UNITED STATES PATENT AND TRADEMARK OFFICE

BEFORE THE PATENT TRIAL AND APPEAL BOARD

ALVOTECH USA INC. and ALVOTECH HF,
Petitioners,

v.

REGENERON PHARMACEUTICALS, INC.,
Patent Owner.

Case: PGR2025-00085
U.S. Patent No. 12,168,036

PATENT OWNER'S UPDATED LIST OF EXHIBITS
(as of June 19, 2026)

Exhibit	Description	Filed
2001	Excerpts from the File History of U.S. Appl. No. 17/384,070	X
2002	US 2018/0326126 ("Fiedler '126")	X
2003	Australian Dep't of Health & Human Aging, Australian Public Assessment Report for Aflibercept (July 2012) ("AusPAR")	X
2004	Ameya U. Borwankar et al., *Viscosity Reduction of a Concentrated Monoclonal Antibody with Arginine.HCl and Arginine.Glutamate*, 55 Ind. Eng. Chem. Rsch. 11225 (2016) ("Borwankar")	X
2005	Neal Whitaker et al., *A Formulation Development Approach to Identify and Select Stable Ultra–High-Concentration Monoclonal Antibody Formulations with Reduced Viscosities*, 106 J. Pharmaceutical Scis. 3230 (2017) ("Whitaker")	X
2006	Ameya U. Borwankar et al., *Viscosity Reduction of a Concentrated Monoclonal Antibody with Arginine.HCl and Arginine.Glutamate: Supplementary Information*, 55 Ind. Eng. Chem. Rsch. 11225 (2016) ("Borwankar SI")	X
2007	Eylea HD® Highlights of Prescribing Information (Oct. 2024)	X
2008	AbbVie, *Allergan, an AbbVie Company, and Molecular Partners Receive Complete Response Letter from FDA on Biologics License Application for Abicipar Pegol*, AbbVie News Center (June 26, 2020), https://news.abbvie.com/2020-06-26-Allergan-an-AbbVie-Company-and-Molecular-Partners-Receive-Complete-Response-Letter-from-FDA-on-Biologics-License-Application-for-Abicipar-pegol ("AbbVie Press Release")	X

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2009	Rod McNeil, *Advances and Developments in Medical Retina*, 25 Eye News 1 (2019) ("McNeil")	X
2010	Rehan M. Hussain & Thomas A. Ciulla, *Emerging Vascular Endothelial Growth Factor Antagonists to Treat Neovascular Age-Related Macular Degeneration*, 22 Expert Op. Emerging Drugs 235 (2017) ("Hussain")	X
2011	Angus Liu, *Novartis Calls off 3 Beovu Trials Testing More Frequent Dosing on Concerns of Vision-Threatening Side Effect*, FiercePharma (June 1, 2021) ("Liu 2021")	X
2012	Beovu Label (2024)	X
2013	Philip J. Rosenfeld & David J. Browning, *Editorial: Is This a 737 Max Moment for Brolucizumab?*, 216 Am. J. Ophthalmology A7 (2020) ("Rosenfeld")	X
2014	J. Gaudreault et al., *Preclinical Pharmacokinetics of Ranibizumab (rhuFabV2) After a Single Intravitreal Administration*, 46 Invest. Ophthalmol Vis. Sci. 726 (2005) ("Gaudreault")	X
2015	Public Trial Transcript Excerpts, *Regeneron Pharmaceuticals, Inc. v. Mylan Pharmaceuticals Inc.*, No. 22-cv-61 (N.D.W. Va.)	X
2016	Wang, *Protein Aggregation and Its Inhibition in Biopharmaceutics*, 289 Int. J. Pharm. 1 (2005) ("Wang 2005")	X
2017	Wang, *Instability, Stabilization, and Formulation of Liquid Protein Pharmaceuticals*, 1856 Int. J. Pharm. 129 (1999) ("Wang 1999")	X

Exhibit	Description	Filed
2018	Philip J. Rosenfeld et al., *Maximum Tolerated Dose of a Humanized Anti-Vascular Endothelial Growth Factor Antibody Fragment for Treating Neovascular Age-Related Macular Degeneration*, 112 Ophthalmology 1048 (2005) ("Rosenfeld 2005")	X
2019	U.S. Patent Application Publication No. 2004/0197324 ("Liu 2004")	X
2020	U.S. Patent No. 7,087,411 ("Daly")	X
2021	WO 2006/138181 ("Gokarn")	X
2022	Rahul N. Khurana et al., *Two-Year Results of the Phase 3 Randomized Controlled Study of Abicipar in Neovascular Age-Related Macular Degeneration*, 128 Ophthalmology 1027 (2021) ("Khurana 2021")	X
2023	James A. Dixon et al., *VEGF Trap-Eye for the Treatment of Neovascular Age-Related Macular Degeneration*, 18 Expert Opinion Investigational Drugs 1573 (2009) ("Dixon 2009")	X
2024	Pravin U. Dugel et al., *HAWK and HARRIER: Ninety-Six-Week Outcomes from the Phase 3 Trials of Brolucizumab for Neovascular Age-Related Macular Degeneration*, 128 Ophthalmology 89 (2021) ("Dugel")	X
2025	John S.Y. Park & David R. Chow, *Clinical Primer on Eylea HD: Review of the Current Data*, Retina Specialist (Apr. 13, 2024), https://www.retina-specialist.com/article/clinical-primer-on-eylea-hd-review-of-the-current-data ("Park")	X
2026	Vabysmo Label 2024 ("Vabysmo Label")	X

Exhibit	Description	Filed
2027	Jonas L. Fast et al., *Physical Instability of a Therapeutic Fc Fusion Protein: Domain Contributions to Conformational and Colloidal Stability*, 48 Biochemistry 11724 (2009) ("Fast")	X
2028	Zaltrap Label 2023 ("Zaltrap Label")	X
2029	Michael Thomas, *Comparative Effectiveness of Aflibercept for the Treatment of Patients with Neovascular Age-related Macular Degeneration*, 7 Clinical Ophthalmology 495 (2013) ("Thomas")	X
2030	Alvotech, Report of Foreign Private Issuer (Form 6-K) (June 23, 2025) ("Alvotech Press Release")	X
2031	Molecular Partners, Molecular Partners to Regain Global Rights to Abicipar, Molecular Partners News Release Details (Aug. 8, 2021) *https://investors.molecularpartners.com/news-releases/news-release-details/molecular-partners-regain-global-rights-abicipar* ("Molecular Partners Press Release")	X
2032	Novartis, *Novartis Reports One Year Results of Phase III MERLIN Study Evaluating Beovu® Every Four Week Dosing and Provides Update on Beovu Clinical Program*, Novartis Media Releases (May 28, 2021) https://www.novartis.com/news/media-releases/novartis-reports-one-year-results-phase-iii-merlin-study-evaluating-beovu-every-four-week-dosing-and-provides-update-beovu-clinical-program ("Novartis Press Release")	X
2033	U.S. Patent Application Publication No. 2023/0416353	X
2034	WO 2014/009465	X
2035	U.S. Food & Drug Admin., Eylea HD Approval Letter (Aug. 18, 2018) ("Eylea HD FDA Approval Letter")	X

Exhibit	Description	Filed
2036	Gerard A. Rodrigues, *Functional Characterization of Abicipar-Pegol, an Anti-VEGF DARPin Therapeutic That Potently Inhibits Angiogenesis and Vascular Permeability*, 59 *Investigative Ophthalmology & Visual Science* 5836 (2018) ("Rodrigues 2018")	X
2037	Declaration of K. Kayali	X
2038	Petition for Post Grant Review of United States Patent No. 12,158,036, Paper No. 2, PGR2025-00085	X
2039	Robert J. Falconer, *Advances in Liquid Formulations of Parenteral Therapeutic Proteins*, 37 Biotechnology Advances 1 (2019) ("Falconer 2019")	X
2040	18th Annual Conference, Bioprocessing Network 2025, Conference Schedule ("Conference Schedule")	X
2041	D'Amico article, American Journal of Ophthalmology, August, 1985	X
2042	Talamo article, American Journal of Ophthalmology, December, 1985	X
2043	Oum article, Graefe's Archive for Clinical and Experimental Ophthalmology 1992	X
2044	Sakarya article, European Journal Ophthalmology 2010	X
2045	Macugen Highlights of Prescribing Information, Bates ALVOTECH 1048	X
2046	Deposition Transcript of Dr. Robert J. Falconer (May 27, 2026)	X
2047	U.S. Patent No. 7,396,664	X

Exhibit	Description	Filed
2048	U.S. Patent No. 7,354,579	X
2049	H. Abe, *Role of Histidine-Related Compounds as Intracellular Proton Buffering Constituents in Vertebrate Muscle*, 65 Biochemistry (Moscow) 757 (2000)	X
2050	U.S. Patent App. Pub. 2011/0027292	X
2051	Declaration of Bernhardt Trout (June 19, 2026)	X
2052	Declaration of Neil Bressler (June 19, 2026)	X
2053	Deposition Transcript of Dr. Donald J. D'Amico (June 2, 2026)	X
2054	Excerpts from File History of U.S. Patent Application No. 16/409,631	X
2055	Excerpts from the Oxford Dictionary of Biochemistry and Molecular Biology (2d ed. 2000)	X
2056	Francis A. Carey & Robert M. Giuliano, Organic Chemistry, Chapter 25 (8th ed. 2011)	X
2057	Excerpts from the Handbook of Pharmaceutical Excipients (4th ed. 2003) ("HPE")	X
2058	Aruffo, *Immunoglobulin Fusion Proteins*, in Antibody Fusion Proteins 221 (Chamow & Ashkenazi eds., 1999) ("Aruffo 1999")	X
2059	Avastin® Prescribing Information (2004) ("Avastin PI")	X
2060	Bogard et al., *Practical Considerations in the Production, Purification, and Formulation of Monoclonal Antibodies for Immunoscintigraphy and Immunotherapy*, 19 Seminars in Nuclear Med. 202 (1989) ("Bogard 1989")	X

Exhibit	Description	Filed
2061	Chang et al., *Practical Approaches to Protein Formulation Development*, in Rational Design of Stable Protein Formulations (J.F. Carpenter & M.C. Manning eds., 2002) ("Chang 2002")	X
2062	Chi et al., *Physical Stability of Proteins in Aqueous Solution: Mechanism and Driving Forces in Nonnative Protein Aggregation*, 20 Pharm. Res. 1325 (2003) ("Chi 2003")	X
2063	S. Duvvuri et al., *Drug Delivery to the Retina: Challenges and Opportunities*, 3 Expert Op. Biological Therapy 45 (2003) ("Duvvuri 2003")	X
2064	Eylea® Prescribing Information (2011) ("Eylea PI 2011")	X
2065	D. Ghate & H.F. Edelhauser, *Ocular Drug Delivery*, 3 Expert Op. Biological Therapy 275 (2006) ("Ghate 2006")	X
2066	Lucentis® Prescribing Information (2006) ("Lucentis PI 2006")	X
2067	Macugen® Prescribing Information (2004) ("Macugen PI 2004")	X
2068	R.P.A. Manzano et al., *Testing Intravitreal Toxicity of Bevacizumab (Avastin)*, 26 Retina 257 (2006) ("Manzano 2006")	X
2069	Moroney et al., *Aflibercept in Epithelial Ovarian Carcinoma*, 5 Future Oncol. 591 (2009) ("Moroney 2009")	X
2070	M. Nagpal et al., *A Comparative Debate on the Various Anti-Vascular Endothelial Growth Factor Drugs*, 55 Indian J. Ophthalmology 437 (2007) ("Nagpal")	X
2071	Int'l Patent Publication No. WO2005/102303 (2005) ("Peyman")	X

Exhibit	Description	Filed
2072	Shukla et al., *Downstream Processing of Fc-Fusion Proteins*, in Therapeutic Fc-Fusion Proteins 97 (Chamow et al. eds., 2014) ("Shukla 2014")	X
2073	Souillac, *Biophysical Characterization of Insoluble Aggregates of a Multi-Domain Protein: An Insight into the Role of the Various Domains*, 94 J. Pharm. Sci. 2066 (2005) ("Souillac 2005")	X
2074	C.A. Stein & D. Castanotto, *FDA-Approved Oligonucleotide Therapies in 2017*, 25 Molecular Therapy 1069 (2017) ("Stein")	X
2075	Wang et al., *Antibody Structure, Instability, and Formulation*, 96 J. Pharm. Sci. 1 (2007) ("Wang 2007")	X
2076	Winter & Milstein, *Man-made Antibodies*, 349 Nature 293 (1991) ("Winter 1991")	X
2077	Wurm et al., *Optimizing Production and Recovery of Immunoadhesins*, in Antibody Fusion Proteins 281 (Chamow & Ashkenazi eds., 1999) ("Wurm 1999")	X
2078	Liu et al., *Reversible Self-Association Increases the Viscosity of a Concentrated Monoclonal Antibody in Aqueous Solution*, 94 J. Pharm. Sci. 1928 (2005) ("Liu 2005")	X
2079	Shire et al., *Challenges in the Development of High Protein Concentration Formulations*, 93 J. Pharm. Sci. 1390 (2004) ("Shire 2004")	X
2080	Neeraj J. Agrawal et al., *Computational Tool for the Early Screening of Monoclonal Antibodies for Their Viscosities*, 8 mAbs 43 (2016) ("Agrawal")	X
2081	Merriam-Webster's Medical Desk Dictionary (2002) ("Merriam-Webster")	X

Exhibit	Description	Filed
2082	Halvor N. Christensen, *Proteins as Buffers*, Ann. 133 N.Y. Acad. Sci. 34 (1966) ("Christensen")	X
2083	Y. Nozaki & C. Tanford, *Examination of Titration Behavior*, 11 Methods in Enzymology 715 (1967) ("Nozaki")	X
2084	J. Wyman, *An Analysis of the Titration Data of Oxyhemoglobin of the Horse by a Thermal Method*, 127 J. Biol. Chem. 1 (1939) ("Wyman")	X
2085	Andrew A. Moshfeghi, *Safety of Intravitreal Anti-VEGF Agents*, Review of Ophthalmology (Nov. 11, 2014), https://www.reviewofophthalmology.com/article/safety-of-intravitreal-antivegf-agents	X
2086	Joseph J. Tseng et al., *Sustained Increased Intraocular Pressure Related to Intravitreal Antivascular Endothelial Growth Factor Therapy for Neovascular Age-Related Macular Degeneration*, 21 J. Glaucoma 241 (2012)	X
2087	Judy E. Kim et al., *Short-Term Intraocular Pressure Changes Immediately After Intravitreal Injections of Anti-Vascular Endothelial Growth Factor Agents*, 146 Am. J. Ophthalmol. 930 (2008).	X
2088	Evangelos S. Gragoudas et al., *Pegaptanib for Neovascular Age-Related Macular Degeneration*, 351 New Eng. J. Med. 2805 (2004)	X
2089	David M. Brown et al., *Primary Endpoint Results of a Phase II Study of Vascular Endothelial Growth Factor Trap-Eye in Wet Age-Related Macular Degeneration*, 118 Ophthalmology 1089 (2011)	X

Exhibit	Description	Filed
2090	Jeffrey S. Heier et al., *Intravitreal Aflibercept (VEGF Trap-Eye) in Wet Age-Related Macular Degeneration*, 119 Ophthalmology 2537 (2012)	X
2091	Christophe Le Tourneau et al., *Dose Escalation Methods in Phase I Cancer Clinical Trials*, 101 J. Nat'l Cancer Inst. 708 (2009)	X
2092	Jeffrey R. Sachs et al., *Optimal Dosing for Targeted Therapies in Oncology: Drug Development Cases Leading by Example*, 22 Clinical Cancer Rsch. 1318 (2016)	X
2093	RESERVED	
2094	Jiunn H. Lin, *Species Similarities and Differences in Pharmacokinetics*, 23 Drug Metabolism & Disposition 1008 (1995)	X
2095	Frank G. Holz et al., *Efficacy and Safety of Lampalizumab for Geographic Atrophy Due to Age-Related Macular Degeneration: Chroma and Spectri Phase 3 Randomized Clinical Trials*, 136 JAMA Ophthalmol. 666 (2018)	X
2096	Neil M. Bressler et al., *Intraocular Pressure Outcomes After Lampalizumab Injections in Patients with Geographic Atrophy*, 142 JAMA Ophthalmol. 772 (2024)	X
2097	LabChem Inc., Material Safety Data Sheet: Congo Red (CAS No. 573-58-0)	X
2098	Agar Scientific Ltd., Safety Data Sheet: Congo Red (C.I. 22120)	X
2099	Santa Cruz Biotechnology, Safety Data Sheet: Amaranth (SC-214532)	X

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2100	Eduardo B. Rodrigues et al., *Preclinical Investigation of the Retinal Biocompatibility of Six Novel Vital Dyes for Chromovitrectomy*, 29 Retina 497 (2009)	X
2101	Carolina Biological Supply Co., Safety Data Sheet: Crystal Violet (CAS No. 548-62-9)	X
2102	Ellen Kurek, *Conflicting Evidence over Stroke Risk from AMD Drug*, Pharmacy Times, available at https://www.pharmacytimes.com/view/conflicting-evidence-over-storke-risk-from-amd-drug (last visited June 12, 2026)	X
2103	Rajendra S. Apte et al., *Pegaptanib 1-Year Systemic Safety Results from a Safety-Pharmacokinetic Trial in Patients with Neovascular Age-Related Macular Degeneration*, 114 Ophthalmology 1702 (2007)	X
2104	Bevacizumab (Avastin®) Label 2017 (Avastin Label)	X
2105	James A. Hanley & Abby Lippman-Hand, *If Nothing Goes Wrong, Is Everything All Right? Interpreting Zero Numerators*, 249 JAMA 1743 (1983)	X
2106	Mehdi Modarreszadeh et al., *Two Different Doses of Intravitreal Bevacizumab for Treatment of Choroidal Neovascularization Associated with Age-Related Macular Degeneration*, 3 J. Ophthalmic & Vision Res. 102 (2008)	X
2107	Deepika Malik et al., *Safety Profiles of Anti-VEGF Drugs: Bevacizumab, Ranibizumab, Aflibercept and Ziv-Aflibercept on Human Retinal Pigment Epithelium Cells in Culture*, 98 Brit. J. Ophthalmol. i11 (2014)	X
2108	Robert L. Avery et al., *Systemic Pharmacokinetics and Pharmacodynamics of Intravitreal Aflibercept, Bevacizumab, and Ranibizumab*, 37 Retina 1847 (2017)	X

Exhibit	Description	Filed
2109	U.S. Food & Drug Admin., *Guidance for Industry: Estimating the Maximum Safe Starting Dose in Initial Clinical Trials for Therapeutics in Adult Healthy Volunteers* (July 2005)	X
2110	Brandon G. Busbee et al., *Twelve-Month Efficacy and Safety of 0.5 mg or 2.0 mg Ranibizumab in Patients with Subfoveal Neovascular Age-Related Macular Degeneration*, 120 Ophthalmology 1046 (2013)	X
2111	U.S. Food & Drug Admin., *Postmarketing Adverse Event Reporting Compliance Program*, https://www.fda.gov/drugs/surveillance-post-drug-approval-activities/postmarketing-adverse-event-reporting-compliance-program (last visited June 16, 2026)	X
2112	U.S. Food & Drug Admin., *Guidance for Industry: Nonclinical Studies for the Safety Evaluation of Pharmaceutical Excipients* (May 2005)	X
2113	Sophie Postel-Vinay et al., *Towards New Methods for the Determination of Dose Limiting Toxicities and the Assessment of the Recommended Dose for Further Studies of Molecularly Targeted Agents—Dose-Limiting Toxicity and Toxicity Assessment Recommendation Group for Early Trials of Targeted Therapies, an European Organisation for Research and Treatment of Cancer-Led Study*, 50 Eur. J. Cancer 2040 (2014)	X
2114	RESERVED	
2115	V. Burckbuchler et al., *Rheological and Syringeability Properties of Highly Concentrated Human Polyclonal Immunoglobulin Solutions*, 76 Eur. J. Pharmaceutics & Biopharmaceutics 351 (2010)	X

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2116	Francesco Cilurzo et al., *Injectability Evaluation: An Open Issue*, 12 AAPS PharmSciTech 604 (2011)	X
2117	*Regeneron Pharm., Inc. v. Mylan Pharm. Inc.*, 714 F. Supp. 3d 652 (N.D.W. Va. 2024) ("*Mylan*")	X
2118	Jeffrey S. Heier et al., *The 1-Year Results of CLEAR-IT 2, a Phase 2 Study of Vascular Endothelial Growth Factor Trap-Eye Dosed As-Needed After 12-Week Fixed Dosing*, 118 Ophthalmology 1098 (2011)	X
2119	RESERVED	
2120	Claudine E. Pang et al., *Association Between Needle Size, Postinjection Reflux, and Intraocular Pressure Spikes After Intravitreal Injections*, 35 Retina 1401 (2015)	X
2121	Vinicius S. De Stefano et al., *Experimental Investigation of Needles, Syringes and Techniques for Intravitreal Injections*, 39 Clinical & Experimental Ophthalmology 236 (2011)	X
2122	Jean-Pierre Hubschman et al., *Experimental Model of Intravitreal Injection Techniques*, 30 Retina 167 (2010)	X
2123	Hussein Hollands, et al., *Short-term Intraocular Pressure Changes after Intravitreal Injection of Bevacizumab*, 42 Can J. Ophthalmology 807 (2007)	X
2124	Daniel Y. Choi, *Sustained Elevated Intraocular Pressures After Intravitreal Injections of Bevacizumab, Ranibizumab, and Pegaptinib*, 31(6) Retina 1028 (2011)	X
2125	Third-Party Confidential Transcript	X
2126	Roche, *Angiogenesis Highlights* (Feb. 16, 2021)	X

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2127	U.S. Food & Drug Admin., *Inactive Ingredients in Approved Drug Products Search: Frequently Asked Questions*, https://www.fda.gov/drugs/drug-approvals-and-databases/inactive-ingredients-approved-drug-products-search-frequently-asked-questions (last visited June 17, 2026) ("Inactive Ingredient FAQ")	X
2128	Mohammed Gabr Kassem, *Sunitinib Malate*, 37 Profiles of Drug Substances, Excipients, and Related Methodology 363 (2021) ("Kassem")	X
2129	Xue-qing Wang, *Bioavailability and pharmacokinetics of sorafenib suspension, nanoparticles and nanomatrix for oral administration to rat*, 419 Int'l J. of Pharmaceutics 339 (2011) ("Wang 2011")	X
2130	*Clinical & Pharmacology & Biopharmaceutics Review(s), Application Number 22-465*, Center for Drug Evaluation & Research (2009) ("No. 22-465")	X
2131	Eylea HD® Highlights of Prescribing Information (Apr. 2026)	X
2132	U.S. Food & Drug Admin., *April 2018: Inactive Ingredient Database File*, downloaded from https://www.fda.gov/drugs/drug-approvals-and-databases/inactive-ingredients-database-download (last visited June 17, 2026) ("Inactive Ingredient Database")	X
2133	Excerpt from T.V. Chirila, et al., *The Vitreous Humor*, Handbook of Biological Properties, 125 (1998) ("Chirila")	X
2134	Mohab Eldeeb, et al., *One-Year Outcomes of Ziv-Aflibercept for Macular Edema in Central Retinal Vein Occlusion*, 8 Am. J. Ophthalmology Case Reports 58 (2017) ("Eldeeb")	X

Exhibit	Description	Filed
2135	Tetsuo Hida, *Experimental and Clinical Observations of the Intraocular Toxicity of Commercial Cortiosteroid Preparations*, 101 Am. J. Ophthalmology 190 (1986) ("Hida")	X
2136	J. García-Carmona, *Morphological Characteristics and Aggregation of Calcite Crystals Obtained by Bubbling CO_2 through a $Ca(OH)_2$ Suspension in the Presence of Additives*, 130 Powder Tech. 307 (2003) ("García-Carmona")	X
2137	E.N. Lee et al., *Stabilizing Peptide Fusion for Solving the Stability and Solubility Problems of Therapeutic Proteins*, 22 Pharm. Res. 1735, 1745 (2005) ("Lee")	X
2138	J.F. Carpenter et al., *Rational Design of Stable Lyophilized Protein Formulations: Theory and Practice*, in Rational Design of Stable Protein Formulations 109, 109 (J.F. Carpenter & M.C. Manning eds., 2002) ("Carpenter")	X
2139	Biochemistry: A Problems Approach, Chapter 2: Amino Acids (2d ed. 1981) ("Biochemistry")	X
2140	U. Schmidt-Erfurth et al., *Intravitreal Aflibercept Injection for Neovascular Age-related Macular Degeneration*, 121 Ophthalmology 193 (2014)	X
2141	Regeneron, Regeneron Announces Encouraging Topline Phase 2 Data of High-dose aflibercept in Wet Age-related Macular Degeneration, Regeneron Press Release Details, (Aug. 24, 2021), https://newsroom.regeneron.com/news-releases/news-release-details/regeneron-announces-encouraging-topline-phase-2-data-high-dose	X
2142	Marina Gühlike, *Taking Subvisible Particle Quantitation to the Limit*, 109 J. Pharm. Sci.505 (2020) ("Gühlike")	X

Exhibit	Description	Filed
2143	J.G. Gross, *Interim Safety Data Comparing Ranibizumab With Panretinal Photocoagulation Among Participants With Proliferative Diabetic Retinopathy*, 135 JAMA Ophthalmology 672 (2017)	X
2144	2026.06.09 Email from M. Chacon	X
2145	2025.09.17 Email from Dr. D'Amico	X

Dated: June 19, 2026

Respectfully submitted,

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CERTIFICATE OF SERVICE
(37 C.F.R. § 42.6(e))

The undersigned hereby certifies that on June 19, 2026, I caused true and correct copies of the foregoing to be served via e-mail on the following:

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