter foot into the sample. Because of the resilience of rubbers and plastics, the indentation reading may change over time, so the indentation time is sometimes reported along with the hardness number. The ASTM test method designation for the Shore Durometer hardness test is ASTM D2240. The results obtained from this test are a useful measure of relative resistance to indentation of various grades of polymers.

Using a durometer in the selection of a sealing member enables selection of a material with optimal durometer hardness that balances the advantages of a lower durometer hardness with the advantages of a higher durometer hardness. For example, when a guide tube (e.g., cannula) is utilized to maintain an opening in a silicone sealing member prior to sensor insertion, a compression set (e.g., some retention of a compressed shape caused by compression of the material over time) within the silicone can result due to compression over time of the sealing member by the guide tube. Compression set can also result from certain sterilization procedures (e.g., radiation sterilization such as electron beam or gamma radiation). Unfortunately, in some circumstances, the compression set of the sealing member may cause gaps or incompleteness of contact between the sealing member and the contacts and/or sensor. In general, a lower durometer hardness provides a better conformation (e.g., seal) surrounding the contacts and/or sensor as compared to a higher durometer hardness. Additionally, a lower durometer hardness enables a design wherein less force is required to create the seal (e.g., to snap the electronics unit into the mounting unit, for example, as in the embodiment illustrated in FIG. **5**A) as compared to a higher durometer hardness, thereby increasing the ease of use of the device. However, the benefits of a lower durometer hardness silicone material must be balanced with potential disadvantages in manufacturing. For example, lower durometer hardness silicones are often produced by compounding with a silicone oil. In some circumstances, it is believed that some silicone oil may leach or migrate during manufacture and/or sterilization, which may corrupt aspects of the manufacturing process (e.g., adhesion of glues and/or effectiveness of coating processes). Additionally, a higher durometer hardness material generally provides greater stability of the material, which may reduce or avoid damage to the sealing member cause by pressure or other forces.

It is generally preferred that a sealing member **36** with a durometer hardness of from about 5 to about 80 Shore A is employed, more preferably a durometer hardness of from about 10 to about 50 Shore A, and even more preferably from about 20 to about 50 Shore A. In one embodiment, of a transcutaneous analyte sensor, the sealing member is fabricated using a silicone of about 20 Shore A to maximize the conformance of the seal around the contacts and/or sensor while minimizing the force required to compress the silicone for that conformance. In another embodiment, the sealing member is formed from a silicone of about 50 Shore A so as to provide increased strength of the sealing member (e.g., its resistance to compression). While a few representative examples have been provided above, one skilled in the art appreciates that higher or lower durometer hardness sealing material may also be suitable for use.

In one alternative embodiment, a sealing member **36** with a durometer hardness of about 10 Shore A is used. In this embodiment, the sealing material tends to "weep" out, further increasing conformance of the seal against the adjacent parts. In another alternative embodiment, a sealing material with a durometer hardness of about 0 (zero) Shore A is used as a sealant and/or in combination with a sealant, also referred to as a lubricant, which in some embodiments is a hydrophobic fluid filling material such as a grease, silicone, petroleum jelly, or the like. Preferably, the sensor and/or contacts are encased in a housing that contains the sealant, causing the material to "squeeze" around contacts and/or sensor. Any suitable hydrophobic fluid filling material can be employed. Especially preferred are synthetic or petroleum hydrocarbon-based materials, silicone-based materials, ester-based greases, and other pharmaceutical-grade materials.

In some embodiments, the sealing member can comprise a material that has been modified to enhance the desirable properties of the sealing member **36**. For example, one or more filler materials or stiffening agents such as glass beads, polymer beads, composite beads, beads comprising various inert materials, carbon black, talc, titanium oxide, silicone dioxide, and the like. In some embodiments, the filler material is incorporated into the sealing member material to mechanically stiffen the sealing member. In general, however, use of a filler material or stiffening agent in the sealing member material can provide a variety of enhanced properties including increased modulus of elasticity, crosslink density, hardness, and stiffness, and decreased creep, for example. In some alternative embodiments, gases are chemically (or otherwise) injected into the sealing member material. For example, the sealing material can comprise a polymeric foam (e.g., a polyurethane foam, a latex foam, a styrene-butadiene foam, and the like), or a dispersion of gas bubbles in a grease or jelly.

In alternative embodiments, the seal **36** is designed to form an interference fit with the electronics unit and can be formed from a variety of materials, for example, flexible plastics, or noble metals. One of ordinary skill in the art appreciates that a variety of designs can be employed to provide a seal surrounding electrical contacts such as described herein. For example, the contact holder **34** can be integrally designed as a part of the mounting unit, rather than as a separate piece thereof. Additionally or alternatively, a sealant can be provided in or around the sensor (e.g., within or on the contact subassembly or sealing member), such as is described in more detail with reference to FIGS. **12**A and

12B. In general, sealing materials with durometer hardnesses in the described ranges can provide improved sealing in a variety of sensor applications. For example, a sealing member as described in the preferred embodiments (e.g., selected using a durometer to ensure optimal durometer hardness, and the like) can be implemented adjacent to and/or to at least partially surrounding the sensor in a variety of sensor designs, including, for example, the sensor designs of the preferred embodiments, as well as a planar substrate such as described in U.S. Pat. No. 6,175,752.

In the illustrated embodiment of FIG. **5**A, the sealing member **36** is formed with a raised portion **37** surrounding the contacts **28**. The raised portion **37** enhances the interference fit surrounding the contacts **28** when the electronics unit **16** is mated to the mounting unit **14**. Namely, the raised portion surrounds each contact and presses against the electronics unit **16** to form a tight seal around the electronics unit. However, a variety of alternative sealing member configurations are described with reference to FIGS. **5**D to **5**H, below.

Contacts **28** fit within the seal **36** and provide for electrical connection between the sensor **32** and the electronics unit **16**. In general, the contacts are designed to ensure a stable mechanical and electrical connection of the electrodes that form the sensor **32** (see FIG. **6**A to **6**C) to mutually engaging contacts **28** thereon. A stable connection can be provided using a variety of known methods, for example, domed metallic contacts, cantilevered fingers, pogo pins, or the like, as is appreciated by one skilled in the art.

In preferred embodiments, the contacts **28** are formed from a conductive elastomeric material, such as a carbon black elastomer, through which the sensor **32** extends (see FIGS. **11**B and **12**B). Conductive elastomers are advantageously employed because their resilient properties create a natural compression against mutually engaging contacts, forming a secure press fit therewith. In some embodiments, conductive elastomers can be molded in such a way that pressing the elastomer against the adjacent contact performs a wiping action on the surface of the contact, thereby creating a cleaning action during initial connection. Additionally, in preferred embodiments, the sensor **32** extends through the contacts **28** wherein the sensor is electrically and mechanically secure by the relaxation of elastomer around the sensor (see FIGS. **8**A to **8**D).

In an alternative embodiment, a conductive, stiff plastic forms the contacts, which are shaped to comply upon application of pressure (for example, a leaf-spring shape). Contacts of such a configuration can be used instead of a metallic spring, for example, and advantageously avoid the need for crimping or soldering through compliant materials; additionally, a wiping action can be incorporated into the design to remove contaminants from the surfaces during connection. Non-metallic contacts can be advantageous because of their seamless manufacturability, robustness to thermal compression, non-corrosive surfaces, and native resistance to electrostatic discharge (ESD) damage due to their higher-than-metal resistance.

FIGS. **5**B and **5**C are perspective views of alternative contact configurations. FIG. **5**B is an illustration of a narrow contact configuration. FIG. **5**C is an illustration of a wide contact configuration. One skilled in the art appreciates that a variety of configurations are suitable for the contacts of the preferred embodiments, whether elastomeric, stiff plastic or other materials are used. In some circumstances, it can be advantageous to provide multiple contact configurations (such as illustrated in FIGS. **5**A to **5**C) to differentiate sensors from each other. In other words, the architecture of the contacts can include one or more configurations each designed (keyed) to fit with a particular electronics unit.

FIGS. **5**D to **5**H are schematic cross-sectional views of a portion of the contact subassembly; namely, a variety of alternative embodiments of the sealing member **36** are illustrated. In each of these embodiments (e.g., FIGS. **5**D to **5**H), a sensor **32** is shown, which is configured for operable connection to sensor electronics for measuring an analyte in a host such as described in more detail elsewhere herein. Additionally, two electrical contacts **28**, as described in more detail elsewhere herein, are configured to operably connect the sensor to the sensor electronics. Thus, the sealing member **36** in each of these alternative configurations (e.g., FIGS. **5**D to **5**H) at least partially surrounds the sensor and/or the electrical contacts to seal the electrical contacts from moisture when the sensor is operably connected to the sensor electronics.

FIG. **5**D is a schematic cross-sectional view of the sealing member **36** in an embodiment similar to FIG. **5**A, including gaps **400** that are maintained when the one or more electrical contacts are operably connected to the sensor electronics. Preferably, these air gaps provide for some flexibility of the sealing member **36** to deform or compress to seal the electrical contacts **28** from moisture or other environmental effects.

In certain circumstances, such as during sensor insertion or needle/guide tube retraction (see FIGS. **8**A to **8**D), a sealing member with a certain elasticity can be compressed or deformed by the insertion and/or retraction forces applied thereto. Accordingly in some embodiments, the sealing member is configured to be maintained (e.g., held substantially in place) on the housing (e.g., contact subassembly **26** or base **34**) without substantial translation, deformation, and/or compression (e.g., during sensor insertion). FIG. **5**D illustrates one such implementation, wherein one or more depressions **402** are configured to receive mating protrusions (e.g., on the base **34** of the contact subassembly **26**, not shown). A variety of male-female or other such mechanical structures can be implemented to hold the sealing member in place, as is appreciated by one skilled in the art. In one alternative embodiment, an adhesive (not shown) is configured to adhere the sealing member **36** to the housing (e.g., base **34** of the contact subassembly **26**) to provide substantially the same benefit of holding the sealing member during sensor insertion/retraction without substantial deformation, as described in more detail, above. In another embodiment, the base **34** of the contact subassembly **26** (or equivalent structure) comprises reinforcing mechanical supports configured to hold the sealing member as described above. One skilled in the art appreciates a variety of mechanical and/or chemical methods that can be implemented to maintain a sealing member substantially stationary (e.g., without substantial translation, deformation and/or compression) when compression and/or deformation forces are applied thereto. Although one exemplary embodiment is illustrated with reference to FIG. **5**D, a wide variety of systems and methods for holding the sealing member can be implemented with a sealing member of any particular design.

FIG. **5**E is a schematic cross-sectional view of the sealing member **36** in an alternative embodiment without gaps. In certain circumstances, full contact between mating members may be preferred.

In certain circumstances moisture may "wick" along the length of the sensor (e.g., from an exposed end) through the sealing member **36** to the contacts **28**. FIG. **5**F is a schematic cross-sectional view of a sealing member **36** in an alternative embodiment wherein one or more gaps **400** are pro-

vided. In this embodiment, the gaps **400** extend into the sealing member and encompass at least a portion of the sensor **32**. The gaps **400** or "deep wells" of FIG. **5**F are designed to interrupt the path that moisture may take, avoiding contact of the moisture at the contacts **28**. If moisture is able to travel along the path of the sensor, the abrupt change of surface tension at the opening **404** of the gap **400** in the sealing member **36** substantially deters the moisture from traveling to the contacts **28**.

FIG. **5**G is a schematic cross-sectional view of the sealing member **36** in another alternative embodiment wherein one or more gaps **400** are provided. In this embodiment, the gaps extend from the bottom side of the sealing member **36**, which can be helpful in maintaining a stable position of the contacts **28** and/or reduces "pumping" of air gaps in some situations.

In some embodiments, gaps **400** can be filled by a sealant, which also may be referred to as a lubricant, for example, oil, grease, or gel. In one exemplary embodiment, the sealant includes petroleum jelly and is used to provide a moisture barrier surrounding the sensor. Referring to FIG. **5**F, filling the gaps **400** with a sealant provides an additional moisture barrier to reduce or avoid moisture from traveling to the contacts **28**. Sealant can be used to fill gaps or crevices in any sealing member configuration.

In some sealing member configurations, it can be advantageous to provide a channel **406** through the sealing member **36** in order to create an additional pathway for sealant (e.g. lubricant) in order to expel air and/or to provide a path for excess sealant to escape. In some embodiments, more than one channel is provided.

FIG. **5**H is a schematic cross-sectional view of a sealing member **36** in an alternative embodiment wherein a large gap **400** is provided between the sealing member upper portion **408** and the sealing member lower portion **410**. These portions **408**, **410** may or may not be connected; however, they are configured to sandwich the sensor and sealant (e.g., grease) therebetween. The sealing member **36** illustrated with reference to FIG. **5**H can provide ease of manufacture and/or product assembly with a comprehensive sealing ability. Additional gaps (with or without sealant) can be provided in a variety of locations throughout the sealing member **36**; these additional gaps, for example, provide space for excess sealant.

Sensor

Preferably, the sensor **32** includes a distal portion **42**, also referred to as the in vivo portion, adapted to extend out of the mounting unit for insertion under the host's skin, and a proximal portion **40**, also referred to as an ex vivo portion, adapted to remain above the host's skin after sensor insertion and to operably connect to the electronics unit **16** via contacts **28**. Preferably, the sensor **32** includes two or more electrodes: a working electrode **44** and at least one additional electrode, which can function as a counter electrode and/or reference electrode, hereinafter referred to as the reference electrode **46**. A membrane system is preferably deposited over the electrodes, such as described in more detail with reference to FIGS. **6**A to **6**C, below.

FIG. **6**A is an expanded cutaway view of a proximal portion **40** of the sensor in one embodiment, showing working and reference electrodes. In the illustrated embodiments, the working and reference electrodes **44**, **46** extend through the contacts **28** to form electrical connection therewith (see FIGS. **11**B and **12**B). Namely, the working electrode **44** is in electrical contact with one of the contacts **28** and the reference electrode **46** is in electrical contact with the other contact **28**, which in turn provides for electrical connection with the electronics unit **16** when it is mated with the mounting unit **14**. Mutually engaging electrical contacts permit operable connection of the sensor **32** to the electronics unit **16** when connected to the mounting unit **14**; however other methods of electrically connecting the electronics unit **16** to the sensor **32** are also possible. In some alternative embodiments, for example, the reference electrode can be configured to extend from the sensor and connect to a contact at another location on the mounting unit (e.g., non-coaxially). Detachable connection between the mounting unit **14** and electronics unit **16** provides improved manufacturability, namely, the relatively inexpensive mounting unit **14** can be disposed of when replacing the sensor system after its usable life, while the relatively more expensive electronics unit **16** can be reused with multiple sensor systems.

In alternative embodiments, the contacts **28** are formed into a variety of alternative shapes and/or sizes. For example, the contacts **28** can be discs, spheres, cuboids, and the like. Furthermore, the contacts **28** can be designed to extend from the mounting unit in a manner that causes an interference fit within a mating cavity or groove of the electronics unit, forming a stable mechanical and electrical connection therewith.

FIG. **6**B is an expanded cutaway view of a distal portion of the sensor in one embodiment, showing working and reference electrodes. In preferred embodiments, the sensor is formed from a working electrode **44** and a reference electrode **46** helically wound around the working electrode **44**. An insulator **45** is disposed between the working and reference electrodes to provide necessary electrical insulation therebetween. Certain portions of the electrodes are exposed to enable electrochemical reaction thereon, for example, a window **43** can be formed in the insulator to expose a portion of the working electrode **44** for electrochemical reaction.

In preferred embodiments, each electrode is formed from a fine wire with a diameter of from about 0.001 or less to about 0.010 inches or more, for example, and is formed from, e.g., a plated insulator, a plated wire, or bulk electrically conductive material. Although the illustrated electrode configuration and associated text describe one preferred method of forming a transcutaneous sensor, a variety of known transcutaneous sensor configurations can be employed with the transcutaneous analyte sensor system of the preferred embodiments, such as U.S. Pat. No. 5,711,861 to Ward et al., U.S. Pat. No. 6,642,015 to Vachon et al., U.S. Pat. No. 6,654,625 to Say et al., U.S. Pat. No. 6,565,509 to Say et al., U.S. Pat. No. 6,514,718 to Heller, U.S. Pat. No. 6,465,066 to Essenpreis et al., U.S. Pat. No. 6,214,185 to Offenbacher et al., U.S. Pat. No. 5,310,469 to Cunningham et al., and U.S. Pat. No. 5,683,562 to Shaffer et al., U.S. Pat. No. 6,579,690 to Bonnecaze et al., U.S. Pat. No. 6,484,046 to Say et al., U.S. Pat. No. 6,512,939 to Colvin et al., U.S. Pat. No. 6,424,847 to Mastrototaro et al., U.S. Pat. No. 6,424,847 to Mastrototaro et al., for example. All of the above patents are not inclusive of all applicable analyte sensors; in general, it should be understood that the disclosed embodiments are applicable to a variety of analyte sensor configurations. Much of the description of the preferred embodiments, for example the membrane system described below, can be implemented not only with in vivo sensors, but also with in vitro sensors, such as blood glucose meters (SMBG).

In preferred embodiments, the working electrode comprises a wire formed from a conductive material, such as platinum, platinum-iridium, palladium, graphite, gold, car-

bon, conductive polymer, alloys, or the like. Although the electrodes can by formed by a variety of manufacturing techniques (bulk metal processing, deposition of metal onto a substrate, or the like), it can be advantageous to form the electrodes from plated wire (e.g., platinum on steel wire) or bulk metal (e.g., platinum wire). It is believed that electrodes formed from bulk metal wire provide superior performance (e.g., in contrast to deposited electrodes), including increased stability of assay, simplified manufacturability, resistance to contamination (e.g., which can be introduced in deposition processes), and improved surface reaction (e.g., due to purity of material) without peeling or delamination.

The working electrode **44** is configured to measure the concentration of an analyte. In an enzymatic electrochemical sensor for detecting glucose, for example, the working electrode measures the hydrogen peroxide produced by an enzyme catalyzed reaction of the analyte being detected and creates a measurable electronic current For example, in the detection of glucose wherein glucose oxidase produces hydrogen peroxide as a byproduct, hydrogen peroxide reacts with the surface of the working electrode producing two protons ($2H^+$), two electrons ($2e^-$) and one molecule of oxygen (O_2), which produces the electronic current being detected.

In preferred embodiments, the working electrode **44** is covered with an insulating material **45**, for example, a non-conductive polymer. Dip-coating, spray-coating, vapor-deposition, or other coating or deposition techniques can be used to deposit the insulating material on the working electrode. In one embodiment, the insulating material comprises parylene, which can be an advantageous polymer coating for its strength, lubricity, and electrical insulation properties. Generally, parylene is produced by vapor deposition and polymerization of para-xylylene (or its substituted derivatives). While not wishing to be bound by theory, it is believed that the lubricious (e.g., smooth) coating (e.g., parylene) on the sensors of the preferred embodiments contributes to minimal trauma and extended sensor life. While parylene coatings are generally preferred, any suitable insulating material can be used, for example, fluorinated polymers, polyethyleneterephthalate, polyurethane, polyimide, other nonconducting polymers, or the like. Glass or ceramic materials can also be employed. Other materials suitable for use include surface energy modified coating systems such as are marketed under the trade names AMC18, AMC148, AMC141, and AMC321 by Advanced Materials Components Express of Bellafonte, Pa. In some alternative embodiments, however, the working electrode may not require a coating of insulator.

The reference electrode **46**, which can function as a reference electrode alone, or as a dual reference and counter electrode, is formed from silver, silver/silver chloride, or the like. Preferably, the reference electrode **46** is juxtapositioned and/or twisted with or around the working electrode **44**; however other configurations are also possible (e.g., an intradermal or on-skin reference electrode). In the illustrated embodiments, the reference electrode **46** is helically wound around the working electrode **44**. The assembly of wires is then optionally coated or adhered together with an insulating material, similar to that described above, so as to provide an insulating attachment.

In some embodiments, a silver wire is formed onto the sensor as described above, and subsequently chloridized to form silver/silver chloride reference electrode. Advantageously, chloridizing the silver wire as described herein enables the manufacture of a reference electrode with optimal in vivo performance. Namely, by controlling the quantity and amount of chloridization of the silver to form silver/silver chloride, improved break-in time, stability of the reference electrode, and extended life has been shown with the preferred embodiments. Additionally, use of silver chloride as described above allows for relatively inexpensive and simple manufacture of the reference electrode.

In embodiments wherein an outer insulator is disposed, a portion of the coated assembly structure can be stripped or otherwise removed, for example, by hand, excimer lasing, chemical etching, laser ablation, grit-blasting (e.g., with sodium bicarbonate or other suitable grit), or the like, to expose the electroactive surfaces. Alternatively, a portion of the electrode can be masked prior to depositing the insulator in order to maintain an exposed electroactive surface area. In one exemplary embodiment, grit blasting is implemented to expose the electroactive surfaces, preferably utilizing a grit material that is sufficiently hard to ablate the polymer material, while being sufficiently soft so as to minimize or avoid damage to the underlying metal electrode (e.g., a platinum electrode). Although a variety of "grit" materials can be used (e.g., sand, talc, walnut shell, ground plastic, sea salt, and the like), in some preferred embodiments, sodium bicarbonate is an advantageous grit-material because it is sufficiently hard to ablate, e.g., a parylene coating without damaging, e.g., an underlying platinum conductor. One additional advantage of sodium bicarbonate blasting includes its polishing action on the metal as it strips the polymer layer, thereby eliminating a cleaning step that might otherwise be necessary.

In the embodiment illustrated in FIG. **6**B, a radial window **43** is formed through the insulating material **45** to expose a circumferential electroactive surface of the working electrode. Additionally, sections **41** of electroactive surface of the reference electrode are exposed. For example, the 41 sections of electroactive surface can be masked during deposition of an outer insulating layer or etched after deposition of an outer insulating layer.

In some applications, cellular attack or migration of cells to the sensor can cause reduced sensitivity and/or function of the device, particularly after the first day of implantation. However, when the exposed electroactive surface is distributed circumferentially about the sensor (e.g., as in a radial window), the available surface area for reaction can be sufficiently distributed so as to minimize the effect of local cellular invasion of the sensor on the sensor signal. Alternatively, a tangential exposed electroactive window can be formed, for example, by stripping only one side of the coated assembly structure. In other alternative embodiments, the window can be provided at the tip of the coated assembly structure such that the electroactive surfaces are exposed at the tip of the sensor. Other methods and configurations for exposing electroactive surfaces can also be employed.

In some embodiments, the working electrode has a diameter of from about 0.001 inches or less to about 0.010 inches or more, preferably from about 0.002 inches to about 0.008 inches, and more preferably from about 0.004 inches to about 0.005 inches. The length of the window can be from about 0.1 mm (about 0.004 inches) or less to about 2 mm (about 0.078 inches) or more, and preferably from about 0.5 mm (about 0.02 inches) to about 0.75 mm (0.03 inches). In such embodiments, the exposed surface area of the working electrode is preferably from about 0.000013 in^2 (0.0000839 cm^2) or less to about 0.0025 in^2 (0.016129 cm^2) or more (assuming a diameter of from about 0.001 inches to about 0.010 inches and a length of from about 0.004 inches to about 0.078 inches). The preferred exposed surface area of the working electrode is selected to produce an analyte

signal with a current in the picoAmp range, such as is described in more detail elsewhere herein. However, a current in the picoAmp range can be dependent upon a variety of factors, for example the electronic circuitry design (e.g., sample rate, current draw, A/D converter bit resolution, etc.), the membrane system (e.g., permeability of the analyte through the membrane system), and the exposed surface area of the working electrode. Accordingly, the exposed electroactive working electrode surface area can be selected to have a value greater than or less than the above-described ranges taking into consideration alterations in the membrane system and/or electronic circuitry. In preferred embodiments of a glucose sensor, it can be advantageous to minimize the surface area of the working electrode while maximizing the diffusivity of glucose in order to optimize the signal-to-noise ratio while maintaining sensor performance in both high and low glucose concentration ranges.

In some alternative embodiments, the exposed surface area of the working (and/or other) electrode can be increased by altering the cross-section of the electrode itself. For example, in some embodiments the cross-section of the working electrode can be defined by a cross, star, cloverleaf, ribbed, dimpled, ridged, irregular, or other non-circular configuration; thus, for any predetermined length of electrode, a specific increased surface area can be achieved (as compared to the area achieved by a circular cross-section). Increasing the surface area of the working electrode can be advantageous in providing an increased signal responsive to the analyte concentration, which in turn can be helpful in improving the signal-to-noise ratio, for example.

In some alternative embodiments, additional electrodes can be included within the assembly, for example, a three-electrode system (working, reference, and counter electrodes) and/or an additional working electrode (e.g., an electrode which can be used to generate oxygen, which is configured as a baseline subtracting electrode, or which is configured for measuring additional analytes). U.S. Publication No. US-2005-0161346-A1 and U.S. Publication No. US-2005-0143635-A1 describe some systems and methods for implementing and using additional working, counter, and/or reference electrodes. In one implementation wherein the sensor comprises two working electrodes, the two working electrodes are juxtapositioned (e.g., extend parallel to each other), around which the reference electrode is disposed (e.g., helically wound). In some embodiments wherein two or more working electrodes are provided, the working electrodes can be formed in a double-, triple-, quad-, etc. helix configuration along the length of the sensor (for example, surrounding a reference electrode, insulated rod, or other support structure). The resulting electrode system can be configured with an appropriate membrane system, wherein the first working electrode is configured to measure a first signal comprising glucose and baseline and the additional working electrode is configured to measure a baseline signal consisting of baseline only (e.g., configured to be substantially similar to the first working electrode without an enzyme disposed thereon). In this way, the baseline signal can be subtracted from the first signal to produce a glucose-only signal that is substantially not subject to fluctuations in the baseline and/or interfering species on the signal.

Although the preferred embodiments illustrate one electrode configuration including one bulk metal wire helically wound around another bulk metal wire, other electrode configurations are also contemplated. In an alternative embodiment, the working electrode comprises a tube with a reference electrode disposed or coiled inside, including an insulator therebetween. Alternatively, the reference electrode comprises a tube with a working electrode disposed or coiled inside, including an insulator therebetween. In another alternative embodiment, a polymer (e.g., insulating) rod is provided, wherein the electrodes are deposited (e.g., electro-plated) thereon. In yet another alternative embodiment, a metallic (e.g., steel) rod is provided, coated with an insulating material, onto which the working and reference electrodes are deposited. In yet another alternative embodiment, one or more working electrodes are helically wound around a reference electrode.

Preferably, the electrodes and membrane systems of the preferred embodiments are coaxially formed, namely, the electrodes and/or membrane system all share the same central axis. While not wishing to be bound by theory, it is believed that a coaxial design of the sensor enables a symmetrical design without a preferred bend radius. Namely, in contrast to prior art sensors comprising a substantially planar configuration that can suffer from regular bending about the plane of the sensor, the coaxial design of the preferred embodiments do not have a preferred bend radius and therefore are not subject to regular bending about a particular plane (which can cause fatigue failures and the like). However, non-coaxial sensors can be implemented with the sensor system of the preferred embodiments.

In addition to the above-described advantages, the coaxial sensor design of the preferred embodiments enables the diameter of the connecting end of the sensor (proximal portion) to be substantially the same as that of the sensing end (distal portion) such that the needle is able to insert the sensor into the host and subsequently slide back over the sensor and release the sensor from the needle, without slots or other complex multi-component designs.

In one such alternative embodiment, the two wires of the sensor are held apart and configured for insertion into the host in proximal but separate locations. The separation of the working and reference electrodes in such an embodiment can provide additional electrochemical stability with simplified manufacture and electrical connectivity. It is appreciated by one skilled in the art that a variety of electrode configurations can be implemented with the preferred embodiments.

In some embodiments, the sensor includes an antimicrobial portion configured to extend through the exit-site when the sensor is implanted in the host. Namely, the sensor is designed with in vivo and ex vivo portions as described in more detail elsewhere herein; additionally, the sensor comprises a transition portion, also referred to as an antimicrobial portion, located between the in vivo and ex vivo portions **42**, **40**. The antimicrobial portion is designed to provide antimicrobial effects to the exit-site and adjacent tissue when implanted in the host.

In some embodiments, the antimicrobial portion comprises silver, e.g., the portion of a silver reference electrode that is configured to extend through the exit-site when implanted. Although exit-site infections are a common adverse occurrence associated with some conventional transcutaneous medical devices, the devices of preferred embodiments are designed at least in part to minimize infection, to minimize irritation, and/or to extend the duration of implantation of the sensor by utilizing a silver reference electrode to extend through the exit-site when implanted in a patient. While not wishing to be bound by theory, it is believed that the silver may reduce local tissue infections (within the tissue and at the exit-site); namely, steady release of molecular quantities of silver is believed to have an antimicrobial effect in biological tissue (e.g., reduc-

ing or preventing irritation and infection), also referred to as passive antimicrobial effects. Although one example of passive antimicrobial effects is described herein, one skilled in the art can appreciate a variety of passive anti-microbial systems and methods that can be implemented with the preferred embodiments. Additionally, it is believed that antimicrobial effects can contribute to extended life of a transcutaneous analyte sensor, enabling a functional lifetime past a few days, e.g., seven days or longer.

In some embodiments, active antimicrobial systems and methods are provided in the sensor system in order to further enhance the antimicrobial effects at the exit-site. In one such embodiment, an auxiliary silver wire is disposed on or around the sensor, wherein the auxiliary silver wire is connected to electronics and configured to pass a current sufficient to enhance its antimicrobial properties (active antimicrobial effects), as is appreciated by one skilled in the art. The current can be passed continuously or intermittently, such that sufficient antimicrobial properties are provided. Although one example of active antimicrobial effects is described herein, one skilled in the art can appreciate a variety of active anti-microbial systems and methods that can be implemented with the preferred embodiments.

Anchoring Mechanism

It is preferred that the sensor remains substantially stationary within the tissue of the host, such that migration or motion of the sensor with respect to the surrounding tissue is minimized. Migration or motion is believed to cause inflammation at the sensor implant site due to irritation, and can also cause noise on the sensor signal due to motion-related artifact, for example. Therefore, it can be advantageous to provide an anchoring mechanism that provides support for the sensor's in vivo portion to avoid the above-mentioned problems. Combining advantageous sensor geometry with an advantageous anchoring minimizes additional parts and allows for an optimally small or low profile design of the sensor. In one embodiment the sensor includes a surface topography, such as the helical surface topography provided by the reference electrode surrounding the working electrode. In alternative embodiments, a surface topography could be provided by a roughened surface, porous surface (e.g. porous parylene), ridged surface, or the like. Additionally (or alternatively), the anchoring can be provided by prongs, spines, barbs, wings, hooks, a bulbous portion (for example, at the distal end), an S-bend along the sensor, a rough surface topography, a gradually changing diameter, combinations thereof, or the like, which can be used alone or in combination with the helical surface topography to stabilize the sensor within the subcutaneous tissue.

Variable Stiffness

As described above, conventional transcutaneous devices are believed to suffer from motion artifact associated with host movement when the host is using the device. For example, when a transcutaneous analyte sensor is inserted into the host, various movements on the sensor (for example, relative movement within and between the subcutaneous space, dermis, skin, and external portions of the sensor) create stresses on the device, which is known to produce artifacts on the sensor signal. Accordingly, there are different design considerations (for example, stress considerations) on various sections of the sensor. For example, the distal portion **42** of the sensor can benefit in general from greater flexibility as it encounters greater mechanical stresses caused by movement of the tissue within the patient and relative movement between the in vivo and ex vivo portions of the sensor. On the other hand, the proximal portion **40** of the sensor can benefit in general from a stiffer, more robust design to ensure structural integrity and/or reliable electrical connections. Additionally, in some embodiments wherein a needle is retracted over the proximal portion **40** of the device (see FIGS. **7** to **9**), a stiffer design can minimize crimping of the sensor and/or ease in retraction of the needle from the sensor. Thus, by designing greater flexibility into the in vivo (distal) portion **42**, the flexibility is believed to compensate for patient movement, and noise associated therewith. By designing greater stiffness into the ex vivo (proximal) portion **40**, column strength (for retraction of the needle over the sensor), electrical connections, and integrity can be enhanced. In some alternative embodiments, a stiffer distal end and/or a more flexible proximal end can be advantageous as described in U.S. Publication No. US-2006-0015024-A1.

The preferred embodiments provide a distal portion **42** of the sensor **32** designed to be more flexible than a proximal portion **40** of the sensor. The variable stiffness of the preferred embodiments can be provided by variable pitch of any one or more helically wound wires of the device, variable cross-section of any one or more wires of the device, and/or variable hardening and/or softening of any one or more wires of the device, such as is described in more detail with reference to U.S. Publication No. US-2006-0015024-A1.

Membrane System

FIG. **6**C is a cross-sectional view through the sensor on line C-C of FIG. **6**B showing the exposed electroactive surface of the working electrode surrounded by the membrane system in one embodiment. Preferably, a membrane system is deposited over at least a portion of the electroactive surfaces of the sensor **32** (working electrode and optionally reference electrode) and provides protection of the exposed electrode surface from the biological environment, diffusion resistance (limitation) of the analyte if needed, a catalyst for enabling an enzymatic reaction, limitation or blocking of interferents, and/or hydrophilicity at the electrochemically reactive surfaces of the sensor interface. Some examples of suitable membrane systems are described in U.S. Publication No. US-2005-0245799-A1.

In general, the membrane system includes a plurality of domains, for example, an electrode domain **47**, an interference domain **48**, an enzyme domain **49** (for example, including glucose oxidase), and a resistance domain **50**, as shown in FIG. **6**C, and can include a high oxygen solubility domain, and/or a bioprotective domain (not shown), such as is described in more detail in U.S. Publication No. US-2005-0245799-A1. The membrane system can be deposited on the exposed electroactive surfaces using known thin film techniques (for example, vapor deposition, spraying, electrodepositing, dipping, or the like). In alternative embodiments, however, other vapor deposition processes (e.g., physical and/or chemical vapor deposition processes) can be useful for providing one or more of the insulating and/or membrane layers, including ultrasonic vapor deposition, electrostatic deposition, evaporative deposition, deposition by sputtering, pulsed laser deposition, high velocity oxygen fuel deposition, thermal evaporator deposition, electron beam evaporator deposition, deposition by reactive sputtering molecular beam epitaxy, atmospheric pressure chemical vapor deposition (CVD), atomic layer CVD, hot wire CVD, low-pressure CVD, microwave plasma-assisted CVD, plasma-enhanced CVD, rapid thermal CVD, remote plasma-enhanced CVD, and ultra-high vacuum CVD, for example. However, the membrane system can be disposed over (or deposited on) the electroactive surfaces using any known method, as will be appreciated by one skilled in the art.

In some embodiments, one or more domains of the membrane systems are formed from materials such as described above in connection with the porous layer, such as silicone, polytetrafluoroethylene, polyethylene-co-tetrafluoroethylene, polyolefin, polyester, polycarbonate, biostable polytetrafluoroethylene, homopolymers, copolymers, terpolymers of polyurethanes, polypropylene (PP), polyvinylchloride (PVC), polyvinylidene fluoride (PVDF), polybutylene terephthalate (PBT), polymethylmethacrylate (PMMA), polyether ether ketone (PEEK), polyurethanes, cellulosic polymers, polysulfones and block copolymers thereof including, for example, di-block, tri-block, alternating, random and graft copolymers. U.S. Publication No. US-2005-0245799-A1 describes biointerface and membrane system configurations and materials that can be applied to the preferred embodiments.

Electrode Domain

In selected embodiments, the membrane system comprises an electrode domain. The electrode domain **47** is provided to ensure that an electrochemical reaction occurs between the electroactive surfaces of the working electrode and the reference electrode, and thus the electrode domain **47** is preferably situated more proximal to the electroactive surfaces than the interference and/or enzyme domain. Preferably, the electrode domain includes a coating that maintains a layer of water at the electrochemically reactive surfaces of the sensor. In other words, the electrode domain is present to provide an environment between the surfaces of the working electrode and the reference electrode which facilitates an electrochemical reaction between the electrodes. For example, a humectant in a binder material can be employed as an electrode domain; this allows for the full transport of ions in the aqueous environment. The electrode domain can also assist in stabilizing the operation of the sensor by accelerating electrode start-up and drifting problems caused by inadequate electrolyte. The material that forms the electrode domain can also provide an environment that protects against pH-mediated damage that can result from the formation of a large pH gradient due to the electrochemical activity of the electrodes.

In one embodiment, the electrode domain **47** includes a flexible, water-swellable, hydrogel film having a "dry film" thickness of from about 0.05 micron or less to about 20 microns or more, more preferably from about 0.05, 0.1, 0.15, 0.2, 0.25, 0.3, 0.35, 0.4, 0.45, 0.5, 1, 1.5, 2, 2.5, 3, or 3.5 microns to about 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 19.5 microns, and more preferably still from about 2, 2.5 or 3 microns to about 3.5, 4, 4.5, or 5 microns. "Dry film" thickness refers to the thickness of a cured film cast from a coating formulation by standard coating techniques.

In certain embodiments, the electrode domain **47** is formed of a curable mixture of a urethane polymer and a hydrophilic polymer. Particularly preferred coatings are formed of a polyurethane polymer having carboxylate or hydroxyl functional groups and non-ionic hydrophilic polyether segments, wherein the polyurethane polymer is crosslinked with a water soluble carbodiimide (e.g., 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC)) in the presence of polyvinylpyrrolidone and cured at a moderate temperature of about 50° C.

In some preferred embodiments, the electrode domain **47** is formed from a hydrophilic polymer such as polyvinylpyrrolidone (PVP). An electrode domain formed from PVP has been shown to reduce break-in time of analyte sensors; for example, a glucose sensor utilizing a cellulosic-based interference domain such as described in more detail below.

Preferably, the electrode domain is deposited by vapor deposition, spray coating, dip coating, or other thin film techniques on the electroactive surfaces of the sensor. In one preferred embodiment, the electrode domain is formed by dip-coating the electroactive surfaces in an electrode layer solution and curing the domain for a time of from about 15 minutes to about 30 minutes at a temperature of from about 40° C. to about 55° C. (and can be accomplished under vacuum (e.g., 20 to 30 mmHg)). In embodiments wherein dip-coating is used to deposit the electrode domain, a preferred insertion rate of from about 1 to about 3 inches per minute into the electrode layer solution, with a preferred dwell time of from about 0.5 to about 2 minutes in the electrode layer solution, and a preferred withdrawal rate of from about 0.25 to about 2 inches per minute from the electrode layer solution provide a functional coating. However, values outside of those set forth above can be acceptable or even desirable in certain embodiments, for example, depending upon solution viscosity and solution surface tension, as is appreciated by one skilled in the art. In one embodiment, the electroactive surfaces of the electrode system are dip-coated one time (one layer) and cured at 50° C. under vacuum for 20 minutes.

Although an independent electrode domain **47** is described herein, in some embodiments sufficient hydrophilicity can be provided in the interference domain and/or enzyme domain (the domain adjacent to the electroactive surfaces) so as to provide for the full transport of ions in the aqueous environment (e.g. without a distinct electrode domain). In these embodiments, an electrode domain is not necessary.

Interference Domain

Interferents are molecules or other species that are reduced or oxidized at the electrochemically reactive surfaces of the sensor, either directly or via an electron transfer agent, to produce a false positive analyte signal. In preferred embodiments, an interference domain **48** is provided that substantially restricts, resists, or blocks the flow of one or more interfering species. Some known interfering species for a glucose sensor, as described in more detail above, include acetaminophen, ascorbic acid, bilirubin, cholesterol, creatinine, dopamine, ephedrine, ibuprofen, L-dopa, methyl dopa, salicylate, tetracycline, tolazamide, tolbutamide, triglycerides, and uric acid. In general, the interference domain of the preferred embodiments is less permeable to one or more of the interfering species than to the analyte, e.g., glucose.

In one embodiment, the interference domain **48** is formed from one or more cellulosic derivatives. In general, cellulosic derivatives include polymers such as cellulose acetate, cellulose acetate butyrate, 2-hydroxyethyl cellulose, cellulose acetate phthalate, cellulose acetate propionate, cellulose acetate trimellitate, and the like.

In one preferred embodiment, the interference domain **48** is formed from cellulose acetate butyrate. Cellulose acetate butyrate with a molecular weight of from about 10,000 daltons to about 75,000 daltons, preferably from about 15,000, 20,000, or 25,000 daltons to about 50,000, 55,000, 60,000, 65,000, or 70,000 daltons, and more preferably about 20,000 daltons is employed. In certain embodiments, however, higher or lower molecular weights can be preferred. Additionally, a casting solution or dispersion of cellulose acetate butyrate at a weight percent of from about 15% to about 25%, preferably from about 15%, 16%, 17%, 18%, 19% to about 20%, 21%, 22%, 23%, 24% or 25%, and more preferably about 18% is preferred. Preferably, the casting solution includes a solvent or solvent system, for

example an acetone:ethanol solvent system. Higher or lower concentrations can be preferred in certain embodiments. A plurality of layers of cellulose acetate butyrate can be advantageously combined to form the interference domain in some embodiments, for example, three layers can be employed. It can be desirable to employ a mixture of cellulose acetate butyrate components with different molecular weights in a single solution, or to deposit multiple layers of cellulose acetate butyrate from different solutions comprising cellulose acetate butyrate of different molecular weights, different concentrations, and/or different chemistries (e.g., functional groups). It can also be desirable to include additional substances in the casting solutions or dispersions, e.g., functionalizing agents, crosslinking agents, other polymeric substances, substances capable of modifying the hydrophilicity/hydrophobicity of the resulting layer, and the like.

In one alternative embodiment, the interference domain **48** is formed from cellulose acetate. Cellulose acetate with a molecular weight of from about 30,000 daltons or less to about 100,000 daltons or more, preferably from about 35,000, 40,000, or 45,000 daltons to about 55,000, 60,000, 65,000, 70,000, 75,000, 80,000, 85,000, 90,000, or 95,000 daltons, and more preferably about 50,000 daltons is preferred. Additionally, a casting solution or dispersion of cellulose acetate at a weight percent of about 3% to about 10%, preferably from about 3.5%, 4.0%, 4.5%, 5.0%, 5.5%, 6.0%, or 6.5% to about 7.5%, 8.0%, 8.5%, 9.0%, or 9.5%, and more preferably about 8% is preferred. In certain embodiments, however, higher or lower molecular weights and/or cellulose acetate weight percentages can be preferred. It can be desirable to employ a mixture of cellulose acetates with molecular weights in a single solution, or to deposit multiple layers of cellulose acetate from different solutions comprising cellulose acetates of different molecular weights, different concentrations, or different chemistries (e.g., functional groups). It can also be desirable to include additional substances in the casting solutions or dispersions such as described in more detail above.

Layer(s) prepared from combinations of cellulose acetate and cellulose acetate butyrate, or combinations of layer(s) of cellulose acetate and layer(s) of cellulose acetate butyrate can also be employed to form the interference domain **48**.

In some alternative embodiments, additional polymers, such as Nafion®, can be used in combination with cellulosic derivatives to provide equivalent and/or enhanced function of the interference domain **48**. As one example, a 5 wt % Nafion® casting solution or dispersion can be used in combination with a 8 wt % cellulose acetate casting solution or dispersion, e.g., by dip coating at least one layer of cellulose acetate and subsequently dip coating at least one layer Nafion® onto a needle-type sensor such as described with reference to the preferred embodiments. Any number of coatings or layers formed in any order may be suitable for forming the interference domain of the preferred embodiments.

In some alternative embodiments, more than one cellulosic derivative can be used to form the interference domain **48** of the preferred embodiments. In general, the formation of the interference domain on a surface utilizes a solvent or solvent system in order to solvate the cellulosic derivative (or other polymer) prior to film formation thereon. In preferred embodiments, acetone and ethanol are used as solvents for cellulose acetate; however one skilled in the art appreciates the numerous solvents that are suitable for use with cellulosic derivatives (and other polymers). Additionally, one skilled in the art appreciates that the preferred relative amounts of solvent can be dependent upon the cellulosic derivative (or other polymer) used, its molecular weight, its method of deposition, its desired thickness, and the like. However, a percent solute of from about 1% to about 25% is preferably used to form the interference domain solution so as to yield an interference domain having the desired properties. The cellulosic derivative (or other polymer) used, its molecular weight, method of deposition, and desired thickness can be adjusted, depending upon one or more other of the parameters, and can be varied accordingly as is appreciated by one skilled in the art.

In some alternative embodiments, other polymer types that can be utilized as a base material for the interference domain **48** including polyurethanes, polymers having pendant ionic groups, and polymers having controlled pore size, for example. In one such alternative embodiment, the interference domain includes a thin, hydrophobic membrane that is non-swellable and restricts diffusion of low molecular weight species. The interference domain **48** is permeable to relatively low molecular weight substances, such as hydrogen peroxide, but restricts the passage of higher molecular weight substances, including glucose and ascorbic acid. Other systems and methods for reducing or eliminating interference species that can be applied to the membrane system of the preferred embodiments are described in U.S. Publication No. US-2005-0115832-A1, U.S. Publication No. US-2005-0176136-A1, U.S. Publication No. US-2005-0161346-A1, and U.S. Publication No. US-2005-0143635-A1. In some alternative embodiments, a distinct interference domain is not included.

In preferred embodiments, the interference domain **48** is deposited directly onto the electroactive surfaces of the sensor for a domain thickness of from about 0.05 microns or less to about 20 microns or more, more preferably from about 0.05, 0.1, 0.15, 0.2, 0.25, 0.3, 0.35, 0.4, 0.45, 0.5, 1, 1.5, 2, 2.5, 3, or 3.5 microns to about 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 19.5 microns, and more preferably still from about 1, 1.5 or 2 microns to about 2.5 or 3 microns. Thicker membranes can also be desirable in certain embodiments, but thinner membranes are generally preferred because they have a lower impact on the rate of diffusion of hydrogen peroxide from the enzyme membrane to the electrodes.

In general, the membrane systems of the preferred embodiments can be formed and/or deposited on the exposed electroactive surfaces (e.g., one or more of the working and reference electrodes) using known thin film techniques (for example, casting, spray coating, drawing down, electro-depositing, dip coating, and the like), however casting or other known application techniques can also be utilized. Preferably, the interference domain is deposited by vapor deposition, spray coating, or dip coating. In one exemplary embodiment of a needle-type (transcutaneous) sensor such as described herein, the interference domain is formed by dip coating the sensor into an interference domain solution using an insertion rate of from about 20 inches/min to about 60 inches/min, preferably 40 inches/min, a dwell time of from about 0 minute to about 5 seconds, preferably 0 seconds, and a withdrawal rate of from about 20 inches/minute to about 60 inches/minute, preferably about 40 inches/minute, and curing (drying) the domain from about 1 minute to about 30 minutes, preferably from about 3 minutes to about 15 minutes (and can be accomplished at room temperature or under vacuum (e.g., 20 to 30 mmHg)). In one exemplary embodiment including cellulose acetate butyrate interference domain, a 3-minute cure (i.e., dry) time is preferred between each layer applied. In another exemplary

embodiment employing a cellulose acetate interference domain, a 15 minute cure (i.e., dry) time is preferred between each layer applied.

The dip process can be repeated at least one time and up to 10 times or more. The preferred number of repeated dip processes depends upon the cellulosic derivative(s) used, their concentration, conditions during deposition (e.g., dipping) and the desired thickness (e.g., sufficient thickness to provide functional blocking of (or resistance to) certain interferents), and the like. In some embodiments, 1 to 3 microns may be preferred for the interference domain thickness; however, values outside of these can be acceptable or even desirable in certain embodiments, for example, depending upon viscosity and surface tension, as is appreciated by one skilled in the art. In one exemplary embodiment, an interference domain is formed from three layers of cellulose acetate butyrate. In another exemplary embodiment, an interference domain is formed from 10 layers of cellulose acetate. In alternative embodiments, the interference domain can be formed using any known method and combination of cellulose acetate and cellulose acetate butyrate, as will be appreciated by one skilled in the art.

In some embodiments, the electroactive surface can be cleaned prior to application of the interference domain **48**. In some embodiments, the interference domain **48** of the preferred embodiments can be useful as a bioprotective or biocompatible domain, namely, a domain that interfaces with host tissue when implanted in an animal (e.g., a human) due to its stability and biocompatibility.

Enzyme Domain

In preferred embodiments, the membrane system further includes an enzyme domain **49** disposed more distally from the electroactive surfaces than the interference domain **48**; however other configurations can be desirable. In the preferred embodiments, the enzyme domain provides an enzyme to catalyze the reaction of the analyte and its co-reactant, as described in more detail below. In the preferred embodiments of a glucose sensor, the enzyme domain includes glucose oxidase; however other oxidases, for example, galactose oxidase or uricase oxidase, can also be used.

For an enzyme-based electrochemical glucose sensor to perform well, the sensor's response is preferably limited by neither enzyme activity nor co-reactant concentration. Because enzymes, including glucose oxidase, are subject to deactivation as a function of time even in ambient conditions, this behavior is compensated for in forming the enzyme domain. Preferably, the enzyme domain is constructed of aqueous dispersions of colloidal polyurethane polymers including the enzyme. However, in alternative embodiments the enzyme domain is constructed from an oxygen enhancing material, for example, silicone, or fluorocarbon, in order to provide a supply of excess oxygen during transient ischemia. Preferably, the enzyme is immobilized within the domain. See, e.g., U.S. Publication No. US-2005-0054909-A1.

In preferred embodiments, the enzyme domain is deposited onto the interference domain for a domain thickness of from about 0.05 micron or less to about 20 microns or more, more preferably from about 0.05, 0.1, 0.15, 0.2, 0.25, 0.3, 0.35, 0.4, 0.45, 0.5, 1, 1.5, 2, 2.5, 3, or 3.5 microns to about 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 19.5 microns, and more preferably still from about 2, 2.5 or 3 microns to about 3.5, 4, 4.5, or 5 microns. However in some embodiments, the enzyme domain can be deposited directly onto the electroactive surfaces. Preferably, the enzyme domain is deposited by spray or dip coating. In one embodiment of needle-type (transcutaneous) sensor such as described herein, the enzyme domain is formed by dip coating the interference domain coated sensor into an enzyme domain solution and curing the domain for from about 15 to about 30 minutes at a temperature of from about 40° C. to about 55° C. (and can be accomplished under vacuum (e.g., 20 to 30 mmHg)). In embodiments wherein dip coating is used to deposit the enzyme domain at room temperature, a preferred insertion rate of from about 0.25 inch per minute to about 3 inches per minute, with a preferred dwell time of from about 0.5 minutes to about 2 minutes, and a preferred withdrawal rate of from about 0.25 inch per minute to about 2 inches per minute provides a functional coating. However, values outside of those set forth above can be acceptable or even desirable in certain embodiments, for example, depending upon viscosity and surface tension, as is appreciated by one skilled in the art. In one embodiment, the enzyme domain is formed by dip coating two times (namely, forming two layers) in an enzyme domain solution and curing at 50° C. under vacuum for 20 minutes. However, in some embodiments, the enzyme domain can be formed by dip coating and/or spray coating one or more layers at a predetermined concentration of the coating solution, insertion rate, dwell time, withdrawal rate, and/or desired thickness.

Resistance Domain

In preferred embodiments, the membrane system includes a resistance domain **50** disposed more distal from the electroactive surfaces than the enzyme domain. Although the following description is directed to a resistance domain for a glucose sensor, the resistance domain can be modified for other analytes and co-reactants as well.

There exists a molar excess of glucose relative to the amount of oxygen in blood; that is, for every free oxygen molecule in extracellular fluid, there are typically more than 100 glucose molecules present (see Updike et al., Diabetes Care 5:207-21 (1982)). However, an immobilized enzyme-based glucose sensor employing oxygen as co-reactant is preferably supplied with oxygen in non-rate-limiting excess in order for the sensor to respond linearly to changes in glucose concentration, while not responding to changes in oxygen concentration. Specifically, when a glucose-monitoring reaction is oxygen limited, linearity is not achieved above minimal concentrations of glucose. Without a semipermeable membrane situated over the enzyme domain to control the flux of glucose and oxygen, a linear response to glucose levels can be obtained only for glucose concentrations of up to about 40 mg/dL. However, in a clinical setting, a linear response to glucose levels is desirable up to at least about 400 mg/dL.

The resistance domain includes a semipermeable membrane that controls the flux of oxygen and glucose to the underlying enzyme domain, preferably rendering oxygen in a non-rate-limiting excess. As a result, the upper limit of linearity of glucose measurement is extended to a much higher value than that which is achieved without the resistance domain. In one embodiment, the resistance domain exhibits an oxygen to glucose permeability ratio of from about 50:1 or less to about 400:1 or more, preferably about 200:1. As a result, one-dimensional reactant diffusion is adequate to provide excess oxygen at all reasonable glucose and oxygen concentrations found in the subcutaneous matrix (see Rhodes et al., Anal. Chem., 66:1520-1529 (1994)).

In alternative embodiments, a lower ratio of oxygen-to-glucose can be sufficient to provide excess oxygen by using a high oxygen solubility domain (for example, a silicone or fluorocarbon-based material or domain) to enhance the

supply/transport of oxygen to the enzyme domain. If more oxygen is supplied to the enzyme, then more glucose can also be supplied to the enzyme without creating an oxygen rate-limiting excess. In alternative embodiments, the resistance domain is formed from a silicone composition, such as is described in U.S. Publication No. US-2005-0090607-A1.

In a preferred embodiment, the resistance domain includes a polyurethane membrane with both hydrophilic and hydrophobic regions to control the diffusion of glucose and oxygen to an analyte sensor, the membrane being fabricated easily and reproducibly from commercially available materials. A suitable hydrophobic polymer component is a polyurethane, or polyetherurethaneurea. Polyurethane is a polymer produced by the condensation reaction of a diisocyanate and a difunctional hydroxyl-containing material. A polyurethaneurea is a polymer produced by the condensation reaction of a diisocyanate and a difunctional amine-containing material. Preferred diisocyanates include aliphatic diisocyanates containing from about 4 to about 8 methylene units. Diisocyanates containing cycloaliphatic moieties can also be useful in the preparation of the polymer and copolymer components of the membranes of preferred embodiments. The material that forms the basis of the hydrophobic matrix of the resistance domain can be any of those known in the art as appropriate for use as membranes in sensor devices and as having sufficient permeability to allow relevant compounds to pass through it, for example, to allow an oxygen molecule to pass through the membrane from the sample under examination in order to reach the active enzyme or electrochemical electrodes. Examples of materials which can be used to make non-polyurethane type membranes include vinyl polymers, polyethers, polyesters, polyamides, inorganic polymers such as polysiloxanes and polycarbosiloxanes, natural polymers such as cellulosic and protein based materials, and mixtures or combinations thereof.

In a preferred embodiment, the hydrophilic polymer component is polyethylene oxide. For example, one useful hydrophobic-hydrophilic copolymer component is a polyurethane polymer that includes about 20% hydrophilic polyethylene oxide. The polyethylene oxide portions of the copolymer are thermodynamically driven to separate from the hydrophobic portions of the copolymer and the hydrophobic polymer component. The 20% polyethylene oxide-based soft segment portion of the copolymer used to form the final blend affects the water pick-up and subsequent glucose permeability of the membrane.

In preferred embodiments, the resistance domain is deposited onto the enzyme domain to yield a domain thickness of from about 0.05 microns or less to about 20 microns or more, more preferably from about 0.05, 0.1, 0.15, 0.2, 0.25, 0.3, 0.35, 0.4, 0.45, 0.5, 1, 1.5, 2, 2.5, 3, or 3.5 microns to about 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 19.5 microns, and more preferably still from about 2, 2.5 or 3 microns to about 3.5, 4, 4.5, or 5 microns. Preferably, the resistance domain is deposited onto the enzyme domain by vapor deposition, spray coating, or dip coating. In one preferred embodiment, spray coating is the preferred deposition technique. The spraying process atomizes and mists the solution, and therefore most or all of the solvent is evaporated prior to the coating material settling on the underlying domain, thereby minimizing contact of the solvent with the enzyme.

In another preferred embodiment, physical vapor deposition (e.g., ultrasonic vapor deposition) is used for coating one or more of the membrane domain(s) onto the electrodes, wherein the vapor deposition apparatus and process include an ultrasonic nozzle that produces a mist of micro-droplets in a vacuum chamber. In these embodiments, the micro-droplets move turbulently within the vacuum chamber, isotropically impacting and adhering to the surface of the substrate. Advantageously, vapor deposition as described above can be implemented to provide high production throughput of membrane deposition processes (e.g., at least about 20 to about 200 or more electrodes per chamber), greater consistency of the membrane on each sensor, and increased uniformity of sensor performance, for example, as described below.

In some embodiments, depositing the resistance domain (for example, as described in the preferred embodiments above) includes formation of a membrane system that substantially blocks or resists ascorbate (a known electrochemical interferant in hydrogen peroxide-measuring glucose sensors). While not wishing to be bound by theory, it is believed that during the process of depositing the resistance domain as described in the preferred embodiments, a structural morphology is formed that is characterized in that ascorbate does not substantially permeate therethrough.

In a preferred embodiment, the resistance domain is deposited on the enzyme domain by spray coating a solution of from about 1 wt. % to about 5 wt. % polymer and from about 95 wt. % to about 99 wt. % solvent. In spraying a solution of resistance domain material, including a solvent, onto the enzyme domain, it is desirable to mitigate or substantially reduce any contact with enzyme of any solvent in the spray solution that can deactivate the underlying enzyme of the enzyme domain. Tetrahydrofuran (THF) is one solvent that minimally or negligibly affects the enzyme of the enzyme domain upon spraying. Other solvents can also be suitable for use, as is appreciated by one skilled in the art.

Although a variety of spraying or deposition techniques can be used, spraying the resistance domain material and rotating the sensor at least one time by 180° can typically provide adequate coverage by the resistance domain. Spraying the resistance domain material and rotating the sensor at least two times by 120° provides even greater coverage (one layer of 360° coverage), thereby ensuring resistivity to glucose, such as is described in more detail above.

In preferred embodiments, the resistance domain is spray coated and subsequently cured for a time of from about 15 minutes to about 90 minutes at a temperature of from about 40° C. to about 60° C. (and can be accomplished under vacuum (e.g., from 20 to 30 mmHg)). A cure time of up to about 90 minutes or more can be advantageous to ensure complete drying of the resistance domain.

In one embodiment, the resistance domain is formed by spray coating at least six layers (namely, rotating the sensor seventeen times by 120° for at least six layers of 360° coverage) and curing at 50° C. under vacuum for 60 minutes. However, the resistance domain can be formed by dip coating or spray coating any layer or plurality of layers, depending upon the concentration of the solution, insertion rate, dwell time, withdrawal rate, and/or the desired thickness of the resulting film. Additionally, curing in a convention oven can also be employed.

In certain embodiments, a variable frequency microwave oven can be used to cure the membrane domains/layers. In general, microwave ovens directly excite the rotational mode of solvents. Consequently, microwave ovens cure coatings from the inside out rather than from the outside in as with conventional convection ovens. This direct rotational mode excitation is responsible for the typically observed "fast" curing within a microwave oven. In contrast

to conventional microwave ovens, which rely upon a fixed frequency of emission that can cause arcing of dielectric (metallic) substrates if placed within a conventional microwave oven, Variable Frequency Microwave (VFM) ovens emit thousands of frequencies within 100 milliseconds, which substantially eliminates arcing of dielectric substrates. Consequently, the membrane domains/layers can be cured even after deposition on metallic electrodes as described herein. While not wishing to be bound by theory, it is believe that VFM curing can increase the rate and completeness of solvent evaporation from a liquid membrane solution applied to a sensor, as compared to the rate and completeness of solvent evaporation observed for curing in conventional convection ovens.

In certain embodiments, VFM is can be used together with convection oven curing to further accelerate cure time. In some sensor applications wherein the membrane is cured prior to application on the electrode (see, for example, U.S. Publication No. US-2005-0245799-A1), conventional microwave ovens (e.g., fixed frequency microwave ovens) can be used to cure the membrane layer.

Treatment of Interference Domain/Membrane System

Although the above-described methods generally include a curing step in formation of the membrane system, including the interference domain, the preferred embodiments further include an additional treatment step, which can be performed directly after the formation of the interference domain and/or some time after the formation of the entire membrane system (or anytime in between). In some embodiments, the additional treatment step is performed during (or in combination with) sterilization of the sensor.

In some embodiments, the membrane system (or interference domain) is treated by exposure to ionizing radiation, for example, electron beam radiation, UV radiation, X-ray radiation, gamma radiation, and the like. Alternatively, the membrane can be exposed to visible light when suitable photoinitiators are incorporated into the interference domain. While not wishing to be bound by theory, it is believed that exposing the interference domain to ionizing radiation substantially crosslinks the interference domain and thereby creates a tighter, less permeable network than an interference domain that has not been exposed to ionizing radiation.

In some embodiments, the membrane system (or interference domain) is crosslinked by forming free radicals, which may include the use of ionizing radiation, thermal initiators, chemical initiators, photoinitiators (e.g., UV and visible light), and the like. Any suitable initiator or any suitable initiator system can be employed, for example, α-hydroxyketone, α-aminoketone, ammonium persulfate (APS), redox systems such as APS/bisulfite, or potassium permanganate. Suitable thermal initiators include but are not limited to potassium persulfate, ammonium persulfate, sodium persulfate, and mixtures thereof.

In embodiments wherein electron beam radiation is used to treat the membrane system (or interference domain), a preferred exposure time is from about 6 k or 12 kGy to about 25 or 50 kGy, more preferably about 25 kGy. However, one skilled in the art appreciates that choice of molecular weight, composition of cellulosic derivative (or other polymer), and/or the thickness of the layer can affect the preferred exposure time of membrane to radiation. Preferably, the exposure is sufficient for substantially crosslinking the interference domain to form free radicals, but does not destroy or significantly break down the membrane or does not significantly damage the underlying electroactive surfaces.

In embodiments wherein UV radiation is employed to treat the membrane, UV rays from about 200 nm to about 400 nm are preferred; however values outside of this range can be employed in certain embodiments, dependent upon the cellulosic derivative and/or other polymer used.

In some embodiments, for example, wherein photoinitiators are employed to crosslink the interference domain, one or more additional domains can be provided adjacent to the interference domain for preventing delamination that may be caused by the crosslinking treatment. These additional domains can be "tie layers" (i.e., film layers that enhance adhesion of the interference domain to other domains of the membrane system). In one exemplary embodiment, a membrane system is formed that includes the following domains: resistance domain, enzyme domain, electrode domain, and cellulosic-based interference domain, wherein the electrode domain is configured to ensure adhesion between the enzyme domain and the interference domain. In embodiments wherein photoinitiators are employed to crosslink the interference domain, UV radiation of greater than about 290 nm is preferred. Additionally, from about 0.01 to about 1 wt % photoinitiator is preferred weight-to-weight with a preselected cellulosic polymer (e.g., cellulose acetate); however values outside of this range can be desirable dependent upon the cellulosic polymer selected.

In general, sterilization of the transcutaneous sensor can be completed after final assembly, utilizing methods such as electron beam radiation, gamma radiation, glutaraldehyde treatment, or the like. The sensor can be sterilized prior to or after packaging. In an alternative embodiment, one or more sensors can be sterilized using variable frequency microwave chamber(s), which can increase the speed and reduce the cost of the sterilization process. In another alternative embodiment, one or more sensors can be sterilized using ethylene oxide (EtO) gas sterilization, for example, by treating with 100% ethylene oxide, which can be used when the sensor electronics are not detachably connected to the sensor and/or when the sensor electronics must undergo a sterilization process. In one embodiment, one or more packaged sets of transcutaneous sensors (e.g., 1, 2, 3, 4, or 5 sensors or more) are sterilized simultaneously.

Mutarotase Enzyme

In some embodiments, mutarotase, an enzyme that converts α D-glucose to β D-glucose, is incorporated into the membrane system. Mutarotase can be incorporated into the enzyme domain and/or can be incorporated into another domain of the membrane system. In general, glucose exists in two distinct isomers, α and β, which are in equilibrium with one another in solution and in the blood or interstitial fluid. At equilibrium, α is present at a relative concentration of about 35.5% and β is present in the relative concentration of about 64.5% (see Okuda et. al., Anal Biochem. 1971 September; 43(1):312-5). Glucose oxidase, which is a conventional enzyme used to react with glucose in glucose sensors, reacts with β D-glucose and not with α D-glucose. Since only the β D-glucose isomer reacts with the glucose oxidase, errant readings may occur in a glucose sensor responsive to a shift of the equilibrium between the α D-glucose and the β D-glucose. Many compounds, such as calcium, can affect equilibrium shifts of α D-glucose and β D-glucose. For example, as disclosed in U.S. Pat. No. 3,964,974 to Banaugh et al., compounds that exert a mutarotation accelerating effect on α D-glucose include histidine, aspartic acid, imidazole, glutamic acid, a hydroxyl pyridine, and phosphate.

Accordingly, a shift in α D-glucose and β D-glucose equilibrium can cause a glucose sensor based on glucose

oxidase to err high or low. To overcome the risks associated with errantly high or low sensor readings due to equilibrium shifts, the sensor of the preferred embodiments can be configured to measure total glucose in the host, including α D-glucose and β D-glucose by the incorporation of the mutarotase enzyme, which converts α D-glucose to β D-glucose.

Although sensors of some embodiments described herein include an interference domain in order to block or reduce one or more interferents, sensors with the membrane systems of the preferred embodiments, including an electrode domain **47**, an enzyme domain **48**, and a resistance domain **49**, have been shown to inhibit ascorbate without an additional interference domain. Namely, the membrane system of the preferred embodiments, including an electrode domain **47**, an enzyme domain **48**, and a resistance domain **49**, has been shown to be substantially non-responsive to ascorbate in physiologically acceptable ranges. While not wishing to be bound by theory, it is believed that the processing process of spraying the depositing the resistance domain by spray coating, as described herein, forms results in a structural morphology that is substantially resistance resistant to ascorbate.

Oxygen Conduit

As described above, certain sensors depend upon an enzyme within the membrane system through which the host's bodily fluid passes and in which the analyte (for example, glucose) within the bodily fluid reacts in the presence of a co-reactant (for example, oxygen) to generate a product. The product is then measured using electrochemical methods, and thus the output of an electrode system functions as a measure of the analyte. For example, when the sensor is a glucose oxidase based glucose sensor, the species measured at the working electrode is H_2O_2. An enzyme, glucose oxidase, catalyzes the conversion of oxygen and glucose to hydrogen peroxide and gluconate according to the following reaction:

$$\text{Glucose}+O_2 \rightarrow \text{Gluconate}+H_2O_2$$

Because for each glucose molecule reacted there is a proportional change in the product, H_2O_2, one can monitor the change in H_2O_2 to determine glucose concentration. Oxidation of H_2O_2 by the working electrode is balanced by reduction of ambient oxygen, enzyme generated H_2O_2 and other reducible species at a counter electrode, for example. See Fraser, D. M., "An Introduction to In vivo Biosensing: Progress and Problems." In "Biosensors and the Body," D. M. Fraser, ed., 1997, pp. 1-56 John Wiley and Sons, New York)).

In vivo, glucose concentration is generally about one hundred times or more that of the oxygen concentration. Consequently, oxygen is a limiting reactant in the electrochemical reaction, and when insufficient oxygen is provided to the sensor, the sensor is unable to accurately measure glucose concentration. Thus, depressed sensor function or inaccuracy is believed to be a result of problems in availability of oxygen to the enzyme and/or electroactive surface(s).

Accordingly, in an alternative embodiment, an oxygen conduit (for example, a high oxygen solubility domain formed from silicone or fluorochemicals) is provided that extends from the ex vivo portion of the sensor to the in vivo portion of the sensor to increase oxygen availability to the enzyme. The oxygen conduit can be formed as a part of the coating (insulating) material or can be a separate conduit associated with the assembly of wires that forms the sensor.

Porous Biointerface Materials

In alternative embodiments, the distal portion **42** includes a porous material disposed over some portion thereof, which modifies the host's tissue response to the sensor. In some embodiments, the porous material surrounding the sensor advantageously enhances and extends sensor performance and lifetime in the short term by slowing or reducing cellular migration to the sensor and associated degradation that would otherwise be caused by cellular invasion if the sensor were directly exposed to the in vivo environment. Alternatively, the porous material can provide stabilization of the sensor via tissue ingrowth into the porous material in the long term. Suitable porous materials include silicone, polytetrafluoroethylene, expanded polytetrafluoroethylene, polyethylene-co-tetrafluoroethylene, polyolefin, polyester, polycarbonate, biostable polytetrafluoroethylene, homopolymers, copolymers, terpolymers of polyurethanes, polypropylene (PP), polyvinylchloride (PVC), polyvinylidene fluoride (PVDF), polyvinyl alcohol (PVA), polybutylene terephthalate (PBT), polymethylmethacrylate (PMMA), polyether ether ketone (PEEK), polyamides, polyurethanes, cellulosic polymers, polysulfones and block copolymers thereof including, for example, di-block, tri-block, alternating, random and graft copolymers, as well as metals, ceramics, cellulose, hydrogel polymers, poly (2-hydroxyethyl methacrylate, pHEMA), hydroxyethyl methacrylate, (HEMA), polyacrylonitrile-polyvinyl chloride (PAN-PVC), high density polyethylene, acrylic copolymers, nylon, polyvinyl difluoride, polyanhydrides, poly(l-lysine), poly (L-lactic acid), hydroxyethylmethacrylate, hydroxyapeptite, alumina, zirconia, carbon fiber, aluminum, calcium phosphate, titanium, titanium alloy, nintinol, stainless steel, and CoCr alloy, or the like, such as are described in U.S. Publication No. US-2005-0031689-A1 and U.S. Publication No. US-2005-0112169-A1.

In some embodiments, the porous material surrounding the sensor provides unique advantages in the short term (e.g., one to 30 days) that can be used to enhance and extend sensor performance and lifetime. However, such materials can also provide advantages in the long term too (e.g., greater than 30 days) and thus are useful in the long term sensors described below. Particularly, the in vivo portion of the sensor (the portion of the sensor that is implanted into the host's tissue) is encased (partially or fully) in a porous material. The porous material can be wrapped around the sensor (for example, by wrapping the porous material around the sensor or by inserting the sensor into a section of porous material sized to receive the sensor). Alternately, the porous material can be deposited on the sensor (for example, by electrospinning of a polymer directly thereon). In yet other alternative embodiments, the sensor is inserted into a selected section of porous biomaterial. Other methods for surrounding the in vivo portion of the sensor with a porous material can also be used as is appreciated by one skilled in the art.

The porous material surrounding the sensor advantageously slows or reduces cellular migration to the sensor and associated degradation that would otherwise be caused by cellular invasion if the sensor were directly exposed to the in vivo environment. Namely, the porous material provides a barrier that makes the migration of cells towards the sensor more tortuous and therefore slower (providing short term advantages). It is believed that this reduces or slows the sensitivity loss normally observed in a short-term sensor over time.

In an embodiment wherein the porous material is a high oxygen solubility material, such as porous silicone, the high oxygen solubility porous material surrounds some of or the

entire in vivo portion **42** of the sensor. High oxygen solubility materials are materials that dynamically retain a high availability of oxygen that can be used to compensate for the local oxygen deficit during times of transient ischemia (e.g., silicone and fluorocarbons). It is believed that some signal noise normally seen by a conventional sensor can be attributed to an oxygen deficit. In one exemplary embodiment, porous silicone surrounds the sensor and thereby effectively increases the concentration of oxygen local (proximal) to the sensor. Thus, an increase in oxygen availability proximal to the sensor as achieved by this embodiment ensures that an excess of oxygen over glucose is provided to the sensor; thereby reducing the likelihood of oxygen limited reactions therein. Accordingly, by providing a high oxygen solubility material (e.g., porous silicone) surrounding the in vivo portion of the sensor, it is believed that increased oxygen availability, reduced signal noise, longevity, and ultimately enhanced sensor performance can be achieved.

Bioactive Agents

In some alternative embodiments, a bioactive agent is incorporated into the above described porous material and/or membrane system, which diffuses out into the environment adjacent to the sensing region, such as is described in U.S. Publication No. US-2005-0031689-A1. Additionally or alternately, a bioactive agent can be administered locally at the exit-site or implantation-site. Suitable bioactive agents are those that modify the host's tissue response to the sensor, for example anti-inflammatory agents, anti-infective agents, anesthetics, inflammatory agents, growth factors, immunosuppressive agents, antiplatelet agents, anti-coagulants, antiproliferates, ACE inhibitors, cytotoxic agents, anti-barrier cell compounds, vascularization-inducing compounds, antisense molecules, or mixtures thereof, such as are described in more detail in co-pending U.S. Patent Publication No. US-2005-0031689-A1.

In embodiments wherein the porous material is designed to enhance short-term (e.g., from about 1 to about 30 days) lifetime or performance of the sensor, a suitable bioactive agent can be chosen to ensure that tissue ingrowth does not substantially occur within the pores of the porous material. Namely, by providing a tissue modifying bioactive agent, such as an anti-inflammatory agent (for example, Dexamethasone), substantially tissue ingrowth can be inhibited, at least in the short term, in order to maintain sufficient glucose transport through the pores of the porous material to maintain a stable sensitivity.

In embodiments wherein the porous material is designed to enhance long-term (e.g., from about a day to about a year or more) lifetime or performance of the sensor, a suitable bioactive agent, such as a vascularization-inducing compound or anti-barrier cell compound, can be chosen to encourage tissue ingrowth without barrier cell formation.

In some alternative embodiments, the in vivo portion of the sensor is designed with porosity therethrough, for example, a design wherein the sensor wires are configured in a mesh, loose helix configuration (namely, with spaces between the wires), or with micro-fabricated holes therethrough. Porosity within the sensor modifies the host's tissue response to the sensor, because tissue ingrowth into and/or through the in vivo portion of the sensor increases stability of the sensor and/or improves host acceptance of the sensor, thereby extending the lifetime of the sensor in vivo.

Sensor Manufacture

In some embodiments, the sensor is manufactured partially or wholly using a continuous reel-to-reel process, wherein one or more manufacturing steps are automated. In such embodiments, a manufacturing process can be provided substantially without the need for manual mounting and fixing steps and substantially without the need human interaction. A process can be utilized wherein a plurality of sensors of the preferred embodiments, including the electrodes, insulator, and membrane system, are continuously manufactured in a semi-automated or automated process.

In one embodiment, a plurality of twisted pairs is continuously formed into a coil, wherein a working electrode is coated with an insulator material around which a plurality of reference electrodes is wound. The plurality of twisted pairs are preferably indexed and subsequently moved from one station to the next whereby the membrane system is serially deposited according to the preferred embodiments. Preferably, the coil is continuous and remains as such during the entire sensor fabrication process, including winding of the electrodes, insulator application, and membrane coating processes. After drying of the membrane system, each individual sensor is cut from the continuous coil.

A continuous reel-to-reel process for manufacturing the sensor eliminates possible sensor damage due to handling by eliminating handling steps, and provides faster manufacturing due to faster trouble shooting by isolation when a product fails. Additionally, a process run can be facilitated because of elimination of steps that would otherwise be required (e.g., steps in a manual manufacturing process). Finally, increased or improved product consistency due to consistent processes within a controlled environment can be achieved in a machine or robot driven operation.

In certain embodiments, vapor deposition (e.g., physical vapor deposition) is utilized to deposit one or more of the membrane domains onto the sensor. Vapor deposition can be used to coat one or more insulating layers onto the electrodes and one or more of the domains of the membrane system onto the electrochemically reactive surfaces. The vapor deposition process can be a part of a continuous manufacturing process, for example, a semi-automated or fully-automated manufacturing process. Physical vapor deposition processes are generally preferred. In such physical vapor deposition processes in the gas phase for forming a thin film, source material is physically transferred in a vacuum to the substrate without any chemical reaction(s) involved. Physical vapor deposition processes include evaporation (e.g., by thermal or e-beam) and sputtering processes. In alternative embodiments, chemical vapor deposition can be used. In chemical vapor deposition processes for depositing a thin film, the substrate is exposed to one or more volatile precursors, which react and/or decompose on the substrate surface to produce the desired deposit. Advantageously, vapor deposition processes can be implemented to provide high production throughput of membrane deposition processes (e.g., deposition on at least about 20 to about 200 or more electrodes per chamber), greater consistency of the membrane on each sensor, and increased uniformity of sensor performance.

Applicator

FIG. **7** is an exploded side view of an applicator, showing the components that enable sensor and needle insertion. In this embodiment, the applicator **12** includes an applicator body **18** that aides in aligning and guiding the applicator components. Preferably, the applicator body **18** includes an applicator body base **60** that matingly engages the mounting unit **14** and an applicator body cap **62** that enables appropriate relationships (for example, stops) between the applicator components.

The guide tube subassembly **20** includes a guide tube carrier **64** and a guide tube **66**. In some embodiments, the guide tube is a cannula. The guide tube carrier **64** slides

along the applicator body **18** and maintains the appropriate relative position of the guide tube **66** during insertion and subsequent retraction. For example, prior to and during insertion of the sensor, the guide tube **66** extends through the contact subassembly **26** to maintain an opening that enables easy insertion of the needle therethrough (see FIGS. **8**A to **8**D). During retraction of the sensor, the guide tube subassembly **20** is pulled back, engaging with and causing the needle and associated moving components to retract back into the applicator **12** (See FIGS. **8**C and **8**D). In some embodiments, a lubricant (e.g., petroleum jelly) is placed within the sealing member **36** of the contact subassembly such that it surrounds the guide tube (e.g., cannula), thereby allowing the guide tube to easily retract back into the applicator, for example, without causing compression or deformation of the sealing member **36**.

A needle subassembly **68** is provided that includes a needle carrier **70** and needle **72**. The needle carrier **70** cooperates with the other applicator components and carries the needle **72** between its extended and retracted positions. The needle can be of any appropriate size that can encompass the sensor **32** and aid in its insertion into the host. Preferred sizes include from about 32 gauge or less to about 18 gauge or more, more preferably from about 28 gauge to about 25 gauge, to provide a comfortable insertion for the host. Referring to the inner diameter of the needle, approximately 0.006 inches to approximately 0.023 inches is preferable, and 0.013 inches is most preferable. The needle carrier **70** is configured to engage with the guide tube carrier **64**, while the needle **72** is configured to slidably nest within the guide tube **66**, which allows for easy guided insertion (and retraction) of the needle through the contact subassembly **26**.

A push rod subassembly **74** is provided that includes a push rod carrier **76** and a push rod **78**. The push rod carrier **76** cooperates with other applicator components to ensure that the sensor is properly inserted into the host's skin, namely the push rod carrier **76** carries the push rod **78** between its extended and retracted positions. In this embodiment, the push rod **78** is configured to slidably nest within the needle **72**, which allows for the sensor **32** to be pushed (released) from the needle **72** upon retraction of the needle, which is described in more detail with reference to FIGS. **8**A through **8**D. In some embodiments, a slight bend or serpentine shape is designed into or allowed in the sensor in order to maintain the sensor within the needle by interference. While not wishing to be bound by theory, it is believed that a slight friction fit of the sensor within the needle minimizes motion of the sensor during withdrawal of the needle and maintains the sensor within the needle prior to withdrawal of the needle.

A plunger subassembly **22** is provided that includes a plunger **80** and plunger cap **82**. The plunger subassembly **22** cooperates with other applicators components to ensure proper insertion and subsequent retraction of the applicator components. In this embodiment, the plunger **80** is configured to engage with the push rod to ensure the sensor remains extended (namely, in the host) during retraction, such as is described in more detail with reference to FIG. **8**C.

Sensor Insertion

FIGS. **8**A through **8**D are schematic side cross-sectional views that illustrate the applicator components and their cooperating relationships at various stages of sensor insertion. FIG. **8**A illustrates the needle and sensor loaded prior to sensor insertion. FIG. **8**B illustrates the needle and sensor after sensor insertion. FIG. **8**C illustrates the sensor and needle during needle retraction. FIG. **8**D illustrates the sensor remaining within the contact subassembly after needle retraction. Although the embodiments described herein suggest manual insertion and/or retraction of the various components, automation of one or more of the stages can also be employed. For example, spring-loaded mechanisms that can be triggered to automatically insert and/or retract the sensor, needle, or other cooperative applicator components can be implemented.

Referring to FIG. **8**A, the sensor **32** is shown disposed within the needle **72**, which is disposed within the guide tube **66**. In this embodiment, the guide tube **66** is provided to maintain an opening within the contact subassembly **26** and/or contacts **28** to provide minimal friction between the needle **72** and the contact subassembly **26** and/or contacts **28** during insertion and retraction of the needle **72**. However, the guide tube is an optional component, which can be advantageous in some embodiments wherein the contact subassembly **26** and/or the contacts **28** are formed from an elastomer or other material with a relatively high friction coefficient, and which can be omitted in other embodiments wherein the contact subassembly **26** and or the contacts **28** are formed from a material with a relatively low friction coefficient (for example, hard plastic or metal). A guide tube, or the like, can be preferred in embodiments wherein the contact subassembly **26** and/or the contacts **28** are formed from a material designed to frictionally hold the sensor **32** (see FIG. **8**D), for example, by the relaxing characteristics of an elastomer, or the like. In these embodiments, the guide tube is provided to ease insertion of the needle through the contacts, while allowing for a frictional hold of the contacts on the sensor **32** upon subsequent needle retraction. Stabilization of the sensor in or on the contacts **28** is described in more detail with reference to FIG. **8**D and following. Although FIG. **8**A illustrates the needle and sensor inserted into the contacts subassembly as the initial loaded configuration, alternative embodiments contemplate a step of loading the needle through the guide tube **66** and/or contacts **28** prior to sensor insertion.

Referring to FIG. **8**B, the sensor **32** and needle **72** are shown in an extended position. In this stage, the pushrod **78** has been forced to a forward position, for example by pushing on the plunger shown in FIG. **7**, or the like. The plunger **22** (FIG. **7**) is designed to cooperate with other of the applicator components to ensure that sensor **32** and the needle **72** extend together to a forward position (as shown); namely, the push rod **78** is designed to cooperate with other of the applicator components to ensure that the sensor **32** maintains the forward position simultaneously within the needle **72**.

Referring to FIG. **8**C, the needle **72** is shown during the retraction process. In this stage, the push rod **78** is held in its extended (forward) position in order to maintain the sensor **32** in its extended (forward) position until the needle **72** has substantially fully retracted from the contacts **28**. Simultaneously, the cooperating applicator components retract the needle **72** and guide tube **66** backward by a pulling motion (manual or automated) thereon. In preferred embodiments, the guide tube carrier **64** (FIG. **7**) engages with cooperating applicator components such that a backward (retraction) motion applied to the guide tube carrier retracts the needle **72** and guide tube **66**, without (initially) retracting the push rod **78**. In an alternative embodiment, the push rod **78** can be omitted and the sensor **32** held it its forward position by a cam, elastomer, or the like, which is in contact with a portion of the sensor while the needle moves over another portion of the sensor. One or more slots can be cut in the needle to maintain contact with the sensor during needle retraction.

Referring to FIG. **8**D, the needle **72**, guide tube **66**, and push rod **78** are all retracted from contact subassembly **26**, leaving the sensor **32** disposed therein. The cooperating applicator components are designed such that when the needle **72** has substantially cleared from the contacts **28** and/or contact subassembly **26**, the push rod **78** is retracted along with the needle **72** and guide tube **66**. The applicator **12** can then be released (manually or automatically) from the contacts **28**, such as is described in more detail elsewhere herein, for example with reference to FIGS. **9**D and **10**A.

The preferred embodiments are generally designed with elastomeric contacts to ensure a retention force that retains the sensor **32** within the mounting unit **14** and to ensure stable electrical connection of the sensor **32** and its associated contacts **28**. Although the illustrated embodiments and associated text describe the sensor **32** extending through the contacts **28** to form a friction fit therein, a variety of alternatives are contemplated. In one alternative embodiment, the sensor is configured to be disposed adjacent to the contacts (rather than between the contacts). The contacts can be constructed in a variety of known configurations, for example, metallic contacts, cantilevered fingers, pogo pins, or the like, which are configured to press against the sensor after needle retraction.

It is generally preferred that a contact **28** is formed from a material with a durometer hardness of from about 5 to about 80 Shore A, more preferably from about 10 to about 50 Shore A, and even more preferably from about 20 to about 50 Shore A. In one implementation of a transcutaneous analyte sensor as described with reference to the preferred embodiments, the contact **28** is formed from a material with a durometer hardness of about 20 Shore A to maximize conformance (e.g., compression) of the contact around the sensor and/or within the sealing member. In another implementation of a transcutaneous analyte sensor as described with reference to the preferred embodiments, the contact **28** is formed from a material with a durometer hardness of about 50 Shore A to increase the strength of the contact **28** (e.g., increase resistance to compression). While a few examples have been provided above, one skilled in the art will appreciate that higher or lower durometer hardness sealing materials can also be advantageously employed.

In some embodiments, the durometer hardness of the elastomeric contacts **28** is higher than the durometer hardness of the sealing member **36**. In one example, the durometer hardness of the contacts is about 50 Shore A and the durometer hardness of the sealing member is about 20 Shore A; however, a variety of durometer hardness materials within the preferred range (typically, from about 5 Shore A to about 80 Shore A) can be chosen. In these embodiments, the higher durometer hardness contacts generally provide greater stability while the lower durometer hardness sealing member generally provides superior compression and/or seal around the contacts.

In some embodiments, the durometer hardness of the sealing member **36** is higher than the durometer hardness of the elastomeric contacts **28**. In one example, the durometer hardness of the sealing member is about 50 Shore A and the durometer hardness of the contacts is about 20 Shore A, however a variety of durometer hardness materials within the preferred range (typically, from about 5 Shore A to about 80 Shore A) can be chosen. In these embodiments, the higher durometer hardness sealing member provides greater stability while the lower durometer hardness contacts provide superior compression and/or seal.

The illustrated embodiments are designed with coaxial contacts **28**; namely, the contacts **28** are configured to contact the working and reference electrodes **44**, **46** axially along the distal portion **42** of the sensor **32** (see FIG. **6**A). As shown in FIG. **6**A, the working electrode **44** extends farther than the reference electrode **46**, which allows coaxial connection of the electrodes **44**, **46** with the contacts **28** at locations spaced along the distal portion of the sensor (see also FIGS. **10**B and **11**B). Although the illustrated embodiments employ a coaxial design, other designs are contemplated within the scope of the preferred embodiments. For example, the reference electrode can be positioned substantially adjacent to (but spaced apart from) the working electrode at the distal portion of the sensor. In this way, the contacts **28** can be designed side-by-side rather than coaxially along the axis of the sensor.

FIG. **9**A is a perspective view of an applicator and mounting unit in one embodiment including a safety latch mechanism **84**. The safety latch mechanism **84** is configured to lock the plunger subassembly **22** in a stationary position such that it cannot be accidentally pushed prior to release of the safety latch mechanism. In this embodiment, the sensor system **10** is preferably packaged (e.g., shipped) in this locked configuration, wherein the safety latch mechanism **84** holds the plunger subassembly **22** in its extended position, such that the sensor **32** cannot be prematurely inserted (e.g., accidentally released). The safety latch mechanism **84** is configured such that a pulling force shown in the direction of the arrow (see FIG. **9**A) releases the lock of the safety latch mechanism on the plunger subassembly, thereby allowing sensor insertion. Although one safety latch mechanism that locks the plunger subassembly is illustrated and described herein, a variety of safety latch mechanism configurations that lock the sensor to prevent it from prematurely releasing (i.e., that lock the sensor prior to release of the safety latch mechanism) are contemplated, as can be appreciated by one skilled in the art, and fall within the scope of the preferred embodiments.

FIG. **9**A additionally illustrates a force-locking mechanism **86** included in certain alternative embodiments of the sensor system, wherein the force-locking mechanism **86** is configured to ensure a proper mate between the electronics unit **16** and the mounting unit **14** (see FIG. **13**A, for example). In embodiments wherein a seal is formed between the mounting unit and the electronics unit, as described in more detail elsewhere herein, an appropriate force may be required to ensure a seal has sufficiently formed therebetween; in some circumstances, it can be advantageous to ensure the electronics unit has been properly mated (e.g., snap-fit or sealingly mated) to the mounting unit. Accordingly, upon release of the applicator **12** from the mounting unit **14** (after sensor insertion), and after insertion of the electronics unit **16** into the mounting unit **14**, the force-locking mechanism **86** allows the user to ensure a proper mate and/or seal therebetween. In practice, a user pivots (e.g., lifts or twists) the force-locking mechanism such that it provides force on the electronics unit **16** by pulling up on the circular tab illustrated in FIG. **9**A; the force-locking mechanism is preferably released thereafter. Although one system and one method for providing a secure and/or sealing fit between the electronics unit and the mounting unit are illustrated, various other force-locking mechanisms can be employed that utilize a variety of systems and methods for providing a secure and/or sealing fit between the electronics unit and the mounting unit (housing).

FIGS. **9**B to **9**D are side views of an applicator and mounting unit in one embodiment, showing various stages of sensor insertion. FIG. **9**B is a side view of the applicator matingly engaged to the mounting unit prior to sensor

insertion. FIG. **9**C is a side view of the mounting unit and applicator after the plunger subassembly has been pushed, extending the needle and sensor from the mounting unit (namely, through the host's skin). FIG. **9**D is a side view of the mounting unit and applicator after the guide tube subassembly has been retracted, retracting the needle back into the applicator. Although the drawings and associated text illustrate and describe embodiments wherein the applicator is designed for manual insertion and/or retraction, automated insertion and/or retraction of the sensor/needle, for example, using spring-loaded components, can alternatively be employed.

The preferred embodiments advantageously provide a system and method for easy insertion of the sensor and subsequent retraction of the needle in a single push-pull motion. Because of the mechanical latching system of the applicator, the user provides a continuous force on the plunger cap **82** and guide tube carrier **64** that inserts and retracts the needle in a continuous motion. When a user grips the applicator, his or her fingers grasp the guide tube carrier **64** while his or her thumb (or another finger) is positioned on the plunger cap **82**. The user squeezes his or her fingers and thumb together continuously, which causes the needle to insert (as the plunger slides forward) and subsequently retract (as the guide tube carrier slides backward) due to the system of latches located within the applicator (FIGS. **7** to **9**) without any necessary change of grip or force, leaving the sensor implanted in the host. In some embodiments, a continuous torque, when the applicator components are configured to rotatingly engage one another, can replace the continuous force. Some prior art sensors, in contrast to the sensors of the preferred embodiments, suffer from complex, multi-step, or multi-component insertion and retraction steps to insert and remove the needle from the sensor system.

FIG. **9**B shows the mounting unit and applicator in the ready position. The sensor system can be shipped in this configuration, or the user can be instructed to mate the applicator **12** with the mounting unit **14** prior to sensor insertion. The insertion angle α is preferably fixed by the mating engagement of the applicator **12**. In the illustrated embodiment, the insertion angle α is fixed in the applicator **12** by the angle of the applicator body base **60** with the shaft of the applicator body **18**. However, a variety of systems and methods of ensuring proper placement can be implemented. Proper placement ensures that at least a portion of the sensor **32** extends below the dermis of the host upon insertion. In alternative embodiments, the sensor system **10** is designed with a variety of adjustable insertion angles. A variety of insertion angles can be advantageous to accommodate a variety of insertion locations and/or individual dermis configurations (for example, thickness of the dermis). In preferred embodiments, the insertion angle α is from about 0 to about 90 degrees, more preferably from about 30 to about 60 degrees, and even more preferably about 45 degrees.

In practice, the mounting unit is placed at an appropriate location on the host's skin, for example, the skin of the arm, thigh, or abdomen. Thus, removing the backing layer **9** from the adhesive pad **8** and pressing the base portion of the mounting unit on the skin adheres the mounting unit to the host's skin.

FIG. **9**C shows the mounting unit and applicator after the needle **72** has been extended from the mounting unit **14** (namely, inserted into the host) by pushing the push rod subassembly **22** into the applicator **12**. In this position, the sensor **32** is disposed within the needle **72** (namely, in position within the host), and held by the cooperating applicator components. In alternative embodiments, the mounting unit and/or applicator can be configured with the needle/sensor initially extended. In this way, the mechanical design can be simplified and the plunger-assisted insertion step can be eliminated or modified. The needle can be simply inserted by a manual force to puncture the host's skin, and only one (pulling) step is required on the applicator, which removes the needle from the host's skin.

FIG. **9**D shows the mounting unit and applicator after the needle **72** has been retracted into the applicator **12**, exposing the sensor **32** to the host's tissue. During needle retraction, the push rod subassembly maintains the sensor in its extended position (namely, within the host). In preferred embodiments, retraction of the needle irreversibly locks the needle within the applicator so that it cannot be accidentally and/or intentionally released, reinserted, or reused. The applicator is preferably configured as a disposable device to reduce or eliminate a possibility of exposure of the needle after insertion into the host. However a reusable or reloadable applicator is also contemplated in some alternative embodiments. After needle retraction, the applicator **12** can be released from the mounting unit, for example, by pressing the release latch(es) **30**, and the applicator disposed of appropriately. In alternative embodiments, other mating and release configurations can be implemented between the mounting unit and the applicator, or the applicator can automatically release from the mounting unit after sensor insertion and subsequent needle retraction. In one alternative embodiment, a retention hold (e.g., ball and detent configuration) holds and releases the electronics unit (or applicator).

In one alternative embodiment, the mounting unit is configured to releasably mate with the applicator and electronics unit in a manner such that when the applicator is releasably mated to the mounting unit (e.g., after sensor insertion), the electronics unit is configured to slide into the mounting unit, thereby triggering release of the applicator and simultaneous mating of the electronics unit to the mounting unit. Cooperating mechanical components, for example, sliding ball and detent type configurations, can be used to accomplish the simultaneous mating of electronics unit and release of the applicator.

FIGS. **9**E to **9**G are perspective views of a sensor system **310** of an alternative embodiment, including an applicator **312**, electronics unit **316**, and mounting unit **314**, showing various stages of applicator release and/or electronic unit mating. FIG. **9**E is a perspective view of the applicator matingly engaged to the mounting unit after sensor insertion. FIG. **9**F is a perspective view of the mounting unit and applicator matingly engaged while the electronics unit is slidingly inserted into the mounting unit. FIG. **9**G is a perspective view of the electronics unit matingly engaged with the mounting unit after the applicator has been released.

In general, the sensor system **310** comprises a sensor adapted for transcutaneous insertion into a host's skin; a housing **314** adapted for placement adjacent to the host's skin; an electronics unit **316** releasably attachable to the housing; and an applicator **312** configured to insert the sensor through the housing **314** and into the skin of the host, wherein the applicator **312** is adapted to releasably mate with the housing **314**, and wherein the system **310** is configured to release the applicator **312** from the housing when the electronics unit **316** is attached to the housing **314**.

FIG. **9**E shows the sensor system **310** after the sensor has been inserted and prior to release of the applicator **312**. In this embodiment, the electronics unit **316** is designed to slide into the mounting unit **314**. Preferably, the electronics unit **316** is configured and arranged to slide into the mounting unit **314** in only one orientation. In the illustrated embodi-

ment, the insertion end is slightly tapered and dovetailed in order to guide insertion of the electronics unit **316** into the housing **314**; however other self-alignment configurations are possible. In this way, the electronics unit **316** self-aligns and orients the electronics unit **316** in the housing, ensuring a proper fit and a secure electronic connection with the sensor.

FIG. **9**F shows the sensor system **310** after the electronics unit **316** has been inserted therein. Preferably, the electronic unit **316** slide-fits into the mounting unit. In some embodiments, the sensor system **310** can be designed to allow the electronics unit **316** to be attached to the mounting unit **314** (i.e., operably connected to the sensor) before the sensor system **310** is affixed to the host. Advantageously, this design provides mechanical stability for the sensor during transmitter insertion.

FIG. **9**G shows the sensor system **310** upon release of the applicator **312** from the mounting unit **314** and electronics unit **316**. In this embodiment, the sensor system **310** is configured such that mating the electronics unit to the mounting unit triggers the release of the applicator **312** from the mounting unit **314**.

Thus, the above described sensor system **310**, also referred to as the slide-in system, allows for self-alignment of the electronics unit, creates an improved seal around the contacts due to greater holding force, provides mechanical stability for the sensor during insertion of the electronics unit, and causes automatic release of the applicator and simultaneous lock of the electronics unit into the mounting unit.

Although the overall design of the sensor system **10** results in a miniaturized volume as compared to numerous conventional devices, as described in more detail below; the sensor system **310** further enables a reduction in volume, as compared to, for example, the sensor system **10** described above.

FIGS. **9**H and **9**I are comparative top views of the sensor system shown in the alternative embodiment illustrated in FIGS. **9**E to **9**G and compared to the embodiments illustrated elsewhere (see FIGS. **1** to **3** and **10** to **12**, for example). Namely, the alternative embodiment described with reference to FIGS. **9**E to **9**G further enables reduced size (e.g., mass, volume, and the like) of the device as compared to certain other devices. It has been discovered that the size (including volume and/or surface area) of the device can affect the function of the device. For example, motion of the mounting unit/electronics unit caused by external influences (e.g., bumping or other movement on the skin) is translated to the sensor in vivo, causing motion artifact (e.g., an effect on the signal, or the like). Accordingly, by enabling a reduction of size, a more stable signal with overall improved patient comfort can be achieved.

Accordingly, slide-in system **310** described herein, including the systems and methods for inserting the sensor and connecting the electronics unit to the mounting unit, enables the mounting unit **316**/electronics unit **314** subassembly to have a volume of less than about 10 cm^3, more preferably less than about 8 cm^3, and even more preferably less than about 6 cm^3, 5 cm^3, or 4 cm^3 or less. In general, the mounting unit **316**/electronics unit **314** subassembly comprises a first major surface and a second major surface opposite the first major surface. The first and second major surfaces together preferably account for at least about 50% of the surface area of the device; the first and second major surfaces each define a surface area, wherein the surface area of each major surface is less than or equal to about 10 cm^2, preferably less than or equal to about 8 cm^2, and more preferably less than or equal to about 6.5 cm^2, 6 cm^2, 5.5 cm^2, 5 cm^2, 4.5 cm^2, or 4 cm^2 or less. Typically, the mounting unit **316**/electronics unit **314** subassembly has a length **320** of less than about 40 mm by a width **322** of less than about 20 mm and a thickness of less than about 10 mm, and more preferably a length **320** less than or equal to about 35 mm by a width **322** less than or equal to about 18 mm by a thickness of less than or equal to about 9 mm.

In some embodiments, the mounting unit **14**/electronics unit **16** assembly has the following dimensional properties: preferably a length of about 6 cm or less, more preferably about 5 cm or less, more preferably still about 4.6 cm or less, even more preferably 4 cm or less, and most preferably about 3 cm or less; preferably a width of about 5 cm or less, more preferably about 4 cm or less, even more preferably 3 cm or less, even more preferably still about 2 cm or less, and most preferably about 1.5 cm or less; and/or preferably a thickness of about 2 cm or less, more preferably about 1.3 cm or less, more preferably still about 1 cm or less, even more preferably still about 0.7 cm or less, and most preferably about 0.5 cm or less. The mounting unit **14**/electronics unit **16** assembly preferably has a volume of about 20 cm^3 or less, more preferably about 10 cm^3 or less, more preferably still about 5 cm^3 or less, and most preferably about 3 cm^3 or less; and preferably weighs 12 g or less, more preferably about 9 g or less, and most preferably about 6 g or less, although in some embodiments the electronics unit may weigh more than about 12 g, e.g., up to about 25 g, 45 g, or 90 g.

In some embodiments, the sensor **32** exits the base of the mounting unit **14** at a location distant from an edge of the base. In some embodiments, the sensor **32** exits the base of the mounting unit **14** at a location substantially closer to the center than the edges thereof. While not wishing to be bound by theory, it is believed that by providing an exit port for the sensor **32** located away from the edges, the sensor **32** can be protected from motion between the body and the mounting unit, snagging of the sensor by an external source, and/or environmental contaminants (e.g., microorganisms) that can migrate under the edges of the mounting unit. In some embodiments, the sensor exits the mounting unit away from an outer edge of the device.

In some alternative embodiments, however, the sensor exits the mounting unit **14** at an edge or near an edge of the device. In some embodiments, the mounting unit is configured such that the exit port (location) of the sensor is adjustable; thus, in embodiments wherein the depth of the sensor insertion is adjustable, six-degrees of freedom can thereby be provided.

Extensible Adhesive Pad

In certain embodiments, an adhesive pad is used with the sensor system. A variety of design parameters are desirable when choosing an adhesive pad for the mounting unit. For example: 1) the adhesive pad can be strong enough to maintain full contact at all times and during all movements (devices that release even slightly from the skin have a greater risk of contamination and infection), 2) the adhesive pad can be waterproof or water permeable such that the host can wear the device even while heavily perspiring, showering, or even swimming in some cases, 3) the adhesive pad can be flexible enough to withstand linear and rotational forces due to host movements, 4) the adhesive pad can be comfortable for the host, 5) the adhesive pad can be easily releasable to minimize host pain, 6) and/or the adhesive pad can be easily releasable so as to protect the sensor during release. Unfortunately, these design parameters are difficult to simultaneously satisfy using known adhesive pads, for

example, strong medical adhesive pads are available but are usually non-precise (for example, requiring significant "ripping" force during release) and can be painful during release due to the strength of their adhesion.

Therefore, the preferred embodiments provide an adhesive pad **8'** for mounting the mounting unit onto the host, including a sufficiently strong medical adhesive pad that satisfies one or more strength and flexibility requirements described above, and further provides a for easy, precise and pain-free release from the host's skin. FIG. **10**A is a side view of the sensor assembly, illustrating the sensor implanted into the host with mounting unit adhered to the host's skin via an adhesive pad in one embodiment. Namely, the adhesive pad **8'** is formed from an extensible material that can be removed easily from the host's skin by stretching it lengthwise in a direction substantially parallel to (or up to about 35 degrees from) the plane of the skin. It is believed that this easy, precise, and painless removal is a function of both the high extensibility and easy stretchability of the adhesive pad.

In one embodiment, the extensible adhesive pad includes a polymeric foam layer or is formed from adhesive pad foam. It is believed that the conformability and resiliency of foam aids in conformation to the skin and flexibility during movement of the skin. In another embodiment, a stretchable solid adhesive pad, such as a rubber-based or an acrylate-based solid adhesive pad can be used. In another embodiment, the adhesive pad comprises a film, which can aid in increasing load bearing strength and rupture strength of the adhesive pad

FIGS. **10**B to **10**C illustrate initial and continued release of the mounting unit from the host's skin by stretching the extensible adhesive pad in one embodiment. To release the device, the backing adhesive pad is pulled in a direction substantially parallel to (or up to about 35 degrees from) the plane of the device. Simultaneously, the extensible adhesive pad stretches and releases from the skin in a relatively easy and painless manner.

In one implementation, the mounting unit is bonded to the host's skin via a single layer of extensible adhesive pad **8'**, which is illustrated in FIGS. **10**A to **10**C. The extensible adhesive pad includes a substantially non-extensible pull-tab **52**, which can include a light adhesive pad layer that allows it to be held on the mounting unit **14** prior to release. Additionally, the adhesive pad can further include a substantially non-extensible holding tab **54**, which remains attached to the mounting unit during release stretching to discourage complete and/or uncontrolled release of the mounting unit from the skin.

In one alternative implementation, the adhesive pad **8'** includes two-sides, including the extensible adhesive pad and a backing adhesive pad (not shown). In this embodiment, the backing adhesive pad is bonded to the mounting unit's back surface **25** while the extensible adhesive pad **8'** is bonded to the host's skin. Both adhesive pads provide sufficient strength, flexibility, and waterproof or water permeable characteristics appropriate for their respective surface adhesion. In some embodiments, the backing and extensible adhesive pads are particularly designed with an optimized bond for their respective bonding surfaces (namely, the mounting unit and the skin).

In another alternative implementation, the adhesive pad **8'** includes a double-sided extensible adhesive pad surrounding a middle layer or backing layer (not shown). The backing layer can comprise a conventional backing film or can be formed from foam to enhance comfort, conformability, and flexibility. Preferably, each side of the double-sided adhesive pad is respectively designed for appropriate bonding surface (namely, the mounting unit and skin). A variety of alternative stretch-release configurations are possible. Controlled release of one or both sides of the adhesive pad can be facilitated by the relative lengths of each adhesive pad side, by incorporation of a non-adhesive pad zone, or the like.

FIGS. **11**A and **11**B are perspective and side cross-sectional views, respectively, of the mounting unit immediately following sensor insertion and release of the applicator from the mounting unit. In one embodiment, such as illustrated in FIGS. **11**A and **11**B, the contact subassembly **26** is held in its insertion position, substantially at the insertion angle α of the sensor. Maintaining the contact subassembly **26** at the insertion angle α during insertion enables the sensor **32** to be easily inserted straight through the contact subassembly **26**. The contact subassembly **26** further includes a hinge **38** that allows movement of the contact subassembly **26** from an angled to a flat position. The term "hinge," as used herein, is a broad term and is used in its ordinary sense, including, without limitation, a mechanism that allows articulation of two or more parts or portions of a device. The term is broad enough to include a sliding hinge, for example, a ball and detent type hinging mechanism.

Although the illustrated embodiments describe a fixed insertion angle designed into the applicator, alternative embodiments can design the insertion angle into other components of the system. For example, the insertion angle can be designed into the attachment of the applicator with the mounting unit, or the like. In some alternative embodiments, a variety of adjustable insertion angles can be designed into the system to provide for a variety of host dermis configurations.

FIG. **11**B illustrates the sensor **32** extending from the mounting unit **14** by a preselected distance, which defines the depth of insertion of the sensor into the host. The dermal and subcutaneous make-up of animals and humans is variable and a fixed depth of insertion may not be appropriate for all implantations. Accordingly, in an alternative embodiment, the distance that the sensor extends from the mounting unit is adjustable to accommodate a variety of host body-types. For example, the applicator **12** can be designed with a variety of adjustable settings, which control the distance that the needle **72** (and therefore the sensor **32**) extends upon sensor insertion. One skilled in the art appreciates a variety of means and mechanisms can be employed to accommodate adjustable sensor insertion depths, which are considered within the scope of the preferred embodiments. The preferred insertion depth is from about 0.1 mm or less to about 2 cm or more, preferably from about 0.15, 0.2, 0.25, 0.3, 0.35, 0.4, or 0.45 mm to about 0.5, 0.6, 0.7, 0.8, 0.9, 1, 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, or 1.9 cm.

FIGS. **12**A and **12**B are perspective and side cross-sectional views, respectively, of the mounting unit after articulating the contact subassembly to its functional position (which is also referred to as an inserted, implanted, or sensing position). The hinge **38** enables the contact subassembly **26** to tilt from its insertion position (FIG. **11**) to its functional position (FIG. **12**) by pressing downward on the contact subassembly, for example. Certain embodiments provide this pivotal movement via two separate pieces (the contact subassembly **26** and the mounting unit **14** connected by a hinge, for example, a mechanical or adhesive pad joint or hinge). A variety of pivoting, articulating, and/or hinging mechanisms can be employed with the sensors of preferred embodiments. For example, the hinge can be formed as a part of the contact subassembly **26**. The contact subassembly

can be formed from a flexible piece of material (such as silicone, urethane rubber, or other flexible or elastomeric material), wherein the material is sufficiently flexible to enable bending or hinging of the contact subassembly from an angle appropriate for insertion (FIGS. **11**A and **11**B) to a lower functional configuration (FIGS. **12**A and **12**B).

The relative pivotal movement of the contact subassembly is advantageous, for example, for enabling the design of a low profile device while providing support for an appropriate needle insertion angle. In its insertion position, the sensor system is designed for easy sensor insertion while forming a stable electrical connection with the associated contacts **28**. In its functional position, the sensor system maintains a low profile for convenience, comfort, and discreetness during use. Thus, the sensor systems of preferred embodiments are advantageously designed with a hinging configuration to provide an optimum guided insertion angle while maintaining a low profile device during sensor use.

In some embodiments, a shock-absorbing member or feature is incorporated into the design of the sensor and configured to absorb movement of the in vivo and/or ex vivo portion of the sensor. Conventional analyte sensors can suffer from motion-related artifact associated with host movement when the host is using the device. For example, when a transcutaneous analyte sensor is inserted into the host, various movements on the sensor (for example, relative movement between the in vivo portion and the ex vivo portion and/or movement within the host) create stresses on the device and can produce noise in the sensor signal. Accordingly in some embodiments, a shock-absorbing member is located on the sensor/mounting unit in a location that absorbs stresses associated with the above-described movement.

In the preferred embodiments, the sensor **32** bends from a substantially straight to substantially bent configuration upon pivoting of the contact subassembly from the insertion to functional position. The substantially straight sensor configuration during insertion advantageously provides ease of sensor insertion, while the substantial bend in the sensor in its functional position advantageously provides stability on the proximal end of the sensor with flexibility/mobility on the distal end of the sensor. Additionally, motion within the mounting unit (e.g., caused by external forces to the mounting unit, movement of the skin, and the like) does not substantially translate to the in vivo portion of the sensor. Namely, the bend formed within the sensor **32** functions to break column strength, causing flexion that effectively absorbs movements on the sensor during use. Additionally, the sensor can be designed with a length such that when the contact subassembly **26** is pivoted to its functional position (FIG. **11**B), the sensor pushes forward and flexes, allowing it to absorb motion between the in vivo and ex vivo portions of the sensor. It is believed that both of the above advantages minimize motion artifact on the sensor signal and/or minimize damage to the sensor caused by movement, both of which (motion artifact and damage) have been observed in conventional transcutaneous sensors.

In some alternative embodiments, the shock-absorbing member can be an expanding and contracting member, such as a spring, accordion, telescoping, or bellows-type device. In general, the shock absorbing member can be located such that relative movement between the sensor, the mounting unit, and the host is absorbed without (or minimally) affecting the connection of the sensor to the mounting unit and/or the sensor stability within the implantation site; for example, the shock-absorbing member can be formed as a part of or connected to the sensor **32**.

FIGS. **13**A to **13**C are perspective and side views of a sensor system including the mounting unit **14** and electronics unit **16** attached thereto. After sensor insertion, the transcutaneous analyte sensor system **10** measures a concentration of an analyte or a substance indicative of the concentration or presence of the analyte as described above. Although the examples are directed to a glucose sensor, the analyte sensor can be a sensor capable of determining the level of any suitable analyte in the body, for example, oxygen, lactase, insulin, hormones, cholesterol, medicaments, viruses, or the like. Once the electronics unit **16** is connected to the mounting unit **14**, the sensor **32** is able to measure levels of the analyte in the host.

Detachable connection between the mounting unit **14** and electronics unit **16** provides improved manufacturability, namely, the relatively inexpensive mounting unit **14** can be disposed of when replacing the sensor system after its usable life, while the relatively more expensive electronics unit **16** can be reusable with multiple sensor systems. In certain embodiments, the electronics unit **16** is configured with programming, for example, initialization, calibration reset, failure testing, or the like, each time it is initially inserted into the cavity and/or each time it initially communicates with the sensor **32**. However, an integral (non-detachable) electronics unit can be configured as is appreciated by one skilled in the art.

Referring to the mechanical fit between the mounting unit **14** and the electronics unit **16** (and/or applicator **12**), a variety of mechanical joints are contemplated, for example, snap fit, interference fit, or slide fit. In the illustrated embodiment of FIGS. **13**A to **13**C, tabs **120** are provided on the mounting unit **14** and/or electronics unit **16** that enable a secure connection therebetween. The tabs **120** of the illustrated embodiment can improve ease of mechanical connection by providing alignment of the mounting unit and electronics unit and additional rigid support for force and counter force by the user (e.g., fingers) during connection. However, other configurations with or without guiding tabs are contemplated, such as illustrated in FIGS. **11** and **12**, for example.

In some circumstances, a drift of the sensor signal can cause inaccuracies in sensor performance and/or require re-calibration of the sensor. Accordingly, it can be advantageous to provide a sealant, whereby moisture (e.g., water and water vapor) cannot substantially penetrate to the sensor and its connection to the electrical contacts. The sealant described herein can be used alone or in combination with the sealing member **36** described in more detail above, to seal the sensor from moisture in the external environment.

Preferably, the sealant fills in holes, crevices, or other void spaces between the mounting unit **14** and electronics unit **16** and/or around the sensor **32** within the mounting unit **32**. For example, the sealant can surround the sensor in the portion of the sensor **32** that extends through the contacts **28**. Additionally, the sealant can be disposed within the additional void spaces, for example a hole **122** that extends through the sealing member **36**.

Preferably, the sealant comprises a water impermeable material or compound, for example, oil, grease, or gel. In one exemplary embodiment, the sealant, which also can be referred to as a lubricant in certain embodiments, comprises petroleum jelly and is used to provide a moisture barrier surrounding the sensor **32**. In one experiment, petroleum jelly was liquefied by heating, after which a sensor **32** was immersed into the liquefied petroleum jelly to coat the outer surfaces thereof. The sensor was then assembled into a housing and inserted into a host, during which deployment

the sensor was inserted through the electrical contacts **28** and the petroleum jelly conforming therebetween. Sensors incorporating petroleum jelly, such as described above, when compared to sensors without the petroleum jelly moisture barrier exhibited less or no signal drift over time when studied in a humid or submersed environment. While not wishing to be bound by theory, it is believed that incorporation of a moisture barrier surrounding the sensor, especially between the sensor and its associated electrical contacts, reduces or eliminates the effects of humidity on the sensor signal. The viscosity of grease or oil-based moisture barriers allows penetration into and through even small cracks or crevices within the sensor and mounting unit, displacing moisture and thereby increasing the sealing properties thereof. U.S. Pat. Nos. 4,259,540 and 5,285,513 disclose materials suitable for use as a water impermeable material (sealant).

Referring to the electrical fit between the sensor **32** and the electronics unit **16**, contacts **28** (through which the sensor extends) are configured to electrically connect with mutually engaging contacts on the electronics unit **16**. A variety of configurations are contemplated; however, the mutually engaging contacts operatively connect upon detachable connection of the electronics unit **16** with the mounting unit **14**, and are substantially sealed from external moisture by sealing member **36**. Even with the sealing member, some circumstances can exist wherein moisture can penetrate into the area surrounding the sensor **32** and or contacts, for example, exposure to a humid or wet environment (e.g., caused by sweat, showering, or other environmental causes). It has been observed that exposure of the sensor to moisture can be a cause of baseline signal drift of the sensor over time. For example in a glucose sensor, the baseline is the component of a glucose sensor signal that is not related to glucose (the amount of signal if no glucose is present), which is ideally constant over time. However, some circumstances my exist wherein the baseline can fluctuate over time, also referred to as drift, which can be caused, for example, by changes in a host's metabolism, cellular migration surrounding the sensor, interfering species, humidity in the environment, and the like.

In some embodiments, the mounting unit is designed to provide ventilation (e.g., a vent hole **124**) between the exit-site and the sensor. In certain embodiments, a filter (not shown) is provided in the vent hole **124** that allows the passage of air, while preventing contaminants from entering the vent hole **124** from the external environment. While not wishing to be bound by theory, it is believed that ventilation to the exit-site (or to the sensor **32**) can reduce or eliminate trapped moisture or bacteria, which can otherwise increase the growth and/or lifetime of bacteria adjacent to the sensor.

In some alternative embodiments, a sealing material is provided, which seals the needle and/or sensor from contamination of the external environment during and after sensor insertion. For example, one problem encountered in conventional transcutaneous devices is infection of the exit-site of the wound. For example, bacteria or contaminants can migrate from ex vivo, for example, any ex vivo portion of the device or the ex vivo environment, through the exit-site of the needle/sensor, and into the subcutaneous tissue, causing contamination and infection. Bacteria and/or contaminants can originate from handling of the device, exposed skin areas, and/or leakage from the mounting unit (external to) on the host. In many conventional transcutaneous devices, there exists some path of migration for bacteria and contaminants to the exit-site, which can become contaminated during sensor insertion or subsequent handling or use of the device. Furthermore, in some embodiments of a transcutaneous analyte sensor, the insertion-aiding device (for example, needle) is an integral part of the mounting unit; namely, the device stores the insertion device after insertion of the sensor, which is isolated from the exit-site (namely, point-of-entry of the sensor) after insertion.

Accordingly, these alternative embodiments provide a sealing material on the mounting unit, interposed between the housing and the skin, wherein the needle and/or sensor are adapted to extend through, and be sealed by, the sealing material. The sealing material is preferably formed from a flexible material that substantially seals around the needle/sensor. Appropriate flexible materials include malleable materials, elastomers, gels, greases, or the like (e.g., see U.S. Pat. Nos. 4,259,540 and 5,285,513). However, not all embodiments include a sealing material, and in some embodiments a clearance hole or other space surrounding the needle and/or sensor is preferred.

In one embodiment, the base **24** of the mounting unit **14** is formed from a flexible material, for example silicone, which by its elastomeric properties seals the needle and/or sensor at the exit port **126**, such as is illustrated in FIGS. **12**A and **12**B. Thus, sealing material can be formed as a unitary or integral piece with the back surface **25** of the mounting unit **14**, or with an adhesive pad **8** on the back surface of the mounting unit, however alternatively can be a separate part secured to the device. In some embodiments, the sealing material can extend through the exit port **126** above or below the plane of the adhesive pad surface, or the exit port **126** can comprise a septum seal such as those used in the medical storage and disposal industries (for example, silica gel sandwiched between upper and lower seal layers, such as layers comprising chemically inert materials such as PTFE). A variety of known septum seals can be implemented into the exit port of the preferred embodiments described herein. Whether the sealing material is integral with or a separate part attached to the mounting unit **14**, the exit port **126** is advantageously sealed so as to reduce or eliminate the migration of bacteria or other contaminants to or from the exit-site of the wound and/or within the mounting unit.

During use, a host or caretaker positions the mounting unit at the appropriate location on or near the host's skin and prepares for sensor insertion. During insertion, the needle aids in sensor insertion, after which the needle is retracted into the mounting unit leaving the sensor in the subcutaneous tissue. In this embodiment, the exit-port **126** includes a layer of sealing material, such as a silicone membrane, that encloses the exit-port in a configuration that protects the exit-site from contamination that can migrate from the mounting unit or spacing external to the exit-site. Thus, when the sensor **32** and/or needle **72** extend through, for example, an aperture or a puncture in the sealing material, to provide communication between the mounting unit and subcutaneous space, a seal is formed therebetween. Elastomeric sealing materials can be advantageous in some embodiments because the elasticity provides a conforming seal between the needle/sensor and the mounting unit and/or because the elasticity provides shock-absorbing qualities allowing relative movement between the device and the various layers of the host's tissue, for example.

In some alternative embodiments, the sealing material includes a bioactive agent incorporated therein. Suitable bioactive agents include those which are known to discourage or prevent bacteria and infection, for example, anti-inflammatory, antimicrobials, antibiotics, or the like. It is

believed that diffusion or presence of a bioactive agent can aid in prevention or elimination of bacteria adjacent to the exit-site.

In practice, after the sensor **32** has been inserted into the host's tissue, and an electrical connection formed by mating the electronics unit **16** to the mounting unit **14**, the sensor measures an analyte concentration continuously or continually, for example, at an interval of from about fractions of a second to about 10 minutes or more.

FIG. **14** is a perspective view of a sensor system, including wireless communication between a sensor and a receiver. Preferably the electronics unit **16** is wirelessly connected to a receiver **158** via one- or two-way RF transmissions or the like. However, a wired connection is also contemplated. The receiver **158** provides much of the processing and display of the sensor data, and can be selectively worn and/or removed at the host's convenience. Thus, the sensor system **10** can be discreetly worn, and the receiver **158**, which provides much of the processing and display of the sensor data, can be selectively worn and/or removed at the host's convenience. Particularly, the receiver **158** includes programming for retrospectively and/or prospectively initiating a calibration, converting sensor data, updating the calibration, evaluating received reference and sensor data, and evaluating the calibration for the analyte sensor, such as described in more detail with reference to U.S. Publication No. US-2005-0027463-A1.

FIGS. **15**A and **15**B are perspective views of a receiver in one preferred embodiment, wherein the receiver, also referred to as a communication station, is provided with a docking station for receiving and holding the electronics unit (from the sensor assembly) when not in use. Preferably, the electronics unit **16** is detachable from and reusable with multiple sensor assemblies **10**, in multiple applications, such as described herein. This embodiment provides a docking station **98** on the receiver **158** that enables storage, electrical connection, and/or battery recharge, for example, for the electronics unit **16** while not in use on a sensor assembly **10**. Complementary contacts or pins can be provided on the electronics unit **16** and the docking station **98** that enable operable connection therebetween. In some embodiments, the receiver includes programming that resets calibration when the electronics unit **16** is docked on the receiver **158**. In some embodiments, the receiver is configured to reset states in the battery, such as initializing a new (unused) electronics unit to work with the receiver when the electronics unit **16** is docked on the receiver **158**.

After a sensor's usable life, the host's removes and disposes of the sensor assembly **10**, saving the reusable electronics unit **16** for use with another sensor assembly, which can be a few minutes to a few days later, or more, after disposing of the previous sensor assembly. In some alternative embodiments, the docketing station is provided on an alternative communication station other than the receiver, for example a personal computer, server, personal digital assistant, or the like; wherein the functionality described below can be implemented in a similar manner. In some embodiments, the receiver is configured to test operation of the electronics unit by stepping the electrodes through different current draws and disabling usage of the electronics unit if a failure is detected.

Thus, the described embodiments provide an analyte sensor assembly that enables a comfortable and reliable system for measuring an analyte level for short term applications, e.g., up to 7 days or more, without surgery. After the usable life of the sensor (for example, due to a predetermined expiration, potential infection, or level of inflammation), the host can remove the sensor and mounting from the skin, dispose of the sensor and mounting unit (preferably saving the electronics unit for reuse). The reusable electronics unit can be inserted with another sensor assembly or be implanted surgically and thus provide continuous sensor output for short or long periods of time in another application. Data provided by the analyte sensor assembly can be used to calibrate other sensors, e.g., long term sensors including the implantable long term glucose sensor described hereinbelow.

Long Term Sensor

FIG. **16** is an exploded perspective view of one exemplary embodiment of a long term continuous glucose sensor **1310**A. In this embodiment, the sensor is preferably wholly implanted into the subcutaneous tissue of a host, such as described in U.S. Publication No. US-2006-0015020-A1; U.S. Publication No. US-2005-0245799-A1; U.S. Publication No. US-2005-0192557-A1; U.S. Publication No. US-2004-0199059-A1; U.S. Publication No. US-2005-0027463-A1; and U.S. Pat. No. 6,001,067. In this exemplary embodiment, a body **1320** and a sensing region **1321** house the electrodes **1322** and sensor electronics (see FIG. **17**). The three electrodes **1322** are operably connected to the sensor electronics (see FIG. **17**) and are covered by a sensing membrane **1323** and a biointerface membrane **1324**, which are attached by a clip **1325**.

In one embodiment, the three electrodes **1322** include a platinum working electrode, a platinum counter electrode, and a silver/silver chloride reference electrode. The top ends of the electrodes are in contact with an electrolyte phase (not shown), which is a free-flowing fluid phase disposed between the sensing membrane **1323** and the electrodes **1322**. The sensing membrane **1323** includes an enzyme, for example, glucose oxidase, and covers the electrolyte phase. The biointerface membrane **1324**, such as described above, covers the sensing membrane **1323** and serves, at least in part, to protect the sensor **1310**A from external forces that can result in environmental stress cracking of the sensing membrane **1323**. U.S. Publication No. US-2005-0112169-A1 describes a biointerface membrane that can be used in conjunction with the preferred embodiments.

In one embodiment, the biointerface membrane **1324** generally includes a cell disruptive domain most distal from the electrochemically reactive surfaces and a cell impermeable domain less distal from the electrochemically reactive surfaces than the cell disruptive domain. The cell disruptive domain is preferably designed to support tissue ingrowth, disrupt contractile forces typically found in a foreign body response, encourage vascularity within the membrane, and disrupt the formation of a barrier cell layer. The cell impermeable domain is preferably resistant to cellular attachment, impermeable to cells, and composed of a biostable material.

In one embodiment, the sensing membrane **1323** generally provides one or more of the following functions: 1) supporting tissue ingrowth; 2) protection of the exposed electrode surface from the biological environment, 3) diffusion resistance (limitation) of the analyte, 4) a catalyst for enabling an enzymatic reaction, 5) limitation or blocking of interfering species, and 6) hydrophilicity at the electrochemically reactive surfaces of the sensor interface, such as described in U.S. Publication No. US-2005-0245799-A1. Accordingly, the sensing membrane **1323** preferably includes a plurality of domains or layers, for example, an electrolyte domain, an interference domain, an enzyme domain (for example, glucose oxidase), a resistance domain, and can additionally include an oxygen domain (not shown), and/or a bioprotective domain (not shown), such as

described in more detail herein and in U.S. Publication No. US-2005-0245799-A1. However, it is understood that a sensing membrane modified for other devices, for example, by including fewer or additional domains is within the scope of the preferred embodiments.

In some embodiments, the domains of the biointerface and sensing membranes are formed from materials such as silicone, polytetrafluoroethylene, polyethylene-co-tetrafluoroethylene, polyolefin, polyester, polycarbonate, biostable polytetrafluoroethylene, homopolymers, copolymers, terpolymers of polyurethanes, polypropylene (PP), polyvinylchloride (PVC), polyvinylidene fluoride (PVDF), polybutylene terephthalate (PBT), polymethylmethacrylate (PMMA), polyether ether ketone (PEEK), polyurethanes, cellulosic polymers, polysulfones and block copolymers thereof including, for example, di-block, tri-block, alternating, random and graft copolymers. U.S. Publication No. US-2005-0245799-A1 describes biointerface and sensing membrane configurations and materials that can be applied to the preferred embodiments.

In the illustrated embodiment, the counter electrode is provided to balance the current generated by the species being measured at the working electrode. In the case of a glucose oxidase based glucose sensor, the species being measured at the working electrode is H_2O_2. Glucose oxidase catalyzes the conversion of oxygen and glucose to hydrogen peroxide and gluconate according to the following reaction:

$$\text{Glucose}+O_2 \rightarrow \text{Gluconate}+H_2O_2$$

The change in H_2O_2 can be monitored to determine glucose concentration because for each glucose molecule metabolized, there is a proportional change in the product H_2O_2. Oxidation of H_2O_2 by the working electrode is balanced by reduction of ambient oxygen, enzyme generated H_2O_2, or other reducible species at the counter electrode. The H_2O_2 produced from the glucose oxidase reaction further reacts at the surface of working electrode and produces two protons ($2H^+$), two electrons ($2e^-$), and one oxygen molecule (O_2).

In one embodiment, a potentiostat is employed to monitor the electrochemical reaction at the electrochemical cell. The potentiostat applies a constant potential to the working and reference electrodes to determine a current value. The current that is produced at the working electrode (and flows through the circuitry to the counter electrode) is substantially proportional to the amount of H_2O_2 that diffuses to the working electrode. Accordingly, a raw signal can be produced that is representative of the concentration of glucose in the user's body, and therefore can be utilized to estimate a meaningful glucose value, such as is described herein.

Sensor Electronics

One example of sensor electronics that may be utilized with either the short term or long term sensors is depicted in the block diagram of FIG. **17**, illustrating the electronics **132** associated with the sensor system in one embodiment. In this embodiment, a potentiostat **134** is shown, which is operably connected to an electrode system (such as described above) and provides a voltage to the electrodes, which biases the sensor to enable measurement of an current signal indicative of the analyte concentration in the host (also referred to as the analog portion). In some embodiments, the potentiostat includes a resistor (not shown) that translates the current into voltage. In some alternative embodiments, a current to frequency converter is provided that is configured to continuously integrate the measured current, for example, using a charge counting device.

An A/D converter **136** digitizes the analog signal into a digital signal, also referred to as "counts" for processing. Accordingly, the resulting raw data stream in counts, also referred to as raw sensor data, is directly related to the current measured by the potentiostat **134**.

A processor module **138** includes the central control unit that controls the processing of the sensor electronics **132**. In some embodiments, the processor module includes a microprocessor, however a computer system other than a microprocessor can be used to process data as described herein, for example an ASIC can be used for some or all of the sensor's central processing. The processor typically provides semi-permanent storage of data, for example, storing data such as sensor identifier (ID) and programming to process data streams (for example, programming for data smoothing and/or replacement of signal artifacts such as is described in U.S. Publication No. US-2005-0043598-A1). The processor additionally can be used for the system's cache memory, for example for temporarily storing recent sensor data. In some embodiments, the processor module comprises memory storage components such as ROM, RAM, dynamic-RAM, static-RAM, non-static RAM, EEPROM, rewritable ROMs, flash memory, or the like.

In some embodiments, the processor module comprises a digital filter, for example, an infinite impulse response (IIR) or finite impulse response (FIR) filter, configured to smooth the raw data stream from the A/D converter. Generally, digital filters are programmed to filter data sampled at a predetermined time interval (also referred to as a sample rate). In some embodiments, wherein the potentiostat is configured to measure the analyte at discrete time intervals, these time intervals determine the sample rate of the digital filter. In some alternative embodiments, wherein the potentiostat is configured to continuously measure the analyte, for example, using a current-to-frequency converter as described above, the processor module can be programmed to request a digital value from the A/D converter at a predetermined time interval, also referred to as the acquisition time. In these alternative embodiments, the values obtained by the processor are advantageously averaged over the acquisition time due the continuity of the current measurement. Accordingly, the acquisition time determines the sample rate of the digital filter. In preferred embodiments, the processor module is configured with a programmable acquisition time, namely, the predetermined time interval for requesting the digital value from the A/D converter is programmable by a user within the digital circuitry of the processor module. An acquisition time of from about 2 seconds to about 512 seconds is preferred; however any acquisition time can be programmed into the processor module. A programmable acquisition time is advantageous in optimizing noise filtration, time lag, and processing/battery power.

Preferably, the processor module is configured to build the data packet for transmission to an outside source, for example, an RF transmission to a receiver as described in more detail below. Generally, the data packet comprises a plurality of bits that can include a preamble, a unique identifier identifying the electronics unit, the receiver, or both, (e.g., sensor ID code), data (e.g., raw data, filtered data, and/or an integrated value) and/or error detection or correction. Preferably, the data (transmission) packet has a length of from about 8 bits to about 128 bits, preferably about 48 bits; however, larger or smaller packets can be desirable in certain embodiments. The processor module can be configured to transmit any combination of raw and/or filtered data. In one exemplary embodiment, the transmission packet

contains a fixed preamble, a unique ID of the electronics unit, a single five-minute average (e.g., integrated) sensor data value, and a cyclic redundancy code (CRC).

In some embodiments, the processor module further comprises a transmitter portion that determines the transmission interval of the sensor data to a receiver, or the like. In some embodiments, the transmitter portion, which determines the interval of transmission, is configured to be programmable. In one such embodiment, a coefficient can be chosen (e.g., a number of from about 1 to about 100, or more), wherein the coefficient is multiplied by the acquisition time (or sampling rate), such as described above, to define the transmission interval of the data packet. Thus, in some embodiments, the transmission interval is programmable from about 2 seconds to about 850 minutes, more preferably from about 30 second to about 5 minutes; however, any transmission interval can be programmable or programmed into the processor module. However, a variety of alternative systems and methods for providing a programmable transmission interval can also be employed. By providing a programmable transmission interval, data transmission can be customized to meet a variety of design criteria (e.g., reduced battery consumption, timeliness of reporting sensor values, etc.)

Conventional glucose sensors measure current in the nanoAmp range. In contrast to conventional glucose sensors, the preferred embodiments are configured to measure the current flow in the picoAmp range, and in some embodiments, femtoAmps. Namely, for every unit (mg/dL) of glucose measured, at least one picoAmp of current is measured. Preferably, the analog portion of the A/D converter **136** is configured to continuously measure the current flowing at the working electrode and to convert the current measurement to digital values representative of the current. In one embodiment, the current flow is measured by a charge counting device (e.g., a capacitor). Preferably, a charge counting device provides a value (e.g., digital value) representative of the current flow integrated over time (e.g., integrated value). In some embodiments, the value is integrated over a few seconds, a few minutes, or longer. In one exemplary embodiment, the value is integrated over 5 minutes; however, other integration periods can be chosen. Thus, a signal is provided, whereby a high sensitivity maximizes the signal received by a minimal amount of measured hydrogen peroxide (e.g., minimal glucose requirements without sacrificing accuracy even in low glucose ranges), reducing the sensitivity to oxygen limitations in vivo (e.g., in oxygen-dependent glucose sensors).

In some embodiments, the electronics unit is programmed with a specific ID, which is programmed (automatically or by the user) into a receiver to establish a secure wireless communication link between the electronics unit and the receiver. Preferably, the transmission packet is Manchester encoded; however, a variety of known encoding techniques can also be employed.

A battery **144** is operably connected to the sensor electronics **132** and provides the power for the sensor. In one embodiment, the battery is a lithium manganese dioxide battery; however, any appropriately sized and powered battery can be used (for example, AAA, nickel-cadmium, zinc-carbon, alkaline, lithium, nickel-metal hydride, lithium-ion, zinc-air, zinc-mercury oxide, silver-zinc, and/or hermetically-sealed). In some embodiments, the battery is rechargeable, and/or a plurality of batteries can be used to power the system. The sensor can be transcutaneously powered via an inductive coupling, for example. In some embodiments, a quartz crystal **96** is operably connected to the processor **138** and maintains system time for the computer system as a whole, for example for the programmable acquisition time within the processor module.

Optional temperature probe **140** is shown, wherein the temperature probe is located on the electronics assembly or the glucose sensor itself. The temperature probe can be used to measure ambient temperature in the vicinity of the glucose sensor. This temperature measurement can be used to add temperature compensation to the calculated glucose value.

An RF module **148** is operably connected to the processor **138** and transmits the sensor data from the sensor to a receiver within a wireless transmission **150** via antenna **152**. In some embodiments, a second quartz crystal **154** provides the time base for the RF carrier frequency used for data transmissions from the RF transceiver. In some alternative embodiments, however, other mechanisms, such as optical, infrared radiation (IR), ultrasonic, or the like, can be used to transmit and/or receive data.

In the RF telemetry module of the preferred embodiments, the hardware and software are designed for low power requirements to increase the longevity of the device (for example, to enable a life of from about 3 to about 24 months, or more) with maximum RF transmittance from the in vivo environment to the ex vivo environment for wholly implantable sensors (for example, a distance of from about one to ten meters or more). Preferably, a high frequency carrier signal of from about 402 MHz to about 433 MHz is employed in order to maintain lower power requirements. In some embodiments, the RF module employs a one-way RF communication link to provide a simplified ultra low power data transmission and receiving scheme. The RF transmission can be OOK or FSK modulated, preferably with a radiated transmission power (EIRP) fixed at a single power level of typically less than about 100 microwatts, preferably less than about 75 microwatts, more preferably less than about 50 microwatts, and most preferably less than about 25 microwatts.

Additionally, in wholly implantable devices, the carrier frequency is adapted for physiological attenuation levels, which is accomplished by tuning the RF module in a simulated in vivo environment to ensure RF functionality after implantation; accordingly, the preferred glucose sensor can sustain sensor function for 3 months, 6 months, 12 months, or 24 months or more.

When a sensor is first implanted into host tissue, the sensor and receiver are initialized. This is referred to as start-up mode, and involves optionally resetting the sensor data and calibrating the sensor **32**. In selected embodiments (e.g., transcutaneous sensors), mating the electronics unit **16** to the mounting unit **14** triggers a start-up mode. In other embodiments, the start-up mode is triggered by the receiver, which is described in more detail with reference to FIG. **21**, below.

Preferably, the electronics unit **16** indicates to the receiver (FIGS. **14** and **17**) that calibration is to be initialized (or re-initialized). The electronics unit **16** transmits a series of bits within a transmitted data packet wherein a sensor code can be included in the periodic transmission of the device. The status code is used to communicate sensor status to the receiving device. The status code can be inserted into any location in the transmitted data packet, with or without other sensor information. In one embodiment, the status code is designed to be unique or near unique to an individual sensor, which can be accomplished using a value that increments, decrements, or changes in some way after the transmitter detects that a sensor has been removed and/or attached to the

transmitter. In an alternative embodiment, the status code can be configured to follow a specific progression, such as a BCD interpretation of a Gray code.

In some embodiments (e.g., transcutaneous sensors), the sensor electronics **132** are configured to detect a current drop to zero in the working electrode **44** associated with removal of a sensor **32** from the host (or the electronics unit **16** from the mounting unit **14**), which can be configured to trigger an increment of the status code. If the incremented value reaches a maximum, it can be designed to roll over to 0. In some embodiments, the sensor electronics are configured to detect a voltage change cycle associated with removal and/or re-insertion of the sensor, which can be sensed in the counter electrode (e.g., of a three-electrode sensor), which can be configured to trigger an increment of the status code.

In some embodiments, the sensor electronics **132** can be configured to send a special value (for example, 0) that indicates that the electronics unit is not attached when removal of the sensor (or electronics unit) is detected. This special value can be used to trigger a variety of events, for example, to halt display of analyte values. Incrementing or decrementing routines can be used to skip this special value.

Data Smoothing

Typically, an analyte sensor (both short and long term) produces a raw data signal that is indicative of the analyte concentration of a user. The above described glucose sensor is only one example of an abundance of analyte sensors that are able to provide a raw data signal output indicative of the concentration of the analyte of interest. The devices and methods of the preferred embodiments, including data smoothing, calibration, evaluation, and other data processing, can be applied to raw data obtained from any analyte sensor capable of producing a output signal.

Raw data signals received from an analyte sensor can include signal noise, which degrades the quality of the data. The use of smoothing algorithms help improve the signal-to-noise ratio in the sensor by reducing signal jitter, for example. One example of a conventional data smoothing algorithms include finite impulse response filter (FIR), which is particularly suited for reducing high-frequency noise (see Steil et al. U.S. Pat. No. 6,558,351). Other analyte sensors have utilized heuristic and moving average type algorithms to accomplish data smoothing of signal jitter in data signals, for example.

It is advantageous to also reduce signal noise by attenuating transient, low frequency, non-analyte related signal fluctuations (e.g., transient ischemia and/or long transient periods of postural effects that interfere with sensor function due to lack of oxygen and/or other physiological effects).

In one embodiment, this attenuation of transient low frequency non-analyte related signal noise is accomplished using a recursive filter. In contrast to conventional non-recursive (e.g., FIR) filters in which each computation uses new input data sets, a recursive filter is an equation that uses moving averages as inputs; that is, a recursive filter includes previous averages as part of the next filtered output. Recursive filters are advantageous at least in part due to their computational efficiency.

FIG. **18** is a graph that illustrates data smoothing of a raw data signal in one embodiment. In this embodiment, the recursive filter is implemented as a digital infinite impulse response filter (IIR) filter, wherein the output is computed using 6 additions and 7 multiplies as shown in the following equation:

$$y(n) = \frac{a_0 * x(n) + a_1 * (n-1) + a_2 * x(n-2) + a_3 * x(n-3) - b_1 * y(n-1) - b_2 * y(n-2) - b_3 * y(n-3)}{b_0}$$

This polynomial equation includes coefficients that are dependent on sample rate and frequency behavior of the filter. In this exemplary embodiment, frequency behavior passes low frequencies up to cycle lengths of 40 minutes, and is based on a 30 second sample rate.

In some embodiments, data smoothing can be implemented in the sensor and the smoothed data transmitted to a receiver for additional processing. In other embodiments, raw data can be sent from the sensor to a receiver for data smoothing and additional processing therein. In yet other embodiments, the sensor is integral with the receiver and therefore no transmission of data is required.

In one exemplary embodiment, where the sensor is an implantable glucose sensor, data smoothing is performed in the sensor to ensure a continuous stream of data. In alternative embodiments, data smoothing can be transmitted from the sensor to the receiver, and the data smoothing performed at the receiver; however that there can be a risk of transmit-loss in the radio transmission from the sensor to the receiver when the transmission is wireless. For example, in embodiments wherein a sensor is implemented in vivo, the raw sensor signal can be more consistent within the sensor (in vivo) than the raw signal transmitted to a source (e.g., receiver) outside the body (e.g., if a patient were to take the receiver off to shower, communication between the sensor and receiver can be lost and data smoothing in the receiver would halt accordingly.) Consequently, a multiple point data loss in the filter can take, for example, anywhere from 25 to 40 minutes for the smoothed data to recover to where it would have been had there been no data loss.

Other systems and methods for data smoothing and/or for detecting and/or replacing certain sensor data (e.g., signal artifacts or system noise) are described in U.S Publication No. US-2005-0043598-A1.

Receiver

FIGS. **19**A to **19**D are schematic views of a receiver in first, second, third, and fourth embodiments, respectively. A receiver **1640** comprises systems necessary to receive, process, and display sensor data from an analyte sensor, such as described elsewhere herein. Particularly, the receiver **1640** can be a pager-sized device, for example, and comprise a user interface that has a plurality of buttons **1642** and a liquid crystal display (LCD) screen **1644**, and which can include a backlight. In some embodiments the user interface can also include a keyboard, a speaker, and a vibrator such as described with reference to FIG. **20**A.

In some embodiments a user is able to toggle through some or all of the screens shown in FIGS. **19**A to **19**D using a toggle button on the receiver. In some embodiments, the user is able to interactively select the type of output displayed on their user interface. In some embodiments, the sensor output can have alternative configurations.

FIG. **20**A is a block diagram that illustrates the configuration of the medical device in one embodiment, including a continuous analyte sensor, a receiver, and an external device. In general, the analyte sensor system is any sensor configuration that provides an output signal indicative of a concentration of an analyte (e.g., invasive, minimally-invasive, and/or non-invasive sensors as described above). The output signal is sent to a receiver **158** and received by an input module **174**, which is described in more detail below.

The output signal is typically a raw data stream that is used to provide a useful value of the measured analyte concentration to a patient or a doctor, for example. In some embodiments, the raw data stream can be continuously or periodically algorithmically smoothed or otherwise modified to diminish outlying points that do not accurately represent the analyte concentration, for example due to signal noise or other signal artifacts, such as described in U.S. Pat. No. 6,931,327.

Referring again to FIG. **20**A, the receiver **158**, which is operatively linked to the sensor system **10**, receives a data stream from the sensor system **10** via the input module **174**. In one embodiment, the input module includes a quartz crystal operably connected to an RF transceiver (not shown) that together function to receive and synchronize data streams from the sensor system **10**. However, the input module **174** can be configured in any manner that is capable of receiving data from the sensor. Once received, the input module **174** sends the data stream to a processor **176** that processes the data stream, such as is described in more detail below.

The processor **176** is the central control unit that performs the processing, such as storing data, analyzing data streams, calibrating analyte sensor data, estimating analyte values, comparing estimated analyte values with time corresponding measured analyte values, analyzing a variation of estimated analyte values, downloading data, and controlling the user interface by providing analyte values, prompts, messages, warnings, alarms, or the like. The processor includes hardware and software that performs the processing described herein, for example flash memory provides permanent or semi-permanent storage of data, storing data such as sensor ID, receiver ID, and programming to process data streams (for example, programming for performing estimation and other algorithms described elsewhere herein) and random access memory (RAM) stores the system's cache memory and is helpful in data processing.

Preferably, the input module **174** or processor module **176** performs a Cyclic Redundancy Check (CRC) to verify data integrity, with or without a method of recovering the data if there is an error. In some embodiments, error correction techniques such as those that use Hamming codes or Reed-Solomon encoding/decoding methods are employed to correct for errors in the data stream. In one alternative embodiment, an iterative decoding technique is employed, wherein the decoding is processed iteratively (e.g., in a closed loop) to determine the most likely decoded signal. This type of decoding can allow for recovery of a signal that is as low as 0.5 dB above the noise floor, which is in contrast to conventional non-iterative decoding techniques (such as Reed-Solomon), which requires approximately 3 dB or about twice the signal power to recover the same signal (e.g., a turbo code).

An output module **178**, which is integral with and/or operatively connected with the processor **176**, includes programming for generating output based on the data stream received from the sensor system **10** and its processing incurred in the processor **176**. In some embodiments, output is generated via a user interface **160**.

The user interface **160** comprises a keyboard **162**, speaker **164**, vibrator **166**, backlight **168**, liquid crystal display (LCD) screen **170**, and one or more buttons **172**. The components that comprise the user interface **160** include controls to allow interaction of the user with the receiver. The keyboard **162** can allow, for example, input of user information about himself/herself, such as mealtime, exercise, insulin administration, customized therapy recommendations, and reference analyte values. The speaker **164** can produce, for example, audible signals or alerts for conditions such as present and/or estimated hyperglycemic or hypoglycemic conditions in a person with diabetes. The vibrator **166** can provide, for example, tactile signals or alerts for reasons such as described with reference to the speaker, above. The backlight **168** can be provided, for example, to aid the user in reading the LCD **170** in low light conditions. The LCD **170** can be provided, for example, to provide the user with visual data output, such as is described in U.S. Publication No. US-2005-0203360-A1. FIGS. **20**B to **20**D illustrate some additional visual displays that can be provided on the screen **170**. In some embodiments, the LCD is a touch-activated screen, enabling each selection by a user, for example, from a menu on the screen. The buttons **172** can provide for toggle, menu selection, option selection, mode selection, and reset, for example. In some alternative embodiments, a microphone can be provided to allow for voice-activated control.

In some embodiments, prompts or messages can be displayed on the user interface to convey information to the user, such as reference outlier values, requests for reference analyte values, therapy recommendations, deviation of the measured analyte values from the estimated analyte values, or the like. Additionally, prompts can be displayed to guide the user through calibration or trouble-shooting of the calibration.

In some embodiments, the receiver and/or a device connected to the receiver is configured to audibly output the user's analyte value(s), trend information (increasing or decreasing analyte values), and the like, hereinafter referred to as the audible output module. In some embodiments, the audible output module additionally includes: high and low blood glucose limits at which the module will audibly output the user's analyte value and/or trend information; English and non-English language versions; and choice of male or female voice. In some embodiments, the audible output is transmitted to an earbud worn by the patient for use where privacy is required or by a patient who is somewhat audibly impaired. The audible output module can be particularly advantageous in applications wherein the user is visually and/or hearing impaired, or is unable to visually check their receiver due to other circumstances (e.g., operating a motor vehicle or machinery, engaged in a business meeting or social event, or the like).

Additionally, data output from the output module **178** can provide wired or wireless, one- or two-way communication between the receiver **158** and an external device **180**. The external device **180** can be any device that wherein interfaces or communicates with the receiver **158**. In some embodiments, the external device **180** is a computer, and the receiver **158** is able to download historical data for retrospective analysis by the patient or physician, for example. In some embodiments, the external device **180** is a modem or other telecommunications station, and the receiver **158** is able to send alerts, warnings, emergency messages, or the like, via telecommunication lines to another party, such as a doctor or family member. In some embodiments, the external device **180** is an insulin pen, and the receiver **158** is able to communicate therapy recommendations, such as insulin amount and time to the insulin pen. In some embodiments, the external device **180** is an insulin pump, and the receiver **158** is able to communicate therapy recommendations, such as insulin amount and time to the insulin pump. The external device **180** can include other technology or medical devices, for example pacemakers, implanted analyte sensor patches, other infusion devices, telemetry devices, or the like.

The user interface **160**, including keyboard **162**, buttons **172**, a microphone (not shown), and the external device **180**, can be configured to allow input of data. Data input can be helpful in obtaining information about the patient (for example, meal time, exercise, or the like), receiving instructions from a physician (for example, customized therapy recommendations, targets, or the like), and downloading software updates, for example. Keyboard, buttons, touchscreen, and microphone are all examples of mechanisms by which a user can input data directly into the receiver. A server, personal computer, personal digital assistant, insulin pump, and insulin pen are examples of external devices that can provide useful information to the receiver. Other devices internal or external to the sensor that measure other aspects of a patient's body (for example, temperature sensor, accelerometer, heart rate monitor, oxygen monitor, or the like) can be used to provide input helpful in data processing. In one embodiment, the user interface can prompt the patient to select an activity most closely related to their present activity, which can be helpful in linking to an individual's physiological patterns, or other data processing. In another embodiment, a temperature sensor and/or heart rate monitor can provide information helpful in linking activity, metabolism, and glucose excursions of an individual. While a few examples of data input have been provided here, a variety of information can be input, which can be helpful in data processing.

FIG. **20**B is an illustration of an LCD screen **170** showing continuous and single point glucose information in the form of a trend graph **184** and a single numerical value **186**. The trend graph shows upper and lower boundaries **182** representing a target range between which the host should maintain his/her glucose values. Preferably, the receiver is configured such that these boundaries **182** can be configured or customized by a user, such as the host or a care provider. By providing visual boundaries **182**, in combination with continuous analyte values over time (e.g., a trend graph **184**), a user can better learn how to control his/her analyte concentration (e.g., a person with diabetes can better learn how to control his/her glucose concentration) as compared to single point (single numerical value **186**) alone. Although FIG. **20**B illustrates a 1 hour trend graph (e.g., depicted with a time range **188** of 1 hour), a variety of time ranges can be represented on the screen **170**, for example, 3 hour, 9 hour, 1 day, and the like.

FIG. **20**C is an illustration of an LCD screen **170** showing a low alert screen that can be displayed responsive to a host's analyte concentration falling below a lower boundary (see boundaries **182**). In this exemplary screen, a host's glucose concentration has fallen to 55 mg/dL, which is below the lower boundary set in FIG. **20**B, for example. The arrow **190** represents the direction of the analyte trend, for example, indicating that the glucose concentration is continuing to drop. The annotation **192** ("LOW") is helpful in immediately and clearly alerting the host that his/her glucose concentration has dropped below a preset limit, and what may be considered to be a clinically safe value, for example. FIG. **20**D is an illustration of an LCD screen **170** showing a high alert screen that can be displayed responsive to a host's analyte concentration rising above an upper boundary (see boundaries **182**). In this exemplary screen, a host's glucose concentration has risen to 200 mg/dL, which is above a boundary set by the host, thereby triggering the high alert screen. The arrow **190** represents the direction of the analyte trend, for example, indicating that the glucose concentration is continuing to rise. The annotation **192** ("HIGH") is helpful in immediately and clearly alerting the host that his/her glucose concentration has above a preset limit, and what may be considered to be a clinically safe value, for example.

Although a few exemplary screens are depicted herein, a variety of screens can be provided for illustrating any of the information described in the preferred embodiments, as well as additional information. A user can toggle between these screens (e.g., using buttons **172**) and/or the screens can be automatically displayed responsive to programming within the receiver **158**, and can be simultaneously accompanied by another type of alert (audible or tactile, for example).

In some embodiments the receiver **158** can have a length of from about 8 cm to about 15 cm, a width of from about 3.5 cm to about 10 cm, and/or a thickness of from about 1 cm to about 3.5 cm. In some embodiments the receiver **158** can have a volume of from about 120 cm^3 to about 180 cm^3, and can have a weight of from about 70 g to 130 g. The dimensions and volume can be higher or lower, depending, e.g., on the type of devices integrated (e.g., finger stick devices, pumps, PDAs, and the like.), the type of user interface employed, and the like.

In some embodiments, the receiver **158** is an application-specific device. In some embodiments the receiver **158** can be a device used for other functions, such as are described in U.S. Pat. No. 6,558,320. For example, the receiver **158** can be integrated into a personal computer (PC), a personal digital assistant (PDA), a cell phone, or another fixed or portable computing device. The integration of the receiver **158** function into a more general purpose device can comprise the addition of software and/or hardware to the device. Communication between the sensor electronics **16** and the receiver **158** function of the more general purpose device can be implemented with wired or wireless technologies. For example, a PDA can be configured with a data communications port and/or a wireless receiver. After the user establishes a communication link between the electronics unit **16** and the PDA, the electronics unit **16** transmits data to the PDA which then processes the data according to software which has been loaded thereon so as to display.

Sensor Calibration

Reference is now made to FIG. **21**, which is a flow chart that illustrates the initial calibration and data output of sensor data from a sensor, e.g., a short term or long term analyte sensor. Calibration of an analyte sensor comprises data processing that uses sensor data and other information to determine an estimated analyte measurement that is meaningful to a user or useful for further data processing. One or more reference values can be used to calibrate the sensor data. In some embodiments, reference values can be obtained from sensor data of another analyte sensor. For example, a substantially continuous short term sensor can be used to calibrate another substantially continuous short term sensor or a substantially continuous long term sensor, or a substantially continuous long term sensor can be used to calibrate another substantially continuous long term sensor or a substantially continuous short term sensor. Reference values can also be obtained from a non-continuous monitoring system, such as a self-monitored blood analyte test (e.g., finger stick blood samples), an optical sensor, etc. However, using sensor data from an already employed substantially continuous short term sensor or a substantially continuous long term sensor can reduce or obviate the need for non-continuous manual measurements to obtain reference data. In addition, reference values can be obtained from in-vitro benchtop sources during a benchtop calibration process. For example, reference solutions of analyte may be

used for in-vitro calibration where the reference value for the concentration is known from the reference solution preparation.

FIG. **21** illustrates initial calibration of an analyte sensor using reference data from any reference analyte source, including another analyte sensor (for example, a previously calibrated analyte sensor). At block **261**, a sensor data receiving module, also referred to as the sensor data module, receives sensor data (e.g., a data stream), including one or more time-spaced sensor data points, from a sensor via the receiver, which can be in wired or wireless communication with the sensor. The sensor data point(s) can be smoothed, such as described with reference to FIG. **18**, above. During the initialization of the sensor, prior to initial calibration, the receiver (e.g., computer system) receives and stores the sensor data. However, the receiver can be configured so that data is not displayed to a user until initial calibration and possibly stabilization of the sensor has been determined.

At block **262**, a reference data receiving module, also referred to as the reference input module, receives reference data from a reference analyte source, including one or more reference data points. In one embodiment, the reference data is based on sensor data from another substantially continuous analyte sensor, e.g., an analyte sensor described herein, or another type of suitable analyte sensor. For example, a previously calibrated short or long term glucose sensor providing sensor data for a host can provide reference data for use in calibrating sensor data from a long term glucose sensor implanted in the same host.

At block **263**, a data matching module, also referred to as the processor module, matches reference data (e.g., one or more reference analyte data points) with substantially time corresponding sensor data (e.g., one or more sensor data points) to provide one or more matched data pairs. In one embodiment, one reference data point is matched to one time corresponding sensor data point to form a matched data pair. In another embodiment, a plurality of reference data points are averaged (e.g., equally or non-equally weighted average, mean-value, median, or the like) and matched to one time corresponding sensor data point to form a matched data pair. In another embodiment, one reference data point is matched to a plurality of time corresponding sensor data points averaged to form a matched data pair. In yet another embodiment, a plurality of reference data points are averaged and matched to a plurality of time corresponding sensor data points averaged to form a matched data pair. In yet another embodiment, a line is produced from the reference data points as a function of time. The line may be produced by connecting individual data points, from smoothed data points, or as a best-fit through the application of an appropriate algorithm. A line produced from a series of sensor data points may be matched to the line produced from the reference data points.

To properly associate the sensor data with reference data, a delay in the sensing process can be taken into account. In one embodiment, the reference data is provided by another substantially continuous source that has similar sensor characteristics so that the delay in the sensor data is similar to the delay in the reference data. If the delay is not similar, the data can be appropriately processed to associate the data taken at the same time. In one embodiment where the sensor data is being associated with data from a non-continuous source, time corresponding sensor data comprises one or more sensor data points that occur 5 min (e.g., +/−2½ minutes) after the reference analyte data timestamp (e.g., the time that the reference analyte data is obtained). In this embodiment, the 15 minute time delay has been chosen to account for an approximately 10 minute delay introduced by the filter used in data smoothing and an approximately 5 minute physiological time-lag (e.g., the time necessary for the analyte to diffusion through a membrane(s) of an analyte sensor). In alternative embodiments, the time corresponding sensor value can be more or less than the above-described embodiment, for example ±60 minutes. Variability in time correspondence of sensor and reference data can be attributed to, for example a longer or shorter time delay introduced by the data smoothing filter, or if the configuration of the analyte sensor incurs a greater or lesser physiological time lag. In one exemplary embodiment, time delay information from a short term sensor can be used by a long term sensor as described above. In another exemplary embodiment, time delay information from a first short term sensor can be used by a second short term sensor as described above. In yet another exemplary embodiment, time delay information from a first long term sensor can be used by a second long term sensor as described above.

In some embodiments, tests are used to evaluate the best matched pair using a reference data point against individual sensor values over a predetermined time period (e.g., about 30 minutes). In one such exemplary embodiment, the reference data point is matched with sensor data points at 5-minute intervals and each matched pair is evaluated. The matched pair with the best correlation can be selected as the matched pair for data processing. In some alternative embodiments, matching a reference data point with an average of a plurality of sensor data points over a predetermined time period can be used to form a matched pair.

At block **264**, a calibration set module, also referred to as the processor module, forms an initial calibration set from a set of one or more matched data pairs, which are used to determine the relationship between the reference analyte data and the sensor analyte data, such as will be described in more detail with reference to block **267**, below.

The matched data pairs, which make up the initial calibration set, can be selected according to predetermined criteria. The criteria for the initial calibration set can be the same as, or different from, the criteria for the update calibration set, which is described in more detail with reference to FIG. **24**. In some embodiments, the number (n) of data pair(s) selected for the initial calibration set is one. In other embodiments, n data pairs are selected for the initial calibration set wherein n is a function of the frequency of the received reference data points. For example, a substantially continuous analyte sensor can provide numerous data points for use as reference data (e.g., dozens or even hundreds). In one exemplary embodiment, a substantially continuous analyte sensor provides about 288 reference data points per day (about every five minutes for twenty-four hours). In one exemplary embodiment, six data pairs make up the initial calibration set.

In some embodiments, the data pairs are selected only within a certain analyte value threshold, for example wherein the reference analyte value is from about 40 and to about 400 mg/dL. In some embodiments, the data pairs that form the initial calibration set are selected according to their time stamp. In some embodiments, the calibration set is selected such as described with reference to FIG. **24**.

At block **265**, an optional stability determination module, also referred to as the start-up module, determines the stability of the analyte sensor over a period of time. Some analyte sensors can have an initial instability time period during which the analyte sensor is unstable for environmental, physiological, or other reasons. One example of initial sensor instability is an embodiment wherein the analyte

sensor is implanted subcutaneously; in this example embodiment, stabilization of the analyte sensor can be dependent upon the maturity of the tissue ingrowth around and within the sensor. Another example of initial sensor instability is in an embodiment wherein the analyte sensor is inserted transdermally (e.g., transcutaneously); in this example embodiment, stabilization of the analyte sensor can be dependent upon electrode stabilization and/or sweat, for example.

Accordingly, in some embodiments, determination of sensor stability can include waiting a predetermined time period (e.g., an implantable sensor is known to require a time period for tissue ingrowth, and a transdermal (e.g., transcutaneous) sensor is known to require time to equilibrate the sensor with the user's tissue); in some embodiments, this waiting period is from about one minute to about six weeks. Although in some embodiments, the sensitivity (e.g., sensor signal strength with respect to analyte concentration) can be used to determine the stability of the sensor; for example, amplitude and/or variability of sensor sensitivity can be evaluated to determine the stability of the sensor. In alternative embodiments, detection of pH levels, oxygen, hypochlorite, interfering species (e.g., ascorbate, urea, and acetaminophen), correlation between sensor and reference values (e.g., R-value), baseline drift and/or offset, and the like can be used to determine the stability of the sensor. In one exemplary embodiment, wherein the sensor is a glucose sensor, it is known to provide a signal that is associated with interfering species (e.g., ascorbate, urea, acetaminophen), which can be used to evaluate sensor stability. In another exemplary embodiment, wherein the sensor is a glucose sensor such as described with reference to FIGS. **8** and **9**, the counter electrode can be monitored for oxygen deprivation, which can be used to evaluate sensor stability or functionality. Numerous other systems and methods for evaluating sensor stability or functionality are described in U.S. Publication No. US-2005-0043598-A1. In one embodiment employing a long term implantable sensor that requires a waiting period, a short term sensor or a series of short term sensors employed for the same user provides sensor data to the user during the waiting period. In an embodiment employing a series of short term sensors, the sensors can be employed so that they provide sensor data in discrete or overlapping periods. In such embodiments, the sensor data from one short term sensor can be used to calibrate another short term sensor, or be used to confirm the validity of a subsequently employed short term sensor. The data can also be used for calibrating the implantable long term sensor so that data from the long term sensor can be used immediately upon completion of the waiting period.

At decision block **266**, the system (e.g., microprocessor) determines whether the analyte sensor is sufficiently stable according to certain criteria, such as described above. In one embodiment wherein the sensor is an implantable glucose sensor, the system waits a predetermined time period believed necessary for sufficient tissue ingrowth and evaluates the sensor sensitivity (e.g., from about one minute to about six weeks). In another embodiment, the receiver determines sufficient stability based on oxygen concentration near the sensor head. In yet another embodiment, the sensor determines sufficient stability based on a reassessment of baseline drift and/or offset. A few examples of determining sufficient stability are given here, however a variety of known tests and parameters can be used to determine sensor stability without departing from the spirit and scope of the preferred embodiments.

If the receiver does not assess that the stability of the sensor is sufficient, then the processing returns to block **261**, wherein the receiver receives sensor data such as described in more detail above. The above-described steps are repeated until sufficient stability is determined.

If the receiver does assess that the stability of the sensor is sufficient, then processing continues to block **267** and the calibration set is used to calibrate the sensor.

At block **267**, the conversion function module uses the calibration set to create a conversion function. The conversion function substantially defines the relationship between the reference analyte data and the analyte sensor data.

A variety of known methods can be used with the preferred embodiments to create the conversion function from the calibration set. In one embodiment, wherein a plurality of matched data points form the initial calibration set, a linear least squares regression is performed on the initial calibration set such as described with reference to FIG. **14**A.

FIG. **22**A is a graph that illustrates a regression performed on a calibration set to create a conversion function in one exemplary embodiment. In this embodiment, a linear least squares regression is performed on the initial calibration set. The x-axis represents reference analyte data; the y-axis represents sensor data. The graph pictorially illustrates regression of the matched pairs **276** in the calibration set. Regression calculates a slope **272** and an offset **274** ($y=mx+b$), which defines the conversion function.

In alternative embodiments other algorithms could be used to determine the conversion function, for example forms of linear and non-linear regression, for example fuzzy logic, neural networks, piece-wise linear regression, polynomial fit, genetic algorithms, and other pattern recognition and signal estimation techniques.

In yet other alternative embodiments, the conversion function can comprise two or more different optimal conversions to account for optimal conversion variability due to dependence on parameters, such as time of day, calories consumed, exercise, or analyte concentration above or below a set threshold, for example. In one such exemplary embodiment, the conversion function is adapted for the estimated glucose concentration (e.g., high vs. low). For example, in an implantable glucose sensor, it has been observed that the cells surrounding the implant will consume at least a small amount of glucose as it diffuses toward the glucose sensor. Assuming the cells consume substantially the same amount of glucose whether the glucose concentration is low or high, this phenomenon will have a greater effect on the concentration of glucose during low blood sugar episodes than the effect on the concentration of glucose during relatively higher blood sugar episodes. Accordingly, the conversion function is adapted to compensate for the sensitivity differences in blood sugar level. In one implementation, the conversion function comprises two different regression lines wherein a first regression line is applied when the estimated blood glucose concentration is at or below a certain threshold (e.g., 150 mg/dL) and a second regression line is applied when the estimated blood glucose concentration is at or above a certain threshold (e.g., 150 mg/dL). In one alternative implementation, a predetermined pivot of the regression line that forms the conversion function can be applied when the estimated blood is above or below a set threshold (e.g., 150 mg/dL), wherein the pivot and threshold are determined from a retrospective analysis of the performance of a conversion function and its performance at a range of glucose concentrations. In another implementation, the regression line that forms the conversion function is pivoted about a point in order to comply with clinical acceptability standards (e.g., Clarke Error Grid, Consensus Grid, mean absolute relative difference, or other

clinical cost function). Although only a few example implementations are described, the preferred embodiments contemplate numerous implementations where the conversion function is adaptively applied based on one or more parameters that can affect the sensitivity of the sensor data over time.

In some alternative embodiments, the sensor is calibrated with a single-point through the use of a dual-electrode system to simplify sensor calibration. In one such dual-electrode system, a first electrode functions as a hydrogen peroxide sensor including a membrane system containing glucose-oxidase disposed thereon, which operates as described herein. A second electrode is a hydrogen peroxide sensor that is configured similar to the first electrode, but with a modified membrane system (with the enzyme domain removed, for example). This second electrode provides a signal composed mostly of the baseline signal, b.

In some dual-electrode systems, the baseline signal is (electronically or digitally) subtracted from the glucose signal to obtain a glucose signal substantially without baseline. Accordingly, calibration of the resultant difference signal can be performed by solving the equation y=mx with a single paired measurement. Calibration of the implanted sensor in this alternative embodiment can be made less dependent on the values/range of the paired measurements, less sensitive to error in manual blood glucose measurements, and can facilitate the sensor's use as a primary source of glucose information for the user. U.S. Publication No. US-2005-0143635-A1 describes systems and methods for subtracting the baseline from a sensor signal.

In some alternative dual-electrode system embodiments, the analyte sensor is configured to transmit signals obtained from each electrode separately (e.g., without subtraction of the baseline signal). In this way, the receiver can process these signals to determine additional information about the sensor and/or analyte concentration. For example, by comparing the signals from the first and second electrodes, changes in baseline and/or sensitivity can be detected and/or measured and used to update calibration (e.g., without the use of a reference analyte value). In one such example, by monitoring the corresponding first and second signals over time, an amount of signal contributed by baseline can be measured. In another such example, by comparing fluctuations in the correlating signals over time, changes in sensitivity can be detected and/or measured.

In some alternative embodiments, a regression equation y=mx+b is used to calculate the conversion function; however, prior information can be provided for m and/or b, thereby enabling calibration to occur with fewer paired measurements. In one calibration technique, prior information (e.g., obtained from in vivo or in vitro tests) determines a sensitivity of the sensor and/or the baseline signal of the sensor by analyzing sensor data from measurements taken by the sensor (e.g., prior to inserting the sensor). For example, if there exists a predictive relationship between in vitro sensor parameters and in vivo parameters, then this information can be used by the calibration procedure. For example, if a predictive relationship exists between in vitro sensitivity and in vivo sensitivity, $m \approx f(m_{in\ vitro})$, then the predicted m can be used, along with a single matched pair, to solve for b (b=y−mx). If, in addition, b can be assumed to be 0, for example with a dual-electrode configuration that enables subtraction of the baseline from the signal such as described above, then both m and b are known a priori, matched pairs are not needed for calibration, and the sensor can be completely calibrated e.g. without the need for reference analyte values (e.g. values obtained after implantation in vivo.)

In another alternative embodiment, prior information can be provided to guide or validate the baseline (b) and/or sensitivity (m) determined from the regression analysis. In this embodiment, boundaries can be set for the regression line that defines the conversion function such that working sensors are calibrated accurately and easily (with two points), and non-working sensors are prevented from being calibrated. If the boundaries are drawn too tightly, a working sensor may not enter into calibration. Likewise, if the boundaries are drawn too loosely, the scheme can result in inaccurate calibration or can permit non-working sensors to enter into calibration. For example, subsequent to performing regression, the resulting slope and/or baseline are tested to determine whether they fall within a predetermined acceptable threshold (boundaries). These predetermined acceptable boundaries can be obtained from in vivo or in vitro tests (e.g., by a retrospective analysis of sensor sensitivities and/or baselines collected from a set of sensors/patients, assuming that the set is representative of future data).

If the slope and/or baseline fall within the predetermined acceptable boundaries, then the regression is considered acceptable. Alternatively, if the slope and/or baseline fall outside the predetermined acceptable boundaries, steps can be taken to either correct the regression or fail-safe such that a system will not process or display errant data. This can be useful in situations wherein regression results in errant slope or baseline values. For example, when points (matched pairs) used for regression are too close in value, the resulting regression statistically is less accurate than when the values are spread farther apart. As another example, a sensor that is not properly deployed or is damaged during deployment can yield a skewed or errant baseline signal.

FIG. **22**B is a graph that illustrates one example of using prior information for slope and baseline. The x-axis represents reference glucose data (blood glucose) from a reference glucose source in mg/dL; the y-axis represents sensor data from a transcutaneous glucose sensor of the preferred embodiments in counts. An upper boundary line **215** is a regression line that represents an upper boundary of "acceptability" in this example; the lower boundary line **216** is a regression line that represents a lower boundary of "acceptability" in this example. The boundary lines **215**, **216** were obtained from retrospective analysis of in vivo sensitivities and baselines of glucose sensors as described in the preferred embodiments.

A plurality of matched data pairs **217** represent data pairs in a calibration set obtained from a glucose sensor as described in the preferred embodiments. The matched data pairs are plotted according to their sensor data and time-corresponding reference glucose data. A regression line **218** represents the result of regressing the matched data pairs **217** using least squares regression. In this example, the regression line falls within the upper and lower boundaries **215**, **216** indicating that the sensor calibration is acceptable.

However, if the slope and/or baseline had fallen outside the predetermined acceptable boundaries, which would be illustrated in this graph by a line that crosses the upper and/or lower boundaries **215**, **216**, then the system may be configured to assume a preset baseline value and re-run the regression (or a modified version of the regression) with the assumed baseline, where the assumed baseline value is derived from in vivo or in vitro testing. Subsequently, the newly derived slope and baseline are again tested to deter-

mine whether they fall within the predetermined acceptable boundaries. The processing continues in response to the results of the boundary test in a similar fashion. In general, for a set of matched pairs (e.g., a calibration set), regression lines with higher slope (sensitivity) have a lower baseline while regression lines with lower slope (sensitivity) have a higher baseline. Accordingly, the step of assuming a baseline and testing against boundaries can be repeated using a variety of different assumed baselines based on the baseline, sensitivity, in vitro testing, and/or in vivo testing. For example, if a boundary test fails due to high sensitivity, then a higher baseline is assumed and the regression re-run and boundary-tested. It is preferred that after about two iterations of assuming a baseline and/or sensitivity and running a modified regression, the system assumes an error has occurred (if the resulting regression lines fall outside the boundaries) and fail-safe. The term "fail-safe" includes modifying the system processing and/or display of data responsive to a detected error to avoid reporting of inaccurate or clinically irrelevant analyte values.

In yet another alternative embodiment, information obtained prior to sensor insertion can be provided for selecting the baseline (b) and/or slope (m) determined from the regression analysis. For example, a distribution of slopes and/or baselines typically obtained may be determined (or estimated) from retrospective analysis of a sample set of implanted sensors. This prior distribution information may be used to enable the sensor system to select a preferred combination of slope and/or baseline to be used in calibration.

In some embodiments, when one or more matched pairs are obtained from a sensor implanted in a host (as described elsewhere herein), a plurality of possible calibration lines can be drawn dependent upon the method of utilizing these matched pairs to draw a calibration line. For example, where only a single matched pair is obtained, numerous calibration lines having various baselines and slopes may be drawn through the single matched pair. The single matched pair alone does not provide enough information to determine the best calibration line. Similarly, where multiple matched pairs are obtained that are clustered closely together, numerous calibration lines may be drawn that pass close to all of the matched pairs in the cluster (e.g., numerous calibration lines are similarly correlated with the matched pairs). In some circumstances, some of these multiple calibration lines may not represent the best calibration (e.g., due to error in the reference glucose, small distribution of pairs, only one matched pair, and the like). Accordingly, in one embodiment, prior information regarding the typical distribution of slopes and/or baselines may be used to select one of the multiple calibration lines.

In one embodiment, a baseline can be selected, during calibration, based on the prior distribution information described above (e.g., by selecting the most probable baseline from the distribution of baselines). In another embodiment, a slope can be selected, during calibration, based on the prior distribution information described above (e.g., by selecting the most probable slope from the distribution of slopes). In a preferred embodiment, both a baseline and slope are selected, during calibration, based on the prior distribution information described above. For example, in one embodiment, of the possible multiple calibration lines, the calibration line is chosen that has a slope and baseline closest to the maximum joint probability of both the slope and baseline distributions. In some embodiments, preference may be given to either the slope or baseline such that having either a slope or baseline closer to its most probable value is weighted more heavily.

In one embodiment, depicted in FIG. **22**C, the slope and baseline closest to the maximum joint probability may be determined geometrically (e.g., when both the slope and baseline have normal distributions). In FIG. **22**C, each slope and baseline determined (or estimated) from retrospective analysis is plotted as a matched pair **230** in a slope-baseline plot. The set of matched pairs **230** form a cloud distribution, with the center of highest density corresponding to the highest probability of slope and baseline combination. The set of possible calibration lines for the sensor to be calibrated (e.g., from regression of a single matched pair or set of closely spaced matched pairs) make up a line **232** in the slope-baseline plot. In one embodiment, the point on this line **234** closest to the center of highest matched pair **230** density may be chosen as providing the slope and baseline of the selected calibration line. Either the slope or the baseline probabilities may be given preference by scaling the cloud distribution along either the baseline or slope axes.

The use of prior distribution information may be extended to more complex cases. For example, in some embodiments, the desired calibration curve is not linear (e.g., it has three or more parameters). In such cases, the most probable set of parameters (e.g., based on the prior distributions) the still correlates with the calibration data may be used to define the calibration curve.

In these various embodiments, utilizing an additional electrode, prior information (e.g., in vitro or in vivo testing), signal processing, or other information for assisting in the calibration process can be used alone or in combination to reduce or eliminate the dependency of the calibration on reference analyte values obtained by the host.

Referring again to FIG. **21**, at block **268**, a sensor data transformation module uses the conversion function to transform sensor data into substantially real-time analyte value estimates, also referred to as calibrated data, as sensor data is continuously (or intermittently) received from the sensor. For example, in the embodiment of FIG. **22**A, the sensor data, which can be provided to the receiver in "counts," is translated to estimate analyte value(s) in mg/dL. In other words, the offset value at any given point in time can be subtracted from the raw value (e.g., in counts) and divided by the slope to obtain the estimated analyte value:

$$\text{mg/dL} = \frac{(rawvalue - \text{offset})}{\text{slope}}$$

In some alternative embodiments, the sensor and/or reference analyte values are stored in a database for retrospective analysis.

At block **269**, an output module provides output to the user via the user interface. The output is representative of the estimated analyte value, which is determined by converting the sensor data into a meaningful analyte value such as described in more detail with reference to block **268**, above. User output can be in the form of a numeric estimated analyte value, an indication of directional trend of analyte concentration, and/or a graphical representation of the estimated analyte data over a period of time, for example. Other representations of the estimated analyte values are also possible, for example audio and tactile. Accordingly, after initial calibration of the sensor, and possibly determination of stability of the sensor data, real-time continuous analyte

information can be displayed on the user interface so that the user can regularly and proactively care for his/her diabetic condition within the bounds set by his/her physician.

In alternative embodiments, the conversion function is used to predict analyte values at future points in time. These predicted values can be used to alert the user of upcoming hypoglycemic or hyperglycemic events. Additionally, predicted values can be used to compensate for the time lag (e.g., 15 minute time lag such as described elsewhere herein), so that an estimate analyte value displayed to the user represents the instant time, rather than a time delayed estimated value.

In some embodiments, the substantially real time estimated analyte value, a predicted future estimate analyte value, a rate of change, and/or a directional trend of the analyte concentration is used to control the administration of a constituent to the user, including an appropriate amount and time, in order to control an aspect of the user's biological system. One such example is a closed loop glucose sensor and insulin pump, wherein the analyte data (e.g., estimated glucose value, rate of change, and/or directional trend) from the glucose sensor is used to determine the amount of insulin, and time of administration, that can be given to a diabetic user to evade hyper- and hypoglycemic conditions.

The conventional analyte meters (e.g., self-monitored blood analyte tests) are known to have a +−20% error in analyte values. For example, gross errors in analyte readings are known to occur due to patient error in self-administration of the blood analyte test. In one such example, if the user has traces of sugar on his/her finger while obtaining a blood sample for a glucose concentration test, then the measured glucose value will likely be much higher than the actual glucose value in the blood. Additionally, it is known that self-monitored analyte tests (e.g., test strips) are occasionally subject to manufacturing error.

Another cause for error includes infrequency and time delay that can occur if a user does not self-test regularly, or if a user self-tests regularly but does not enter the reference value at the appropriate time or with the appropriate time stamp. Providing reference data using a substantially continuous sensor overcomes those errors associated with irregular self-tests. To ensure the validity of the sensor data, the receiver can be configured to evaluate the clinical acceptability of received reference analyte data prior to their acceptance as a valid reference value.

In one embodiment, the reference analyte data (and/or sensor analyte data) is evaluated with respect to substantially time corresponding sensor data (and/or substantially time corresponding reference analyte data) to determine the clinical acceptability of the reference analyte and/or sensor analyte data. Clinical acceptability considers a deviation between time corresponding glucose measurements (e.g., data from a glucose sensor and data from a reference glucose monitor) and the risk (e.g., to the decision making of a diabetic patient) associated with that deviation based on the glucose value indicated by the sensor and/or reference data. Evaluating the clinical acceptability of reference and sensor analyte data, and controlling the user interface dependent thereon, can minimize clinical risk.

In one embodiment, the receiver evaluates clinical acceptability each time reference data is obtained. In another embodiment, the receiver evaluates clinical acceptability after the initial calibration and stabilization of the sensor, such as described with reference to FIG. **21**, above. In some embodiments, the receiver evaluates clinical acceptability as an initial pre-screen of reference analyte data, for example after determining if the reference glucose measurement is from about 40 to about 400 mg/dL. In other embodiments, other methods of pre-screening data can be used, for example by determining if a reference analyte data value is physiologically feasible based on previous reference analyte data values (e.g., below a maximum rate of change).

After initial calibration such as described in more detail with reference to FIG. **21**, the sensor data receiving module **261** receives substantially continuous sensor data (e.g., a data stream) via a receiver and converts that data into estimated analyte values. As used herein, "substantially continuous" is broad enough to include a data stream of individual measurements taken at time intervals (e.g., time-spaced) of from fractions of a second up to, e.g., 1, 2, or 5 minutes. As sensor data is continuously converted, it can be occasionally recalibrated such as described in more detail below with reference to FIG. **24**. Initial calibration and re-calibration of the sensor requires a reference analyte value. Accordingly, the receiver can receive reference analyte data at any time for appropriate processing. These reference analyte values can be evaluated for clinical acceptability such as described below as a fail-safe against reference analyte test errors.

In some embodiments, the reference data is pre-screened according to environmental and physiological issues, such as time of day, oxygen concentration, postural effects, and patient-entered environmental data. In one example embodiment, wherein the sensor comprises an implantable glucose sensor, an oxygen sensor within the glucose sensor is used to determine if sufficient oxygen is being provided to successfully complete the necessary enzyme and electrochemical reactions for glucose sensing. In another example embodiment wherein the sensor comprises an implantable glucose sensor, the counter electrode could be monitored for a "rail-effect", that is, when insufficient oxygen is provided at the counter electrode causing the counter electrode to reach operational (e.g., circuitry) limits.

In some embodiments, a clinical acceptability evaluation module in the receiver (not shown), also referred to as a clinical module, evaluates newly received reference data and/or time corresponding sensor data. In some embodiments of evaluating clinical acceptability, the rate of change of the reference data as compared to previous data is assessed for clinical acceptability. That is, the rate of change and acceleration (or deceleration) of many analytes has certain physiological limits within the body. Accordingly, a limit can be set to determine if the new matched pair is within a physiologically feasible range, indicated by a rate of change from the previous data that is within known physiological and/or statistical limits. Similarly, in some embodiments any algorithm that predicts a future value of an analyte can be used to predict and then compare an actual value to a time corresponding predicted value to determine if the actual value falls within a clinically acceptable range based on the predictive algorithm, for example.

In one exemplary embodiment, the clinical acceptability evaluation module matches the reference data with a substantially time corresponding converted sensor value such as described with reference to FIG. **21** above, and plots the matched data on a Clarke Error Grid such as described in more detail with reference to FIG. **23**.

FIG. **23** is a graph of two data pairs on a Clarke Error Grid to illustrate the evaluation of clinical acceptability in one exemplary embodiment. The Clarke Error Grid can be used by the clinical acceptability evaluation module to evaluate the clinical acceptability of the disparity between a reference glucose value and a sensor glucose (e.g., estimated glucose)

value, if any, in an embodiment wherein the sensor is a glucose sensor. The x-axis represents glucose reference glucose data and the y-axis represents estimated glucose sensor data. Matched data pairs are plotted accordingly to their reference and sensor values, respectively. In this embodiment, matched pairs that fall within the A and B regions of the Clarke Error Grid are considered clinically acceptable, while matched pairs that fall within the C, D, and E regions of the Clarke Error Grid are not considered clinically acceptable. Particularly, FIG. **23** shows a first matched pair **1992** is shown which falls within the A region of the Clarke Error Grid, therefore is it considered clinically acceptable. A second matched pair **94** is shown which falls within the C region of the Clarke Error Grid, therefore it is not considered clinically acceptable. Further description of evaluating reference data for its clinical acceptability is included in U.S. Publication No. US-2005-0027180-A1.

Reference is now made to FIG. **24**, which is a flow chart that illustrates the process of evaluation of calibration data for best calibration based on inclusion criteria of matched data pairs in one embodiment.

Calibration of analyte sensors can be variable over time; that is, the conversion function suitable for one point in time may not be suitable for another point in time (e.g., hours, days, weeks, or months later). For example, in an embodiment wherein the analyte sensor is subcutaneously implantable, the maturation of tissue ingrowth over time can cause variability in the calibration of the analyte sensor. As another example, physiological changes in the user (e.g., metabolism, interfering blood constituents, lifestyle changes) can cause variability in the calibration of the sensor. Accordingly, a continuously updating calibration algorithm is disclosed that includes reforming the calibration set, and thus recalculating the conversion function, over time according to a set of inclusion criteria.

At block **101**, the reference data receiving module, also referred to as the reference input module, receives one or more new reference analyte values (e.g., data point) from the reference analyte source. In some embodiments, the reference analyte value can be pre-screened according to criteria such as described in more detail with reference to FIG. **21**, block **62**. In some embodiments, the reference analyte values can be evaluated for clinical acceptability such as described in more detail with reference to FIG. **23**.

At block **102**, the data matching module, also referred to as the processor module, forms one or more updated matched data pairs by matching new reference data to substantially time corresponding sensor data, such as described in more detail with reference to FIG. **21**, block **63**.

At block **103**, a calibration evaluation module evaluates the new matched pair(s) inclusion into the calibration set. In some embodiments, the receiver simply adds the updated matched data pair into the calibration set, displaces the oldest and/or least concordant matched pair from the calibration set, and proceeds to recalculate the conversion function accordingly (block **105**).

In some embodiments, the calibration evaluation includes evaluating only the new matched data pair. In some embodiments, the calibration evaluation includes evaluating all of the matched data pairs in the existing calibration set and including the new matched data pair; in such embodiments not only is the new matched data pair evaluated for inclusion (or exclusion), but additionally each of the data pairs in the calibration set are individually evaluated for inclusion (or exclusion). In some alternative embodiments, the calibration evaluation includes evaluating all possible combinations of matched data pairs from the existing calibration set and including the new matched data pair to determine which combination best meets the inclusion criteria. In some additional alternative embodiments, the calibration evaluation includes a combination of at least two of the above-described embodiments.

Inclusion criteria comprise one or more criteria that define a set of matched data pairs that form a substantially optimal calibration set. One inclusion criterion comprises ensuring the time stamp of the matched data pairs (that make up the calibration set) span at least a set time period (e.g., three hours). Another inclusion criterion comprises ensuring that the time stamps of the matched data pairs are not more than a set age (e.g., one week old). Another inclusion criterion ensures that the matched pairs of the calibration set have a substantially distributed amount of high and low raw sensor data, estimated sensor analyte values, and/or reference analyte values. Another criterion comprises ensuring all raw sensor data, estimated sensor analyte values, and/or reference analyte values are within a predetermined range (e.g., 40 to 400 mg/dL for glucose values). Another criterion comprises evaluating the rate of change of the analyte concentration (e.g., from sensor data) during the time stamp of the matched pair(s). For example, sensor and reference data obtained during the time when the analyte concentration is undergoing a slow rate of change can be less susceptible inaccuracies caused by time lag and other physiological and non-physiological effects. Another criterion comprises evaluating the congruence of respective sensor and reference data in each matched data pair; the matched pairs with the most congruence can be chosen. Another criterion comprises evaluating physiological changes (e.g., low oxygen due to a user's posture that can effect the function of a subcutaneously implantable analyte sensor, or other effects such as described with reference to FIG. **21**) to ascertain a likelihood of error in the sensor value. Valuation of calibration set criteria can comprise evaluating one, some, or all of the above described inclusion criteria. It is contemplated that additional embodiments can comprise additional inclusion criteria not explicitly described herein.

At block **104**, the evaluation of the calibration set determines whether to maintain the previously established calibration set, or if the calibration set can be updated (e.g., modified) with the new matched data pair. In some embodiments, the oldest matched data pair is simply displaced when a new matched data pair is included. A new calibration set can include not only the determination to include the new matched data pair, but in some embodiments, can also determine which of the previously matched data pairs can be displaced from the calibration set.

At block **105**, the conversion function module recreates the conversion function using the modified calibration set. The calculation of the conversion function is described in more detail with reference to FIG. **21**.

At block **106**, the sensor data transformation module converts sensor data to calibrated data using the updated conversion function. Conversion of raw sensor data into estimated analyte values is described in more detail with reference to FIG. **21**.

Reference is now made to FIG. **25**, which is a flow chart that illustrates the process of evaluating the quality of the calibration in one embodiment. The calibration quality can be evaluated by determining the statistical association of data that forms the calibration set, which determines the confidence associated with the conversion function used in calibration and conversion of raw sensor data into estimated analyte values.

In one embodiment calibration quality can be evaluated after initial or updated calculation of the conversion function such as described elsewhere herein. However, calibration quality can be performed at any time during the data processing.

At block **111**, a sensor data receiving module, also referred to as the sensor data module, receives the sensor data from the sensor such as described in more detail with reference to FIG. **21**.

At block **112**, a reference data receiving module, also referred to as the reference input module, receives reference data from a reference analyte source, such as a substantially continuous analyte monitor, and as described in more detail with reference to FIG. **21**.

At block **113**, the data matching module, also referred to as the processor module, matches received reference data with substantially time corresponding sensor data to provide one or more matched data pairs, such as described in more detail with reference to FIG. **21**.

At block **114**, the calibration set module, also referred to as the processor module, forms a calibration set from one or more matched data pairs such as described in more detail with reference to FIGS. **21** and **24**.

At block **115**, the conversion function module calculates a conversion function using the calibration set, such as described in more detail with reference to FIGS. **21** and **16**.

At block **116**, the sensor data transformation module continuously (or intermittently) converts received sensor data into estimated analyte values, also referred to as calibrated data, such as described in more detail with reference to FIGS. **21** and **24**.

At block **117**, a quality evaluation module evaluates the quality of the calibration. In one embodiment, the quality of the calibration is based on the association of the calibration set data using statistical analysis. Statistical analysis can comprise any known cost function such as linear regression, non-linear mapping/regression, rank (e.g., non-parametric) correlation, least mean square fit, mean absolute deviation (MAD), mean absolute relative difference, and the like. The result of the statistical analysis provides a measure of the association of data used in calibrating the system. A threshold of data association can be set to determine if sufficient quality is exhibited in a calibration set.

In another embodiment, the quality of the calibration is determined by evaluating the calibration set for clinical acceptability, such as described with reference to blocks **82** and **83** (e.g., Clarke Error Grid, Consensus Grid, or clinical acceptability test). As an example, the matched data pairs that form the calibration set can be plotted on a Clarke Error Grid, such that when all matched data pairs fall within the A and B regions of the Clarke Error Grid, then the calibration is determined to be clinically acceptable.

In yet another alternative embodiment, the quality of the calibration is determined based initially on the association of the calibration set data using statistical analysis, and then by evaluating the calibration set for clinical acceptability. If the calibration set fails the statistical and/or the clinical test, the processing returns to block **115** to recalculate the conversion function with a new (e.g., optimized) set of matched data pairs. In this embodiment, the processing loop (block **115** to block **117**) iterates until the quality evaluation module 1) determines clinical acceptability, 2) determines sufficient statistical data association, 3) determines both clinical acceptability and sufficient statistical data association, or 4) surpasses a threshold of iterations; after which the processing continues to block **118**.

FIGS. **26**A and **26**B illustrate one exemplary embodiment wherein the accuracy of the conversion function is determined by evaluating the correlation coefficient from linear regression of the calibration set that formed the conversion function. In this exemplary embodiment, a threshold (e.g., 0.79) is set for the R-value obtained from the correlation coefficient.

FIGS. **26**A and **26**B are graphs that illustrate an evaluation of the quality of calibration based on data association in one exemplary embodiment using a correlation coefficient. Particularly, FIGS. **26**A and **26**B pictorially illustrate the results of the linear least squares regression performed on a first and a second calibration set (FIGS. **26**A and **26**B, respectively). The x-axis represents reference analyte data; the y-axis represents sensor data. The graph pictorially illustrates regression that determines the conversion function.

The regression line (and thus the conversion function) formed by the regression of the first calibration set of FIG. **26**A is the same as the regression line (and thus the conversion function) formed by the regression of the second calibration set of FIG. **26**B. However, the correlation of the data in the calibration set to the regression line in FIG. **26**A is significantly different than the correlation of the data in the calibration set to the regression line in FIG. **26**A. In other words, there is a noticeably greater deviation of the data from the regression line in FIG. **26**B than the deviation of the data from the regression line in FIG. **26**A.

In order to quantify this difference in correlation, an R-value can be used to summarize the residuals (e.g., root mean square deviations) of the data when fitted to a straight line via least squares method, in this exemplary embodiment. R-value can be calculated according to the following equation:

$$R = \frac{\sum_i (x_i - \bar{x})(y_i - \bar{y})}{\sqrt{\sum_i (x_i - x)^2} \sqrt{\sum_i y_i - y)^2}}$$

In the above equation: i is an index (1 to n), x is a reference analyte value, y is a sensor analyte value, $\bar{x}$ is an average of 1/n reference analyte values, and $\bar{y}$ is an average of 1/n sensor analyte values.

In the exemplary calibration set shown in FIG. **26**A, the calculated R-value is about 0.99, which can also be expressed as the correlation coefficient of regression. Accordingly, the calibration exhibits sufficient data association (and thus insufficient quality) because it falls above the 0.79 threshold set in this exemplary embodiment.

In the exemplary calibration set shown in FIG. **26**B, the calculated R-value is about 0.77, which can also be expressed as the correlation coefficient of regression. Accordingly, the calibration exhibits insufficient data association (and thus insufficient quality) because it falls below the 0.79 threshold set in this exemplary embodiment.

Reference is again made to FIG. **25**, at block **118**, the interface control module, also referred to as the fail-safe module, controls the user interface based upon the quality of the calibration. If the calibration is exhibits sufficient quality, then the then the first reference analyte value is discarded, and the repeated reference analyte value is accepted the process continues, and the user interface can function as normal; that is providing output for the user such as described in more detail with reference to FIG. **21**. If

however the calibration is not deemed sufficient in quality, then fail-safe module begins the initial stages of fail-safe mode, which are described in more detail in U.S. Publication No. US-2005-0027463-A1.

Example I

As described above, benchtop testing of a sensor prior to insertion can be used to calibrate sensor data after the sensor is inserted. FIG. **27** shows in-vivo sensitivity (as calculated using retrospective calibration) vs. in-vitro sensitivity (collected prior to implant), of 16 short term sensors implanted in humans. Least-squares regression was used to define the predictive relationship between in-vitro and in-vivo sensitivities (e.g., $m_{in\ vivo}=2.99*m_{in\ vitro}+26.62$), as shown in FIG. **27**. This relationship was applied to a sensor used in the above-described experiment. The comparison of blood glucose values with the in-vitro calibrated sensor is shown in FIGS. **28** and **29**. Predicted in-vivo sensitivity was assumed for calibration throughout the first day. The mean absolute relative difference found across 43 matched pairs over 3 days was 20.17%. In-vivo sensitivity is usually greater than in-vitro sensitivity (typically 2-3× higher). This may be due to enhanced transport of glucose when the membranes interact with molecules in the tissues, such as proteins or lipid in the interstitial fluid.

Example II

In this example, one substantially continuous glucose sensor was used to calibrate another substantially continuous glucose sensor, where both sensors were used in the same host. The data specifically illustrates the use of a short-term (e.g., transcutaneous) sensor to calibrate a long term (e.g., implantable) sensor, but the methods described are applicable to other combinations, including (1) use of a short-term sensor to calibrate another short term sensor; (2) use of a long-term sensor to calibrate another long-term sensor; (3) use of a long term sensor to calibrate a short term sensor.

As discussed herein, calibration of an enzyme-electrode based glucose sensor consists of solving the line

$$y=mx+b$$

for m and b, where y denotes the sensor signal (in units of A/D counts), x the estimated glucose concentration (mg/dl), m the sensor sensitivity to glucose (counts/mg/dl), and b the baseline signal unrelated to glucose (counts). If two glucose sensors are used concurrently in the same host, and one sensor is calibrated, the time-corresponding glucose values (x) from the calibrated sensor can be matched to the signal values (y) of the un-calibrated sensor. These matched pairs are then used to draw the calibration line for the second sensor using a regression relationship (e.g. ordinary least-squares).

In this example, a short term substantially continuous sensor was employed in a host in which a long term sensor was already implanted. The data from the substantially continuous short term sensor was used to run a prospective calibration on the substantially continuous long term sensor. Referring to FIG. **30**, a plurality of data points (e.g., about 20) from the short term sensor (referred to in FIG. **30** as "STS" and depicted with a diamond symbol) were used as reference data to calibrate the sensor data from the long term sensor (referred to in FIG. **30** as "LTS" and depicted with a dot symbol). The top graph in FIG. **30** shows sensor data (counts (left y-axis)) from an un-calibrated long term sensor and sensor data (glucose measurements (right y-axis)) from a calibrated short term sensor. The top graph in FIG. **30** shows roughly 6 days of un-calibrated long term sensor data for a particular host. On day five, a calibrated short term sensor was employed on the same host. The lower graph in FIG. **30** shows a close-up of the overlay of the sensor data from the long term sensor and, beginning on day five, the sensor data from the short term sensor.

After matching time-corresponding sensor values from the short term sensor data and the long term sensor data, a regression analysis yielded a result for (m, b) of (134.24, 11922.53), with 5 min time-lag of the long term sensor relative to short term sensor. When this calibration was prospectively applied to the long term sensor on day 6, the (calibrated) long term sensor glucose values were compared to reference glucose measurements (e.g. meter readings) to demonstrate calibration accuracy, as shown in FIG. **23**. In FIG. **31**, the long term sensor data is referred to as "LTS" and depicted as small dots, and the reference glucose measurements (self-monitoring blood glucose) are referred to as "SHBG" and depicted with large ovals. The results show the accuracy of the short term sensor calibration. Further improvement in accuracy would likely result if more points were used.

Comparison of the proposed calibration scheme (long term sensor calibration based on short term sensor) vs. a conventional scheme, for example, a 6-pt moving window regression, using 2/day self-monitoring blood glucose (SMBG) is shown in the Table 1.

TABLE 1

Calibration long term sensor calibration schemes: retrospective, with SMBG, and with short term sensor.

Calibration Scheme	Retrospective	With SMBG (conventional)	With STS
m	127.12	197.38	134.24
b	12196.48	13508.15	11922.53

The results suggest that calibration of a long term sensor with a short term sensor can be more accurate than with 2/day SMBG. The m and b found with short term sensor based calibration are closer to the values found with retrospective calibration ('gold standard') than with 2/day SMBG.

Example III

Referring now to FIG. **32**, in this example, sensor data from a short term analyte sensor was used to calibrate a subsequently employed short term sensor. Short term data was collected over a two and a half-day time period from a substantially continuous short term sensor employed on a human host. Finger stick glucose measurements (referred to in FIG. **32** as "Calpts" and depicted with large round dots) were used to calibrate Sensor No. 1. Glucose values (read off of the left y-axis) were derived from the Calibrated Sensor No. 1 sensor data. The Calibrated Sensor No. 1 data is referred to in FIG. **32** as "Calglucose 1" and depicted as small dots.

On Day 3, a second substantially continuous short term sensor (referred to in FIG. **32** as "Sensor2") was employed on the same host as Sensor No. 1, and sensor data was collected about every 5 minutes for one day (e.g., 288 data points). Digital count data (read off of the right y-axis) from the Sensor No. 2 is depicted in FIG. **32** as diamonds.

Using the glucose values resulting from the data of Sensor No. 1, the sensor data from Sensor No. 2 was calibrated using a retrospective calibration analysis, i.e., least squares regression correlation coefficient, and evaluated using a mean absolute relative difference (MARD). The results of the regression are illustrated in FIG. **33**, which shows 288 digital count data points from Sensor No. 2 matched to glucose values from Sensor No. 1 at corresponding times. The regression data was then prospectively applied to the data collected by Sensor No. 2 on Day 4. The Calibrated Sensor 2 data is depicted in FIG. **32** as medium-sized round dots and referred to in FIG. **32** as "Calglucose2." After one day, a finger stick glucose measurement was taken at about 12:00:00. The sensor data from FIG. **32** closely corresponded to the glucose measurement from the finger stick glucose measurement.

Methods and devices that are suitable for use in conjunction with aspects of the preferred embodiments are disclosed in U.S. Pat. Nos. 4,994,167; 4,757,022; 6,001,067; 6,741,877; 6,702,857; 6,558,321; 6,931,327; and 6,862,465.

Methods and devices that are suitable for use in conjunction with aspects of the preferred embodiments are disclosed in U.S. Publication No. US-2005-0176136-A1; U.S. Publication No. US-2005-0251083-A1; U.S. Publication No. US-2005-0143635-A1; U.S. Publication No. US-2005-0181012-A1; U.S. Publication No. US-2005-0177036-A1; U.S. Publication No. US-2005-0124873-A1; U.S. Publication No. US-2005-0051440-A1; U.S. Publication No. US-2005-0115832-A1; U.S. Publication No. US-2005-0245799-A1; U.S. Publication No. US-2005-0245795-A1; U.S. Publication No. US-2005-0242479-A1; U.S. Publication No. US-2005-0182451-A1; U.S. Publication No. US-2005-0056552-A1; U.S. Publication No. US-2005-0192557-A1; U.S. Publication No. US-2005-0154271-A1; U.S. Publication No. US-2004-0199059-A1; U.S. Publication No. US-2005-0054909-A1; U.S. Publication No. US-2005-0112169-A1; U.S. Publication No. US-2005-0051427-A1; U.S. Publication No. US-2003-0032874-A1; U.S. Publication No. US-2005-0103625-A1; U.S. Publication No. US-2005-0203360-A1; U.S. Publication No. US-2005-0090607-A1; U.S. Publication No. US-2005-0187720-A1; U.S. Publication No. US-2005-0161346-A1; U.S. Publication No. US-2006-0015020-A1; U.S. Publication No. US-2005-0043598-A1; U.S. Publication No. US-2003-0217966-A1; U.S. Publication No. US-2005-0033132-A1; U.S. Publication No. US-2005-0031689-A1; U.S. Publication No. US-2004-0045879-A1; U.S. Publication No. US-2004-0186362-A1; U.S. Publication No. US-2005-0027463-A1; U.S. Publication No. US-2005-0027181-A1; U.S. Publication No. US-2005-0027180-A1; U.S. Publication No. US-2006-0020187-A1; U.S. Publication No. US-2006-0036142-A1; U.S. Publication No. US-2006-0020192-A1; U.S. Publication No. US-2006-0036143-A1; U.S. Publication No. US-2006-0036140-A1; U.S. Publication No. US-2006-0019327-A1; U.S. Publication No. US-2006-0020186-A1; U.S. Publication No. US-2006-0020189-A1; U.S. Publication No. US-2006-0036139-A1; U.S. Publication No. US-2006-0020191-A1; U.S. Publication No. US-2006-0020188-A1; U.S. Publication No. US-2006-0036141-A1; U.S. Publication No. US-2006-0020190-A1; U.S. Publication No. US-2006-0036145-A1; U.S. Publication No. US-2006-0036144-A1; and U.S. Publication No. US-2006-0016700-A1.

Methods and devices that are suitable for use in conjunction with aspects of the preferred embodiments are disclosed in U.S. application Ser. No. 09/447,227 filed Nov. 22, 1999 and entitled "DEVICE AND METHOD FOR DETERMINING ANALYTE LEVELS"; U.S. application Ser. No. 11/280,672 filed Nov. 16, 2005, and entitled "TECHNIQUES TO IMPROVE POLYURETHANE MEMBRANES FOR IMPLANTABLE GLUCOSE SENSORS"; U.S. application Ser. No. 11/280,102 filed Nov. 16, 2005, and entitled "TECHNIQUES TO IMPROVE POLYURETHANE MEMBRANES FOR IMPLANTABLE GLUCOSE SENSORS"; U.S. application Ser. No. 11/201,445 filed Aug. 10, 2005 and entitled "SYSTEM AND METHODS FOR PROCESSING ANALYTE SENSOR DATA"; U.S. application Ser. No. 11/335,879 filed Jan. 18, 2006 and entitled "CELLULOSIC-BASED INTERFERENCE DOMAIN FOR AN ANALYTE SENSOR"; U.S. application Ser. No. 11/334,876 filed Jan. 18, 2006 and entitled "TRANSCUTANEOUS ANALYTE SENSOR"; U.S. application Ser. No. 11/333,837 filed Jan. 17, 2006 and entitled "LOW OXYGEN IN VIVO ANALYTE SENSOR".

All references cited herein, including but not limited to published and unpublished applications, patents, and literature references are incorporated herein by reference in their entirety and are hereby made a part of this specification. To the extent publications and patents or patent applications incorporated by reference contradict the disclosure contained in the specification, the specification is intended to supersede and/or take precedence over any such contradictory material.

The term "comprising" as used herein is synonymous with "including," "containing," or "characterized by," and is inclusive or open-ended and does not exclude additional, unrecited elements or method steps.

All numbers expressing quantities of ingredients, reaction conditions, and so forth used in the specification are to be understood as being modified in all instances by the term "about." Accordingly, unless indicated to the contrary, the numerical parameters set forth herein are approximations that may vary depending upon the desired properties sought to be obtained. At the very least, and not as an attempt to limit the application of the doctrine of equivalents to the scope of any claims in any application claiming priority to the present application, each numerical parameter should be construed in light of the number of significant digits and ordinary rounding approaches.

The above description discloses several methods and materials of the present invention. This invention is susceptible to modifications in the methods and materials, as well as alterations in the fabrication methods and equipment. Such modifications will become apparent to those skilled in the art from a consideration of this disclosure or practice of the invention disclosed herein. Consequently, it is not intended that this invention be limited to the specific embodiments disclosed herein, but that it cover all modifications and alternatives coming within the true scope and spirit of the invention.

What is claimed is:

1. A glucose monitoring system comprising:

a transcutaneous electrochemical glucose sensor comprising an in vivo portion and an ex vivo portion, wherein the transcutaneous electrochemical sensor is characterized by an in vivo sensitivity and an in vitro sensitivity, wherein the in vivo sensitivity is different from the in vitro sensitivity, and wherein the in vitro sensitivity is predictive of the in vivo sensitivity due to a relationship therebetween;

sensor electronics configured to be operably connected to the transcutaneous electrochemical glucose sensor;

a processor configured to be operably connected to the sensor electronics;

a memory configured to be operably connected to the processor, wherein the memory has stored therein information associated with the in vivo sensitivity of the transcutaneous electrochemical glucose sensor, wherein the information associated with the in vivo sensitivity is based at least in part on corrections for the difference between the in vivo sensitivity and the in vitro sensitivity of the transcutaneous electrochemical glucose sensor.

2. The glucose monitoring system of claim **1**, wherein the relationship between the in vivo sensitivity and the in vitro sensitivity is established empirically by performing a clinical study and comparing in vivo sensitivities with in vitro sensitivities of a plurality of transcutaneous glucose sensors.

3. The glucose monitoring system of claim **1**, wherein the in vivo sensitivity is greater than the in vitro sensitivity.

4. The glucose monitoring system of claim **1**, wherein the processor is configured to calibrate sensor data from the transcutaneous electrochemical glucose sensor using the information associated with in vivo sensitivity of the transcutaneous electrochemical glucose sensor without reliance on reference glucose concentration values obtained after insertion of at least a portion of the transcutaneous electrochemical glucose sensor in a host.

5. A method for calibrating data from a transcutaneous electrochemical glucose sensor, the method comprising:

receiving sensor data from the transcutaneous electrochemical glucose sensor after at least a portion of the transcutaneous electrochemical glucose sensor is inserted in a host, wherein the sensor data is indicative of a glucose concentration in the host; and

calibrating, using a processor, the sensor data received from the transcutaneous electrochemical glucose sensor, wherein the calibrating is based at least in part on information associated with an in vivo sensitivity of the transcutaneous electrochemical glucose sensor, wherein the information associated with the in vivo sensitivity is based at least in part on corrections for a difference between the in vivo sensitivity and an in vitro sensitivity for the transcutaneous electrochemical glucose sensor.

6. The method for calibrating data from a transcutaneous electrochemical glucose sensor of claim **5**, wherein the calibrating requires no user calibrations after the at least a portion of the transcutaneous electrochemical glucose sensor is inserted in the host.

7. The method for calibrating data from a transcutaneous electrochemical glucose sensor of claim **5**, wherein the difference between the in vivo sensitivity and the in vitro sensitivity for the transcutaneous electrochemical glucose sensor is determined prior to the insertion of the at least a portion of the transcutaneous electrochemical glucose sensor into the host.

8. The method for calibrating data from a transcutaneous electrochemical glucose sensor of claim **5**, wherein the in vitro sensitivity for the transcutaneous electrochemical glucose sensor is based at least in part on sampling a plurality of transcutaneous electrochemical glucose sensors prior to inserting the at least a portion of the transcutaneous electrochemical glucose sensor in the host.

9. The method for calibrating data from a transcutaneous electrochemical glucose sensor of claim **8**, wherein the in vitro sensitivity is calculated at least in part by examining a signal response of each of the plurality of transcutaneous electrochemical glucose sensors to a set of known glucose concentrations.

10. A method for providing calibrated glucose information based on sensor data from a transcutaneous electrochemical glucose sensor, the method comprising:

receiving, by a processor, post-implantation sensor data from the transcutaneous electrochemical glucose sensor, wherein the sensor data is indicative of a glucose concentration in a host;

receiving, by the processor, information associated with an in vivo sensitivity of the transcutaneous electrochemical glucose sensor, wherein the information associated with the in vivo sensitivity is based at least in part on corrections for a difference between the in vivo sensitivity and an in vitro sensitivity for the transcutaneous electrochemical glucose sensor;

generating calibrated sensor data, using the processor, by using the information associated with the in vivo sensitivity of the transcutaneous electrochemical glucose sensor to calibrate the sensor data received from the transcutaneous electrochemical glucose sensor; and

providing the calibrated sensor data to a user to facilitate monitoring a glucose level of the host.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE

CERTIFICATE OF CORRECTION

PATENT NO. : 10,709,364 B2
APPLICATION NO. : 16/690977
DATED : July 14, 2020
INVENTOR(S) : Apurv Ullas Kamath et al.

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

On the Title Page

On page 7, Column 1, item (56), U.S. Patent Documents, Line 19, delete "Roper" and insert --Roeper--.

On page 9, Column 2, item (56), U.S. Patent Documents, Line 59, delete "Duchonet" and insert --Duchon et--.

On page 22, Column 1, item (56), Other Publications, Line 12, delete "Actan," and insert --Acta,--.

On page 22, Column 1, item (56), Other Publications, Line 43, delete "Immoblized" and insert --Immobilized--.

On page 22, Column 2, item (56), Other Publications, Line 5, delete "Biosenors" and insert --Biosensors--.

On page 22, Column 2, item (56), Other Publications, Line 25, delete "Chemica" and insert --Chimica--.

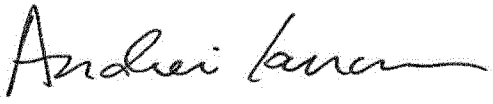
On page 22, Column 2, item (56), Other Publications, Line 42, delete "Chemica" and insert --Chimica--.

On page 23, Column 1, item (56), Other Publications, Line 43, delete "SurFACTS in Biomaterials," and insert --SurFACES in Biomaterials,--.

On page 23, Column 2, item (56), Other Publications, Line 57, delete "Biollogy" and insert --Biology--.

On page 23, Column 2, item (56), Other Publications, Line 66, delete "Self Monitoring" and insert --Self-Monitoring--.

Signed and Sealed this
Sixth Day of October, 2020



Andrei Iancu
Director of the United States Patent and Trademark Office

On page 24, Column 2, item (56), Other Publications, Line 1, delete "1-Butyrul-Glycerol:" and insert --1-Butyryl-Glycerol:--.

On page 24, Column 2, item (56), Other Publications, Line 22, delete "El-Sa'ad" and insert --El-Saad--.

On page 25, Column 1, item (56), Other Publications, Line 37, delete "Biosonsors" and insert --Biosensors--.

On page 25, Column 1, item (56), Other Publications, Line 45, delete "EnzymeTM" and insert --Enzyme™--.

On page 27, Column 1, item (56), Other Publications, Line 2, delete "519pages." and insert --519 pages.--.

On page 28, Column 1, item (56), Other Publications, Line 65, delete "Sensivity" and insert --Sensitivity--.

On page 30, Column 1, item (56), Other Publications, Line 35, delete "Coimmoblized" and insert --Coimmobilized--.

On page 30, Column 2, item (56), Other Publications, Line 68, delete ""Hydrogels" and insert --"Hydrogels--.

On page 30, Column 2, item (56), Other Publications, Line 69, delete "Eng," and insert --Eng.--.

On page 32, Column 1, item (56), Other Publications, Line 2, delete "Glucoseoxidase" and insert --Glucose oxidase--.

On page 32, Column 1, item (56), Other Publications, Line 50, delete "Moatti-Sirat.," and insert --Moatti-Sirat D.,--.

On page 32, Column 1, item (56), Other Publications, Line 66, delete "Biochimica" and insert --Biochimica et--.

On page 33, Column 2, item (56), Other Publications, Line 55, delete "datd" and insert --dated--.

On page 34, Column 1, item (56), Other Publications, Line 61, delete "Decarbozylase" and insert --Decarboxylase--.

On page 34, Column 2, item (56), Other Publications, Line 41, delete "Hypogylcemia" and insert --Hypoglycemia--.

On page 34, Column 2, item (56), Other Publications, Line 68, delete "Quinn C.P.," and insert --Quinn C.A.P.,--.

On page 35, Column 1, item (56), Other Publications, Line 38, delete ""Subcutaenous" and insert --"Subcutaneous--.

On page 36, Column 1, item (56), Other Publications, Line 52, delete "al.,"A" and insert --al., "A--.

On page 36, Column 1, item (56), Other Publications, Line 65, delete "Hypo-Hyperglycemia" and insert --Hypo/Hyperglycemia--.

On page 36, Column 2, item (56), Other Publications, Line 53, delete "J.Exp.Med," and insert --J. Exp. Med,--.

On page 37, Column 1, item (56), Other Publications, Line 35, delete "Gluco Watch®" and insert --GlucoWatch®--.

On page 37, Column 2, item (56), Other Publications, Line 22, delete "Tranducers," and insert --Transducers,--.

On page 37, Column 2, item (56), Other Publications, Line 64, delete "Dogs:" and insert --Dogs.--.

In the Drawings

In sheet 34 of 48, FIG. 22A, (Along Y-axis), Line 1, delete "SENSPR" and insert --SENSOR--.

In sheet 44 of 48, FIG. 29, Line 13, delete "dotx." and insert --dots.--.

In the Specification

Column 6, Line 16, delete "sensor" and insert --sensor.--.

Column 7, Line 57, delete "andrenostenedione;" and insert --androstenedione;--.

Column 7, Line 61, delete "hydroxy-cholic" and insert --hydroxycholic--.

Column 8, Line 2, delete "HCMV," and insert --HCMV, HIV-1,--.

Column 8, Lines 4-5, delete "deoxycorti sol);" and insert --deoxycortisol);--.

Column 8, Line 6, delete "diptheria/tetanus" and insert --diphtheria/tetanus--.

Column 8, Line 12, delete "perioxidase;" and insert --peroxidase;--.

Column 8, Lines 21-22, delete "sissomicin;" and insert --sisomicin;--.

Column 8, Line 26, delete "Giardia duodenalisa," and insert --Giardia duodenalis,--.

Column 8, Line 33, delete "Trepenoma pallidium," and insert --Treponema pallidum,--.

Column 8, Line 34, delete "stomatis" and insert --stomatitis--.

Column 8, Line 55, delete "(barbituates," and insert --(barbiturates,--.

Column 9, Line 4, delete "(FHIAA)." and insert --(5-HIAA).--.

Column 11, Line 22, delete "by product," and insert --byproduct,--.

Column 13, Line 41, delete "the a" and insert --the--.

Column 13, Line 49, delete "can not" and insert --cannot--.

Column 15, Line 56, delete "Sensory" and insert --SensorA--.

Column 16, Line 16, delete "(e.g.," and insert --e.g.,--.

Column 25, Line 25, delete "FIG." and insert --FIGS.--.

Column 28, Lines 55-56, delete "U.S. Pat. No. 6,424,847 to Mastrototaro et al., for example." and insert --for example.--.

Column 29, Line 2, delete "can by" and insert --can be--.

Column 29, Line 47, delete "Bellafonte, Pa." and insert --Bellefonte, PA.--.

Column 36, Lines 42-43, delete "methyl dopa," and insert --methyldopa,--.

Column 40, Line 37, delete "5:207-21 (1982))." and insert --5:207-21(1982)).--.

Column 42, Line 16, delete "interferant" and insert --interferent--.

Column 43, Line 29, delete "some time" and insert --sometime--.

Column 44, Line 52, delete "et." and insert --et--.

Column 44, Line 64, delete "a" and insert --α--.

Column 45, Lines 49-50, delete "New York))." and insert --New York.--.

Column 46, Line 29, delete "hydroxyapeptite," and insert --hydroxyapatite,--.

Column 46, Line 31, delete "nintinol," and insert --nitinol,--.

Column 50, Line 21, delete "and or" and insert --and/or--.

Column 57, Line 30, delete "pad" and insert --pad.--.

Column 61, Line 28, delete “and or” and insert --and/or--.

Column 70, Line 39, delete “U.S” and insert --U.S.--.

Column 71, Line 66, delete “mealtime,” and insert --meal time,--.

Column 82, Line 20, delete “scaling” and insert --sealing--.

Column 90, Line 21, delete ““SHBG”” and insert --“SMBG”--.