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Valneva, Ex. 1010

961713

THE UNITED STATES OF AMERICA

TO ALL TO WHOM THESE PRESENTS SHALL COME:

UNITED STATES DEPARTMENT OF COMMERCE

United States Patent and Trademark Office

November 21, 2018

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APPLICATION NUMBER: *62/581,500*
FILING DATE: *November 03, 2017*
RELATED PCT APPLICATION NUMBER: *PCT/US18/59219*

THE COUNTRY CODE AND NUMBER OF YOUR PRIORITY APPLICATION, TO BE USED FOR FILING ABROAD UNDER THE PARIS CONVENTION, IS *US62/581,500*

Certified by

Michelle K. Lee

Under Secretary of Commerce
for Intellectual Property
and Director of the United States
Patent and Trademark Office

FIG. 1

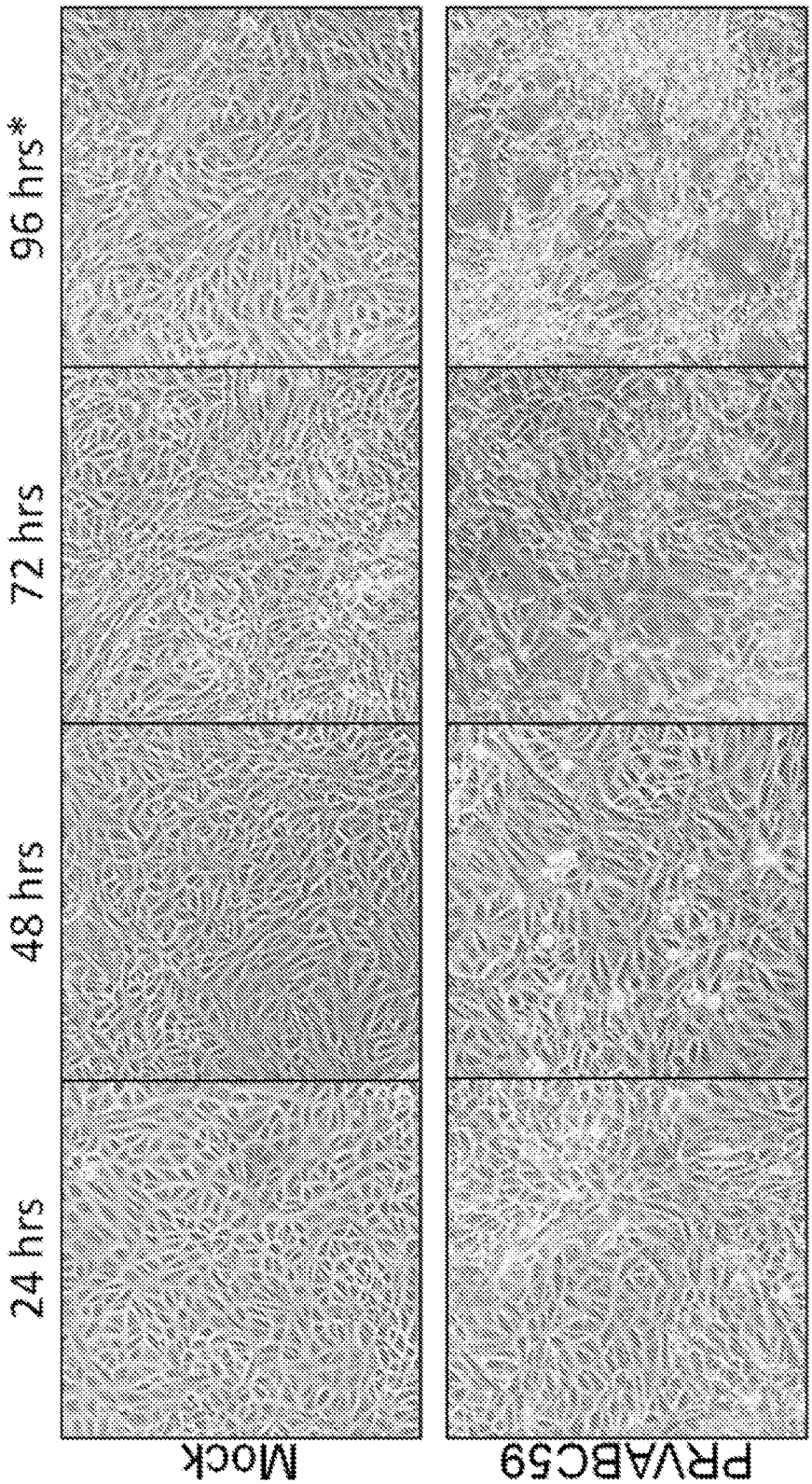


FIG. 2

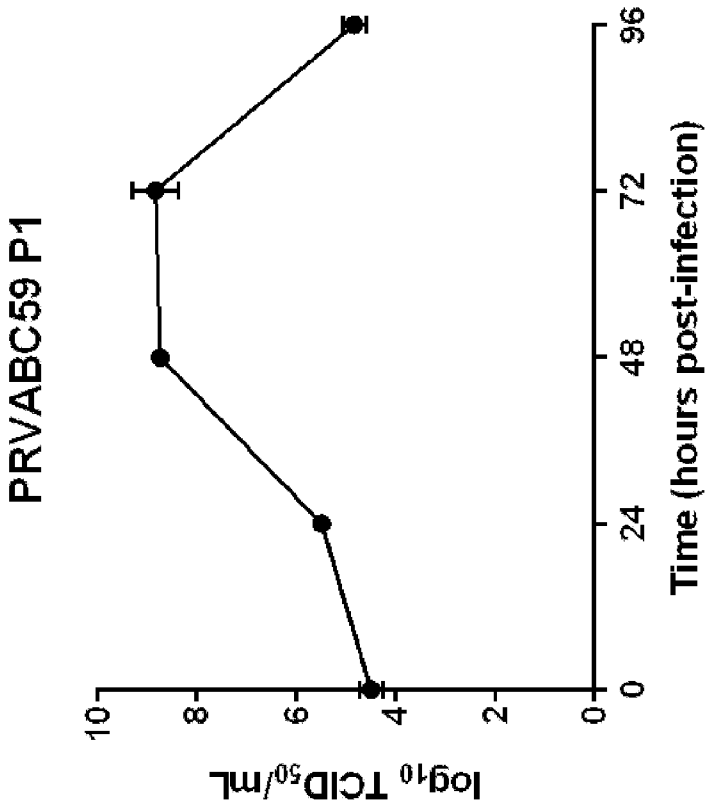

PRVABC59 P1
log10 TCID50/mL
10
8
6
4
2
0
0
24
48
72
96
Time (hours post-infection)

FIG. 3

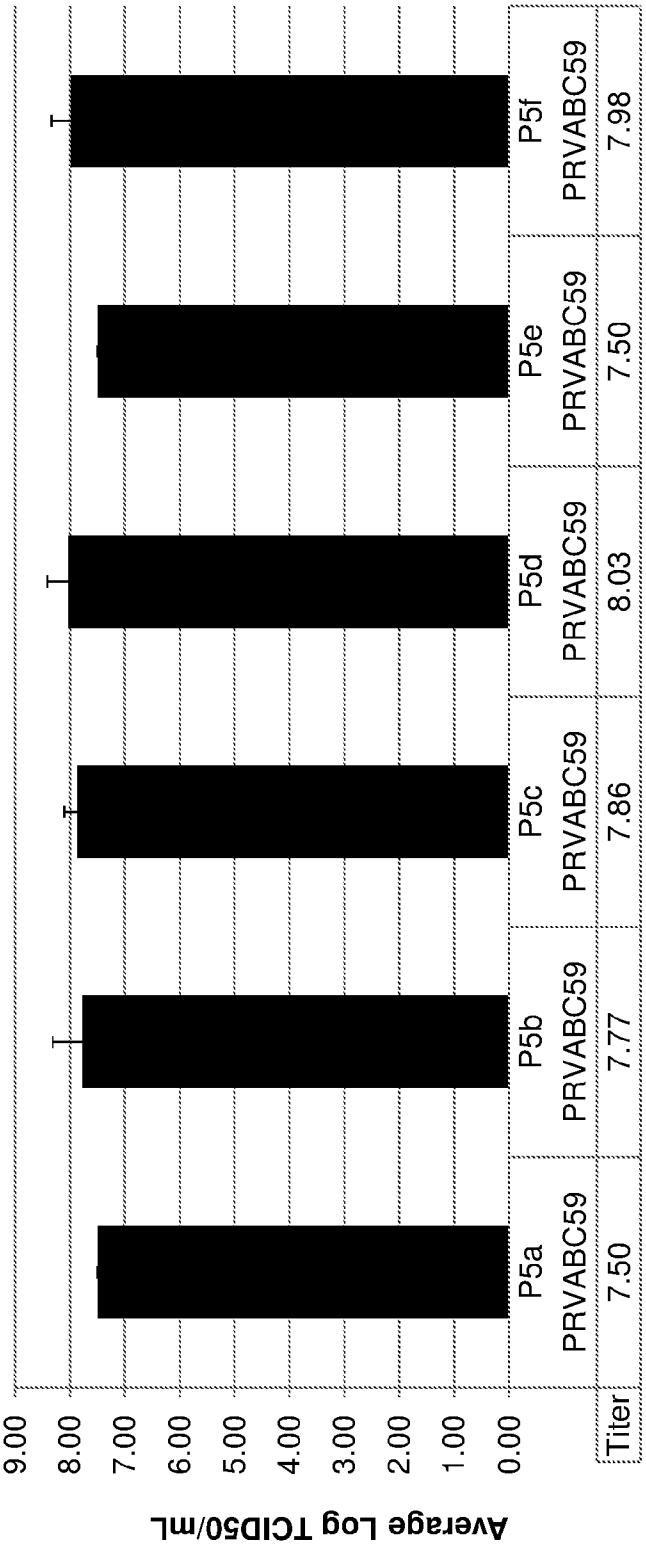

Average Log TCID50/mL
9.00
8.00
7.00
6.00
5.00
4.00
3.00
2.00
1.00
0.00
P5a PRVABC59
P5b PRVABC59
P5c PRVABC59
P5d PRVABC59
P5e PRVABC59
P5f PRVABC59
Titer
7.50
7.77
7.86
8.03
7.50
7.98

FIG. 4

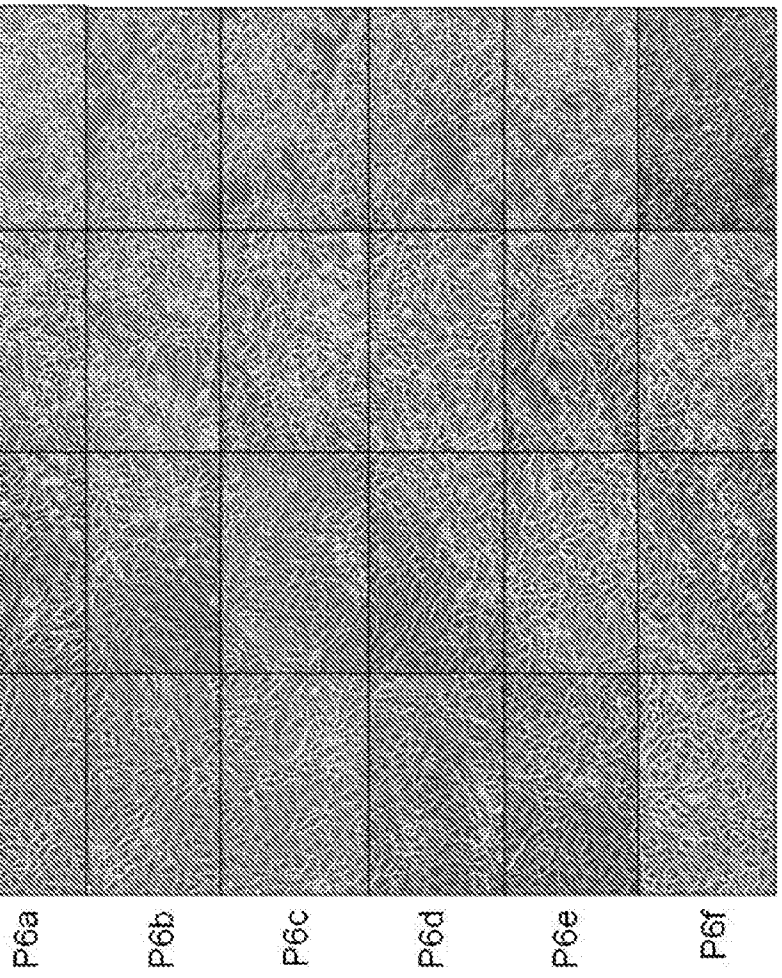

Harvest 1
Harvest 2
Day 1
Day 2
Day 3
Day 5
P6a
P6b
P6c
P6d
P6e
P6f

FIG. 5

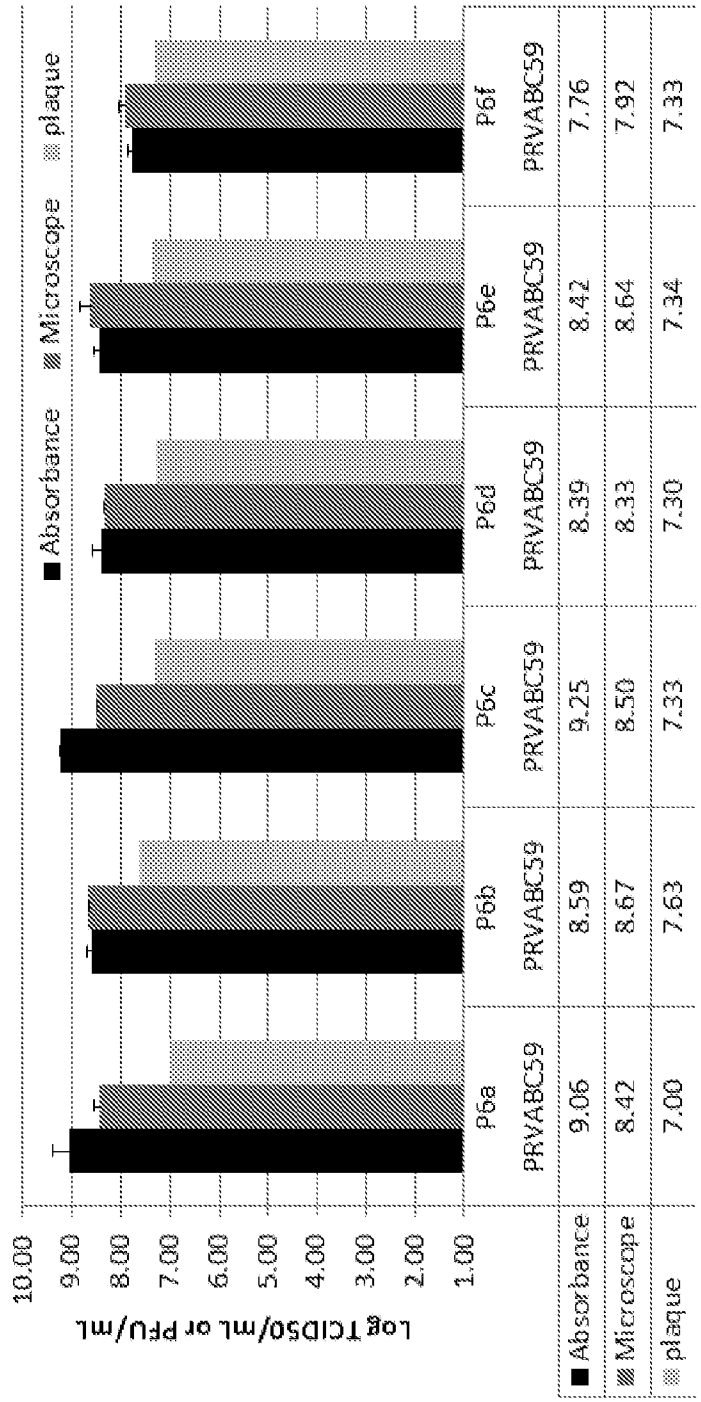

Absorbance
Microscope
plaque
Log TCID50/mL or PFU/mL
10.00
9.00
8.00
7.00
6.00
5.00
4.00
3.00
2.00
1.00
P6a PRVABC59
P6b PRVABC59
P6c PRVABC59
P6d PRVABC59
P6e PRVABC59
P6f PRVABC59
Absorbance 9.06 8.59 9.25 8.39 8.42 7.76
Microscope 8.42 8.67 8.50 8.33 8.64 7.92
plaque 7.00 7.63 7.33 7.30 7.34 7.33

FIG. 6

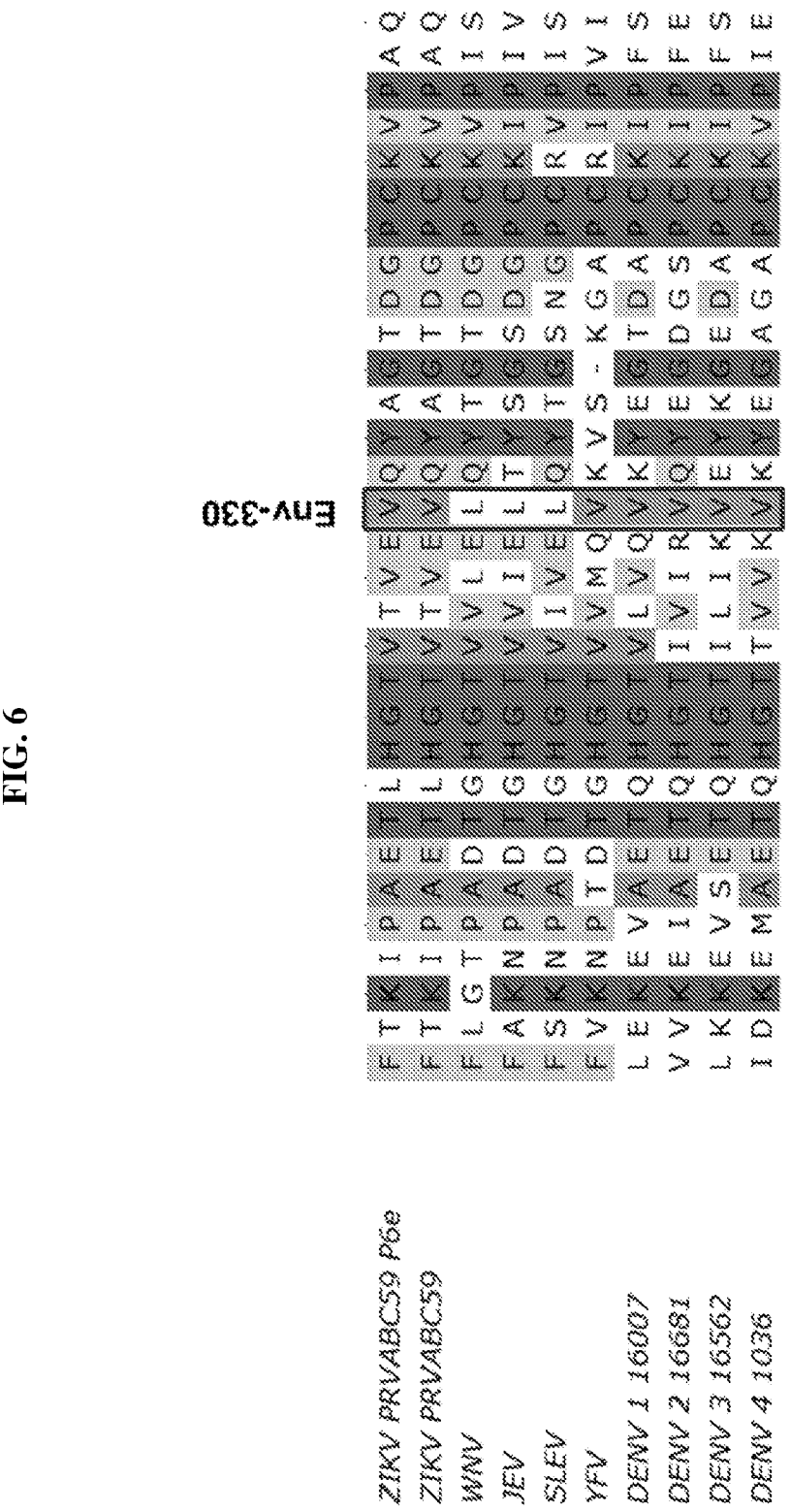

Env-330
ZIKV PRVABC59 P6e
ZIKV PRVABC59
WNV
JEV
SLEV
YFV
DENV 1 16007
DENV 2 16681
DENV 3 16562
DENV 4 1036

FIG. 7

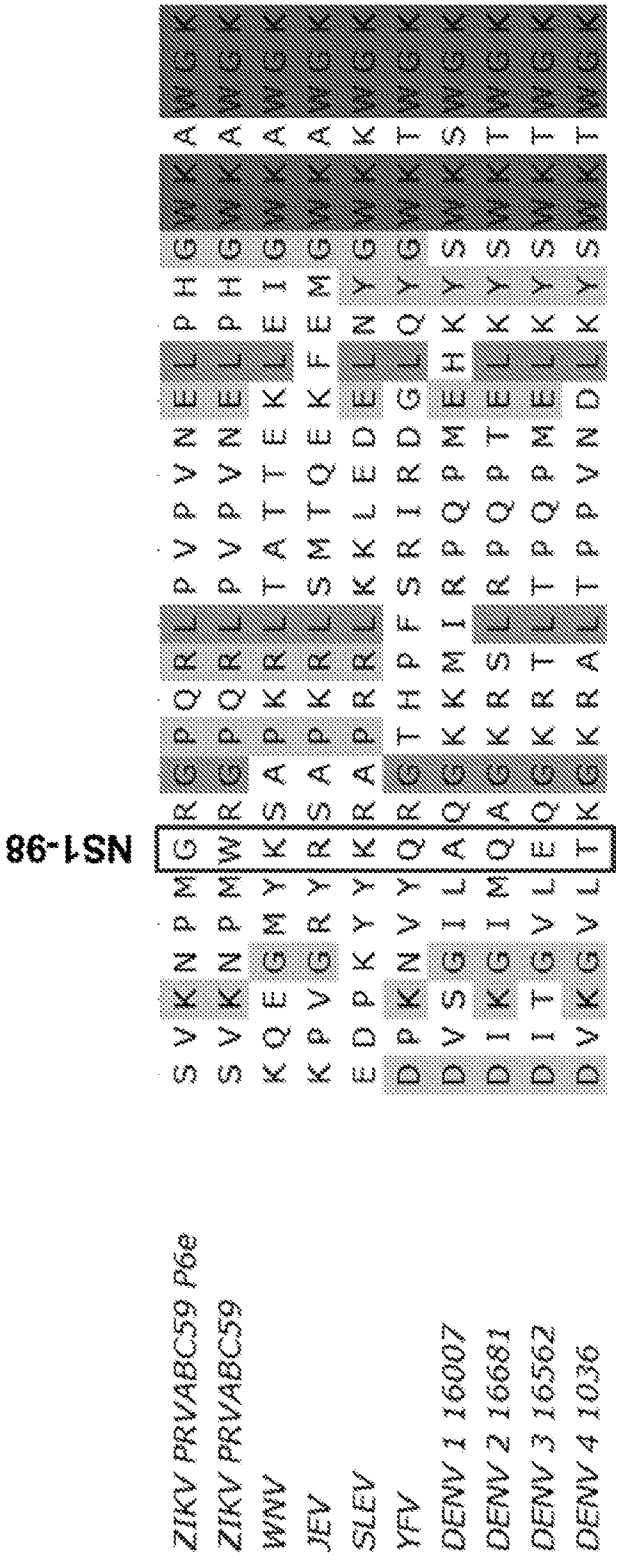

NS1-98
ZIKV PRVABC59 P6e
ZIKV PRVABC59
WNV
JEV
SLEV
YFV
DENV 1 16007
DENV 2 16681
DENV 3 16562
DENV 4 1036

FIG. 8

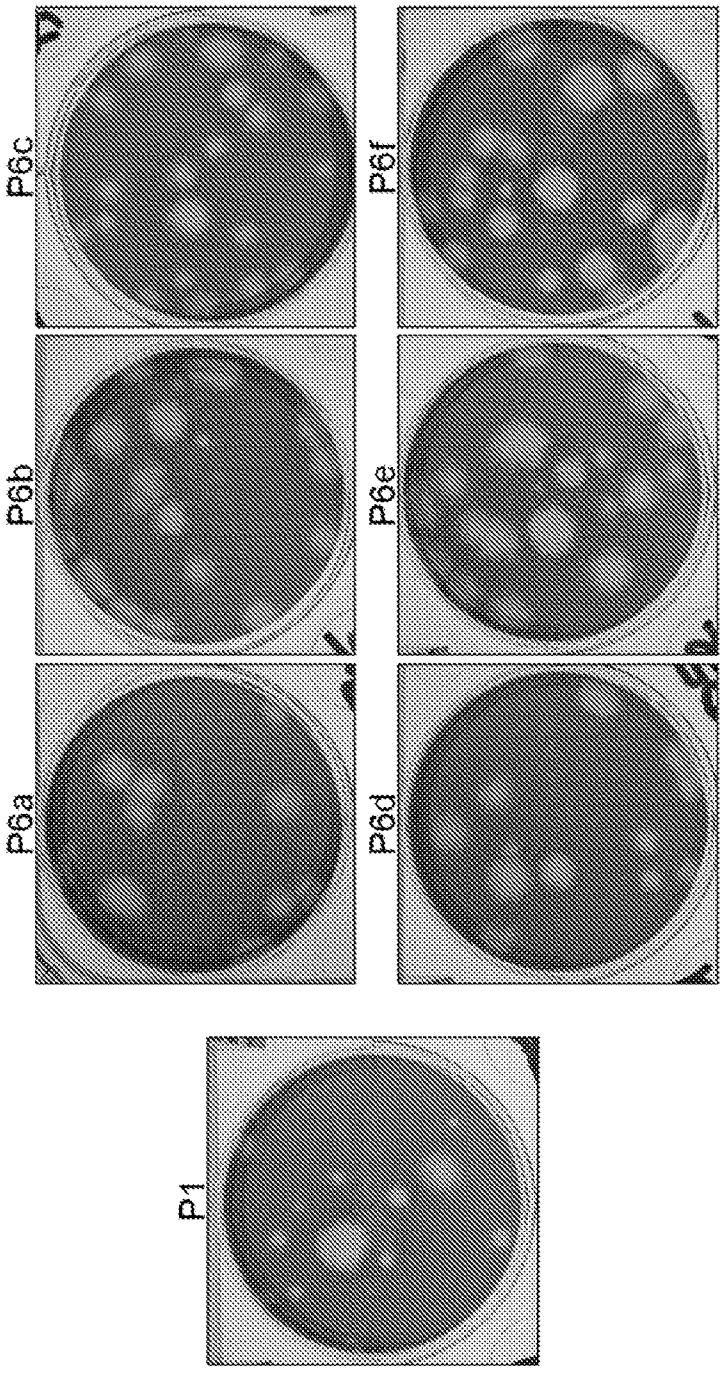


FIG. 9

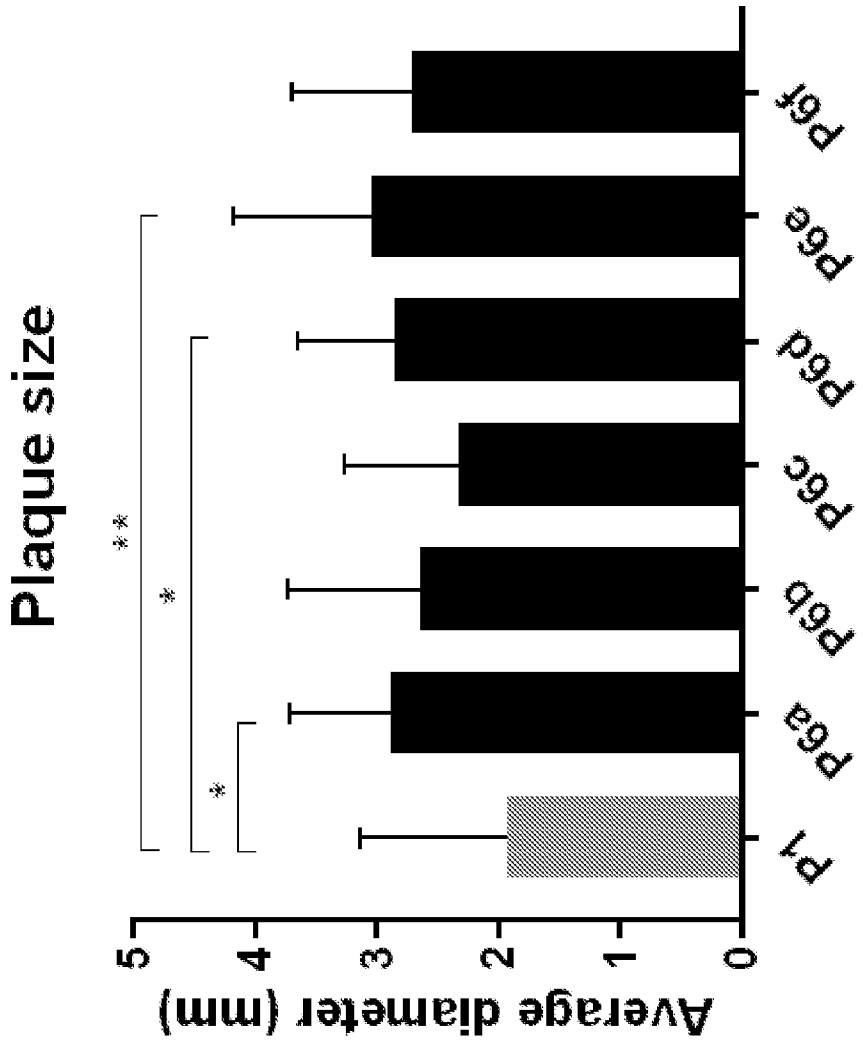

Plaque size
Average diameter (mm)
0
1
2
3
4
5
P1
P6a
P6b
P6c
P6d
P6e
P6f
*
*
**

FIG. 10

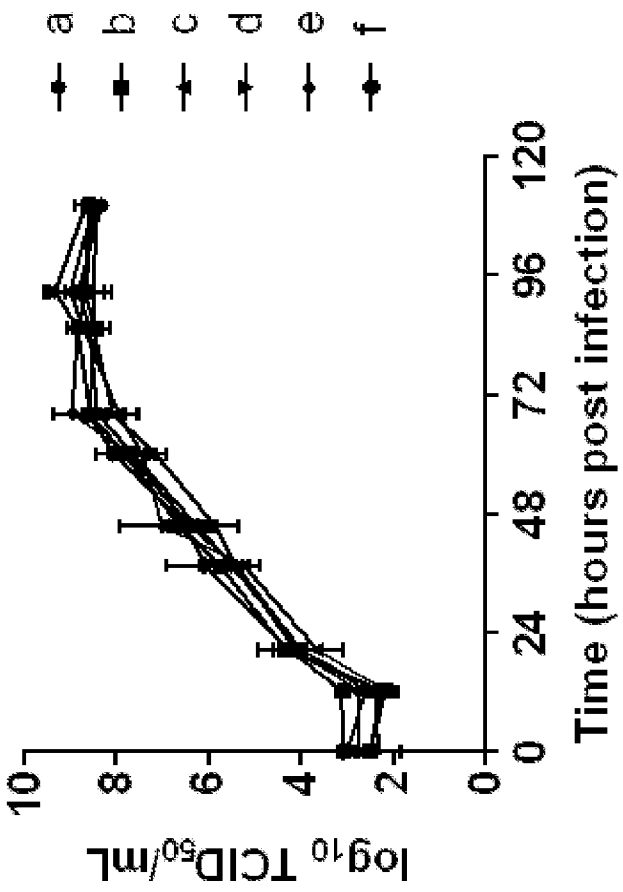

a
b
c
d
e
f
10
8
6
4
2
0
log10 TCID50/mL
0
24
48
72
96
120
Time (hours post infection)

FIG. 11

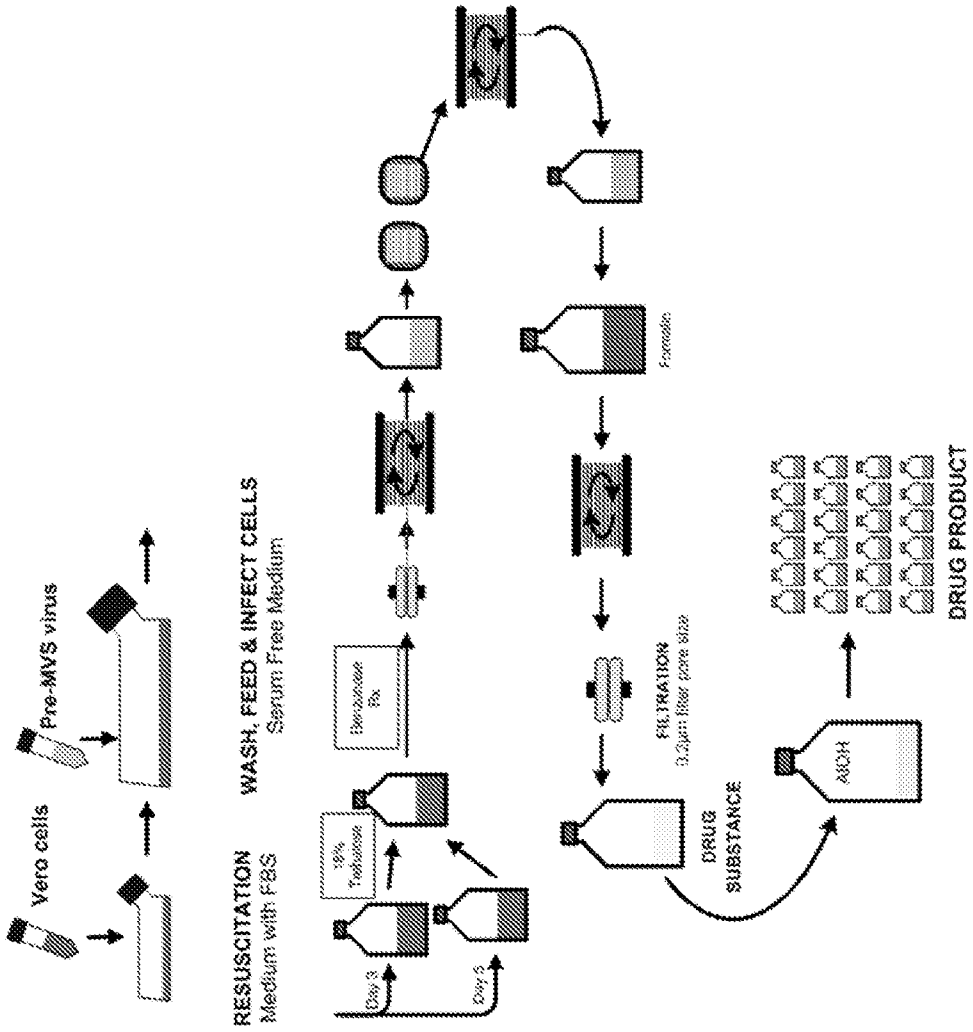

Vero cells
Pre-MVS virus
RESUSCITATION
Medium with FBS
WASH, FEED & INFECT CELLS
Serum Free Medium
Day 3
Day 5
FILTRATION
DRUG SUBSTANCE
DRUG PRODUCT

FIG. 12A



FIG. 12B

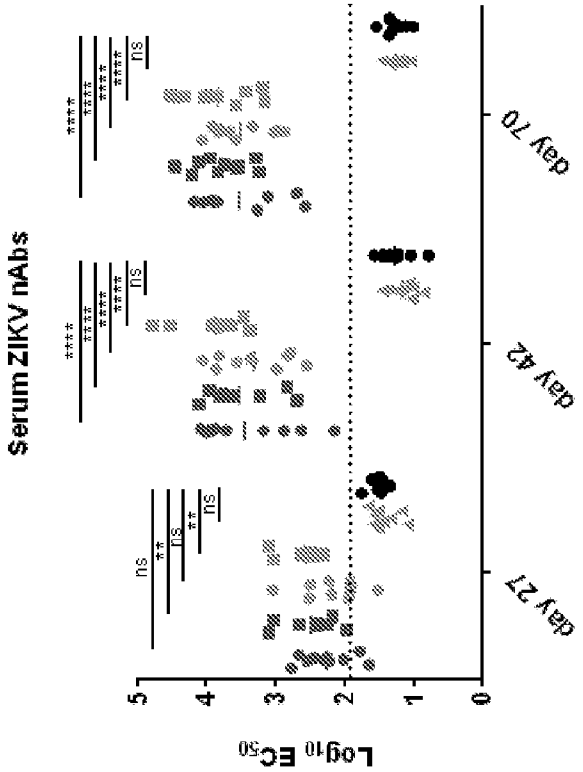


FIG. 13A

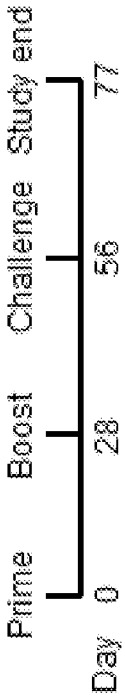


FIG. 13B

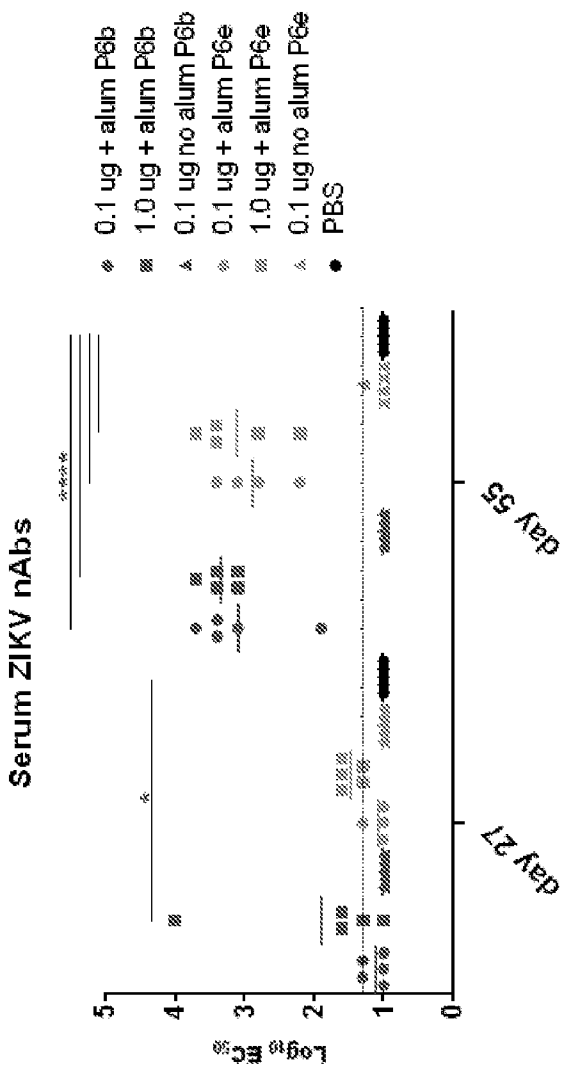


FIG. 14

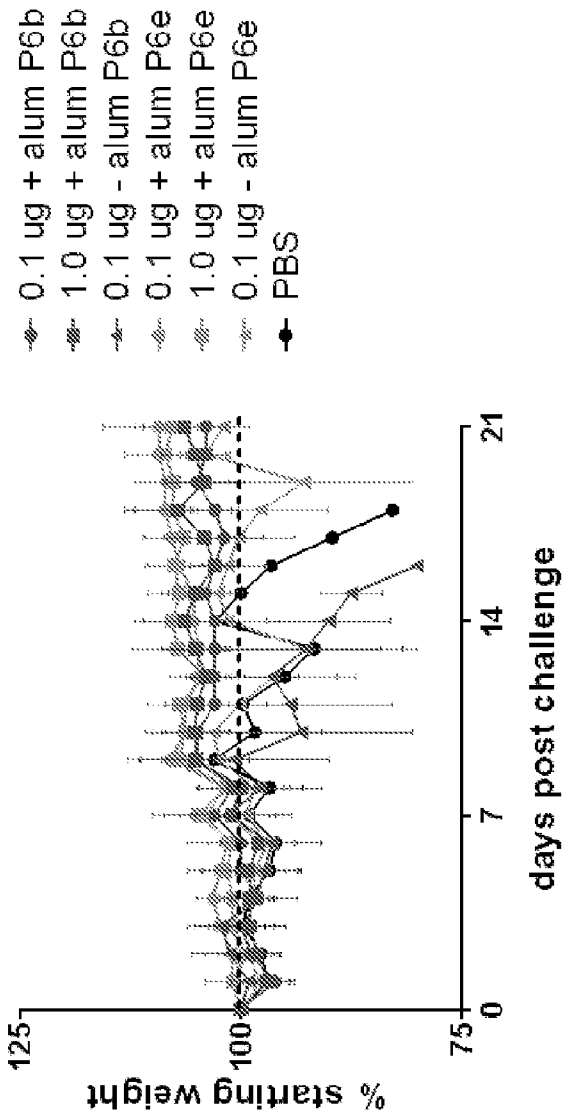

0.1 ug + alum P6b
1.0 ug + alum P6b
0.1 ug - alum P6b
0.1 ug + alum P6e
1.0 ug + alum P6e
0.1 ug - alum P6e
PBS
% starting weight
125
100
75
0
7
14
21
days post challenge

FIG. 15

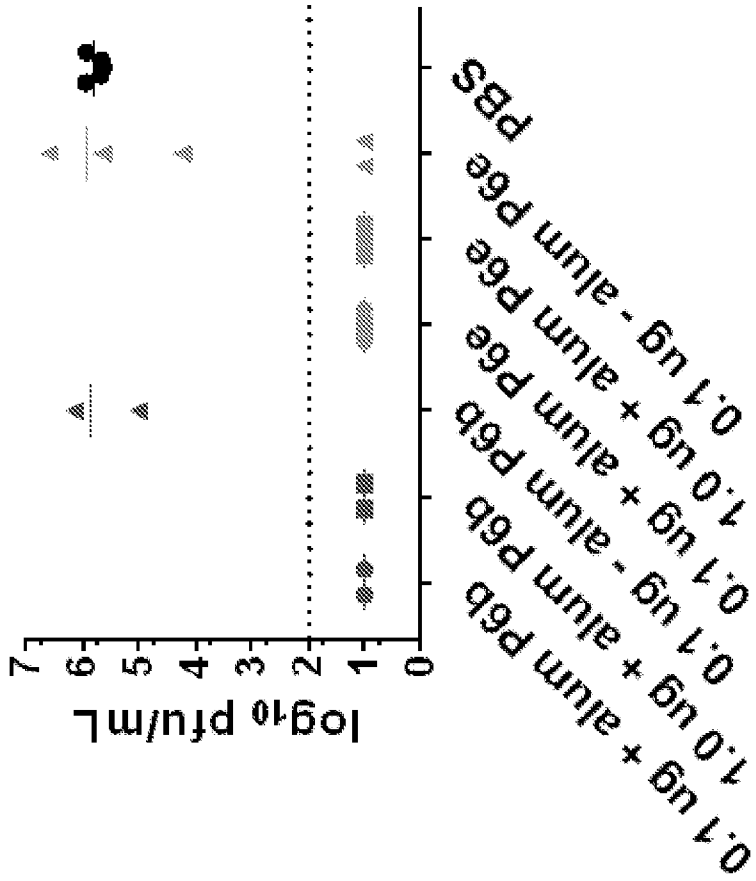

log10 pfu/mL
0
1
2
3
4
5
6
7
0.1 ug + alum P6b
1.0 ug + alum P6b
0.1 ug - alum P6b
1.0 ug + alum P6e
0.1 ug - alum P6e
PBS

FIG. 16

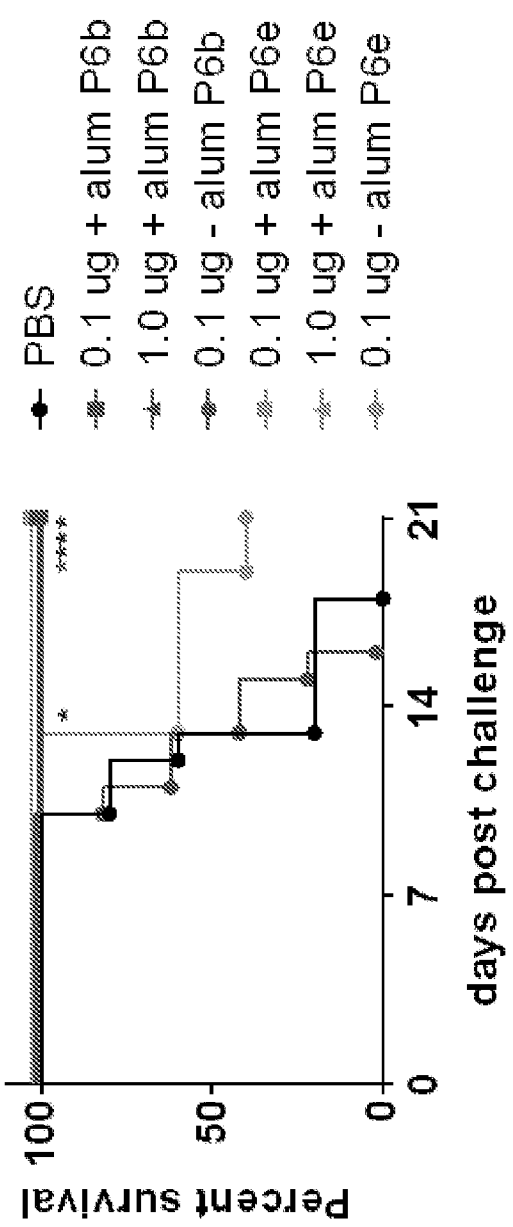

PBS
0.1 ug + alum P6b
1.0 ug + alum P6b
0.1 ug - alum P6b
0.1 ug + alum P6e
1.0 ug + alum P6e
0.1 ug - alum P6e
Percent survival
100
50
0
0
7
14
21
days post challenge
****
*

FIG. 17

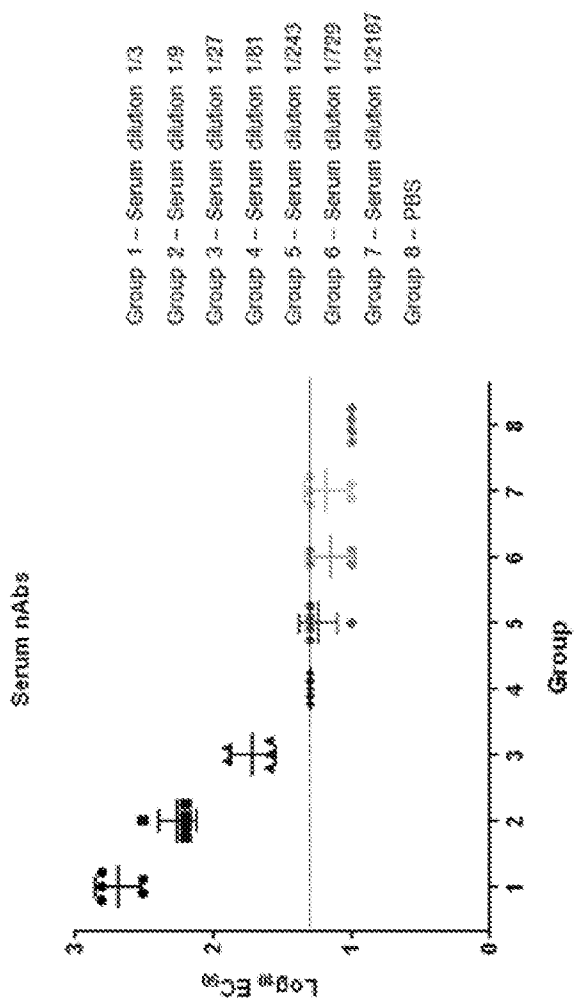

Group 1 -- Serum dilution 1/3
Group 2 -- Serum dilution 1/9
Group 3 -- Serum dilution 1/27
Group 4 -- Serum dilution 1/81
Group 5 -- Serum dilution 1/243
Group 6 -- Serum dilution 1/729
Group 7 -- Serum dilution 1/2187
Group 8 -- PBS
Serum nAbs
$\log_{10}$ EC_{50}
Group
0
1
2
3
4
5
6
7
8

FIG. 18

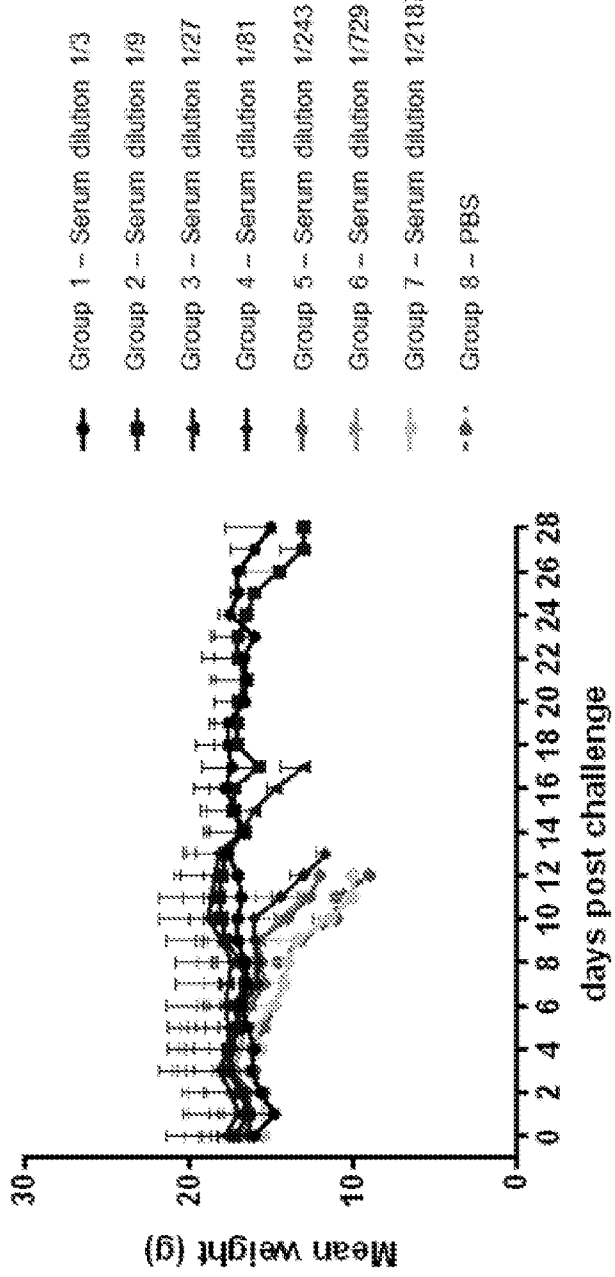

Weight
Mean weight (g)
30
20
10
0
0 2 4 6 8 10 12 14 16 18 20 22 24 26 28
days post challenge
Group 1 – Serum dilution 1/3
Group 2 – Serum dilution 1/9
Group 3 – Serum dilution 1/27
Group 4 – Serum dilution 1/81
Group 5 – Serum dilution 1/243
Group 6 – Serum dilution 1/729
Group 7 – Serum dilution 1/2187
Group 8 – PBS

FIG. 19

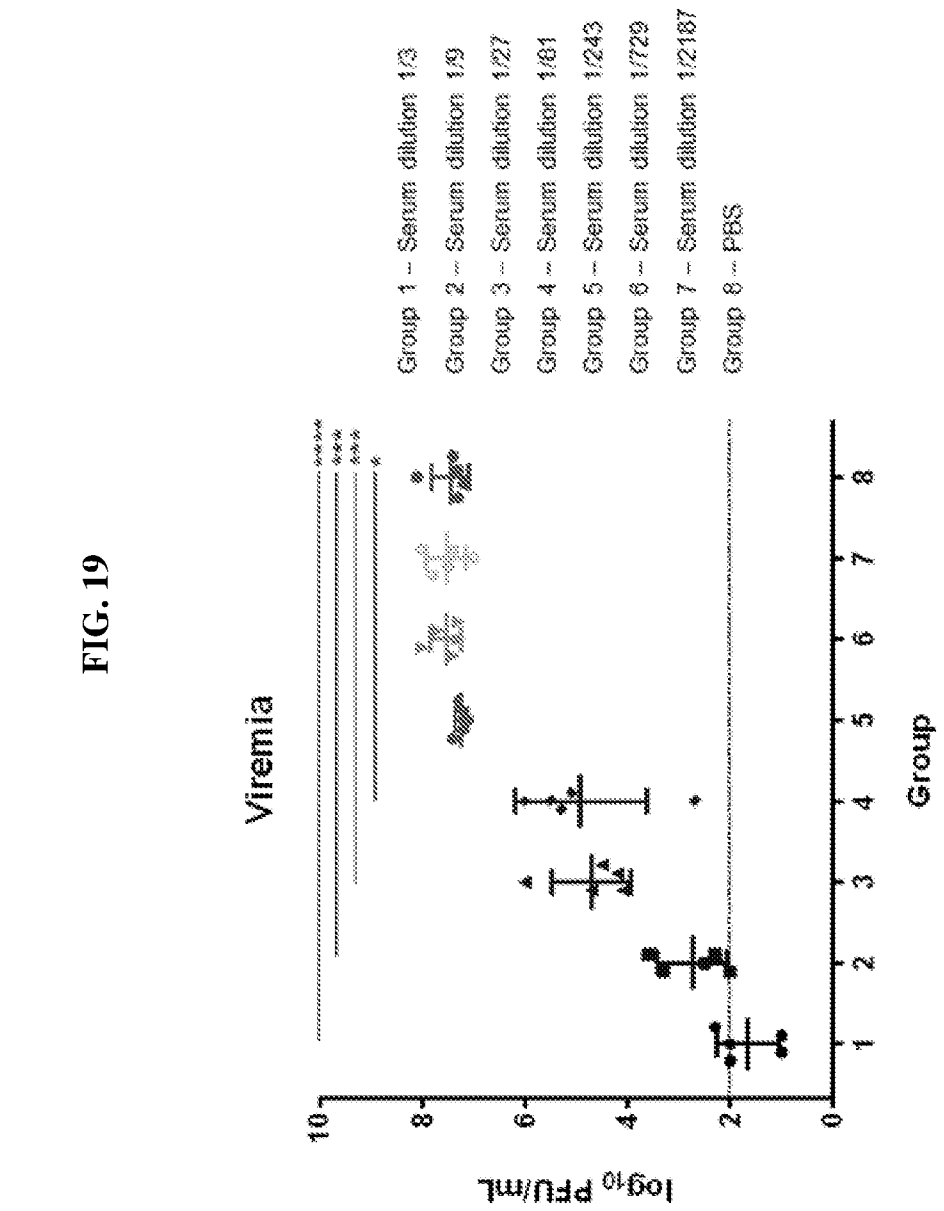

Viremia
log10 PFU/mL
Group
Group 1 – Serum dilution 1/3
Group 2 – Serum dilution 1/9
Group 3 – Serum dilution 1/27
Group 4 – Serum dilution 1/81
Group 5 – Serum dilution 1/243
Group 6 – Serum dilution 1/729
Group 7 – Serum dilution 1/2187
Group 8 – PBS
****
***
*

FIG. 20

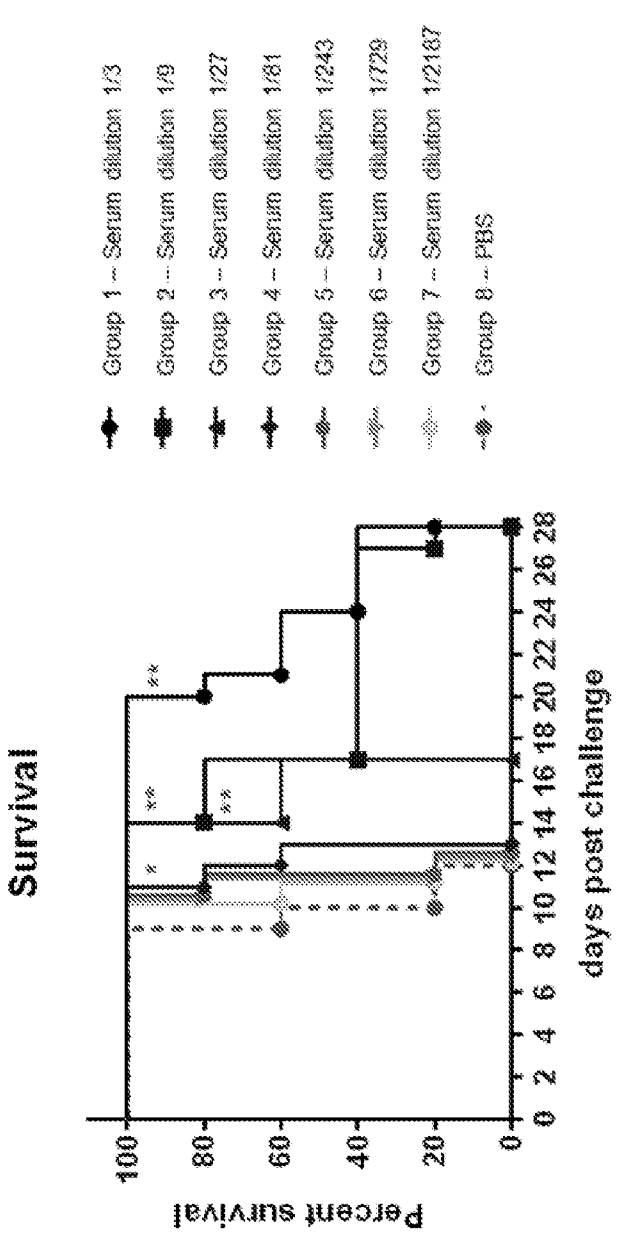
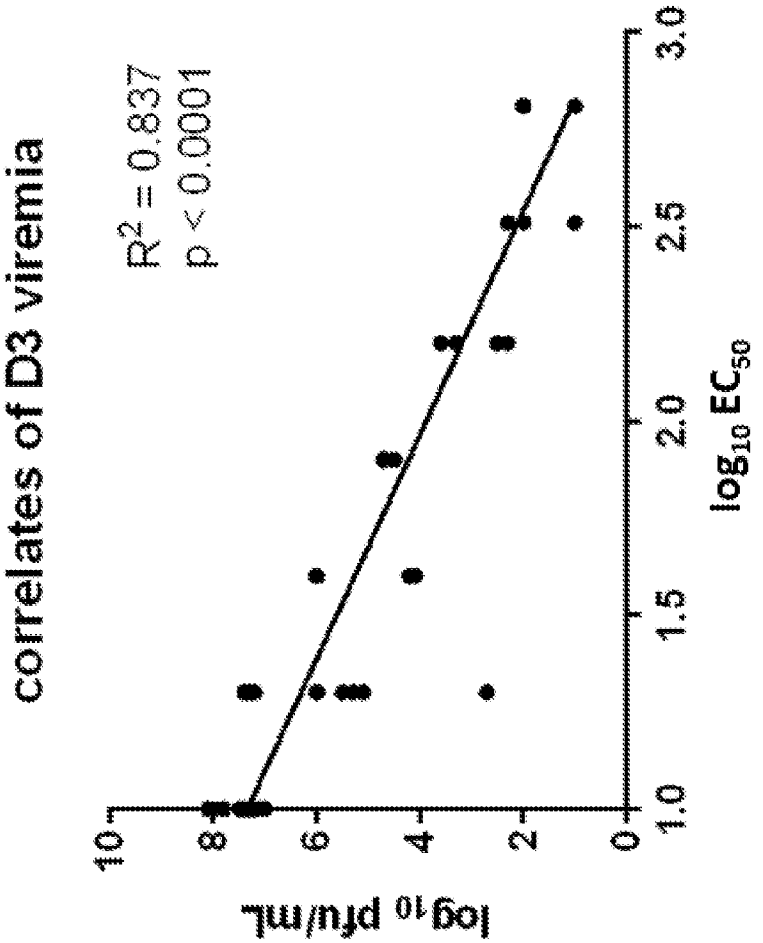

Survival
Percent survival
days post challenge
0
20
40
60
80
100
0 2 4 6 8 10 12 14 16 18 20 22 24 26 28
*
**
**
**
Group 1 -- Serum dilution 1/3
Group 2 -- Serum dilution 1/9
Group 3 -- Serum dilution 1/27
Group 4 -- Serum dilution 1/81
Group 5 -- Serum dilution 1/243
Group 6 -- Serum dilution 1/729
Group 7 -- Serum dilution 1/2187
Group 8 -- PBS

FIG. 21


correlates of D3 viremia
$R^2 = 0.837$
$p < 0.0001$
$\log_{10}$ pfu/mL
$\log_{10} EC_{50}$
10
8
6
4
2
0
1.0
1.5
2.0
2.5
3.0

PTO/AIA/14 (11-15)
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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

The application data sheet is part of the provisional or nonprovisional application for which it is being submitted. The following form contains the bibliographic data arranged in a format specified by the United States Patent and Trademark Office as outlined in 37 CFR 1.76.
This document may be completed electronically and submitted to the Office in electronic format using the Electronic Filing System (EFS) or the document may be printed and included in a paper filed application.

Secrecy Order 37 CFR 5.2:

☐ Portions or all of the application associated with this Application Data Sheet may fall under a Secrecy Order pursuant to 37 CFR 5.2 (Paper filers only. Applications that fall under Secrecy Order may not be filed electronically.)

Inventor Information:

Inventor 1 Remove

Legal Name

Prefix	Given Name	Middle Name	Family Name	Suffix
	Jill	A.	Livengood	

Residence Information (Select One) ● US Residency Non US Residency Active US Military Service

City	State/Province	Country of Residence
Fort Collins	CO	US

Mailing Address of Inventor:

Address 1	c/o Takeda Vaccines, Inc.
Address 2	75 Sidney Street
City	Cambridge
State/Province	MA
Postal Code	02139
Country	US

Inventor 2 Remove

Legal Name

Prefix	Given Name	Middle Name	Family Name	Suffix
	Holli		Giebler	

Residence Information (Select One) ◉ US Residency Non US Residency Active US Military Service

City	State/Province	Country of Residence
Fort Collins	CO	US

Mailing Address of Inventor:

Address 1	c/o Takeda Vaccines, Inc.
Address 2	75 Sidney Street
City	Cambridge
State/Province	MA
Postal Code	02139
Country	US

Inventor 3 Remove

Legal Name

Prefix	Given Name	Middle Name	Family Name	Suffix
	Whitney		Baldwin	

Residence Information (Select One) ◉ US Residency Non US Residency Active US Military Service

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

City	Fort Collins	State/Province	CO	Country of Residence	US

Mailing Address of Inventor:

Address 1	c/o Takeda Vaccines, Inc.		
Address 2	75 Sidney Street		
City	Cambridge	State/Province	MA
Postal Code	02139	Country	US

Inventor 4 — Remove

Legal Name

Prefix	Given Name	Middle Name	Family Name	Suffix
	Claire	Y.H.	Kinney	

Residence Information (Select One) ◉ US Residency ○ Non US Residency ○ Active US Military Service

City	Fort Collins	State/Province	CO	Country of Residence	US

Mailing Address of Inventor:

Address 1	c/o Centers For Disease Control And Prevention		
Address 2	Technology Transfer Office, 4770 Buford Hwy (K79)		
City	Atlanta	State/Province	GA
Postal Code	30341	Country	US

All Inventors Must Be Listed - Additional Inventor Information blocks may be generated within this form by selecting the **Add** button. Add

Correspondence Information:

Enter either Customer Number or complete the Correspondence Information section below.
For further information see 37 CFR 1.33(a).

☐ **An Address is being provided for the correspondence Information of this application.**

Customer Number	20872		
Email Address	PatentDocket@mofo.com	Add Email	Remove Email

Application Information:

Title of the Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME		
Attorney Docket Number	606773001300	Small Entity Status Claimed	☐
Application Type	Provisional		
Subject Matter	Utility		
Total Number of Drawing Sheets (if any)	21	Suggested Figure for Publication (if any)	

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

Filing By Reference:

Only complete this section when filing an application by reference under 35 U.S.C. 111(c) and 37 CFR 1.57(a). Do not complete this section if application papers including a specification and any drawings are being filed. Any domestic benefit or foreign priority information must be provided in the appropriate section(s) below (i.e., "Domestic Benefit/National Stage Information" and "Foreign Priority Information").

For the purposes of a filing date under 37 CFR 1.53(b), the description and any drawings of the present application are replaced by this reference to the previously filed application, subject to conditions and requirements of 37 CFR 1.57(a).

Application number of the previously filed application	Filing date (YYYY-MM-DD)	Intellectual Property Authority or Country

Publication Information:

☐ Request Early Publication (Fee required at time of Request 37 CFR 1.219)

☐ **Request Not to Publish.** I hereby request that the attached application not be published under 35 U.S.C. 122(b) and certify that the invention disclosed in the attached application **has not and will not** be the subject of an application filed in another country, or under a multilateral international agreement, that requires publication at eighteen months after filing.

Representative Information:

Representative information should be provided for all practitioners having a power of attorney in the application. Providing this information in the Application Data Sheet does not constitute a power of attorney in the application (see 37 CFR 1.32).
Either enter Customer Number or complete the Representative Name section below. If both sections are completed the customer Number will be used for the Representative Information during processing.

Please Select One:	● Customer Number	US Patent Practitioner	○ Limited Recognition (37 CFR 11.9)
Customer Number	20872		

Domestic Benefit/National Stage Information:

This section allows for the applicant to either claim benefit under 35 U.S.C. 119(e), 120, 121, 365(c), or 386(c) or indicate National Stage entry from a PCT application. Providing benefit claim information in the Application Data Sheet constitutes the specific reference required by 35 U.S.C. 119(e) or 120, and 37 CFR 1.78.
When referring to the current application, please leave the "Application Number" field blank.

Prior Application Status			Remove
Application Number	Continuity Type	Prior Application Number	Filing or 371(c) Date (YYYY-MM-DD)

Additional Domestic Benefit/National Stage Data may be generated within this form by selecting the **Add** button. Add

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

Foreign Priority Information:

This section allows for the applicant to claim priority to a foreign application. Providing this information in the application data sheet constitutes the claim for priority as required by 35 U.S.C. 119(b) and 37 CFR 1.55. When priority is claimed to a foreign application that is eligible for retrieval under the priority document exchange program (PDX)[i] the information will be used by the Office to automatically attempt retrieval pursuant to 37 CFR 1.55(i)(1) and (2). Under the PDX program, applicant bears the ultimate responsibility for ensuring that a copy of the foreign application is received by the Office from the participating foreign intellectual property office, or a certified copy of the foreign priority application is filed, within the time period specified in 37 CFR 1.55(g)(1).

Remove

Application Number	Country[i]	Filing Date (YYYY-MM-DD)	Access Code[i] (if applicable)

Additional Foreign Priority Data may be generated within this form by selecting the **Add** button. Add

Statement under 37 CFR 1.55 or 1.78 for AIA (First Inventor to File) Transition Applications

☐ This application (1) claims priority to or the benefit of an application filed before March 16, 2013 and (2) also contains, or contained at any time, a claim to a claimed invention that has an effective filing date on or after March 16, 2013.
NOTE: By providing this statement under 37 CFR 1.55 or 1.78, this application, with a filing date on or after March 16, 2013, will be examined under the first inventor to file provisions of the AIA.

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

Authorization or Opt-Out of Authorization to Permit Access:

When this Application Data Sheet is properly signed and filed with the application, applicant has provided written authority to permit a participating foreign intellectual property (IP) office access to the instant application-as-filed (see paragraph A in subsection 1 below) and the European Patent Office (EPO) access to any search results from the instant application (see paragraph B in subsection 1 below).

Should applicant choose not to provide an authorization identified in subsection 1 below, applicant **must opt-out** of the authorization by checking the corresponding box A or B or both in subsection 2 below.

NOTE: This section of the Application Data Sheet is **ONLY** reviewed and processed with the **INITIAL** filing of an application. After the initial filing of an application, an Application Data Sheet cannot be used to provide or rescind authorization for access by a foreign IP office(s). Instead, Form PTO/SB/39 or PTO/SB/69 must be used as appropriate.

1. Authorization to Permit Access by a Foreign Intellectual Property Office(s)

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

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Providing assignment information in this section does not substitute for compliance with any requirement of part 3 of Title 37 of CFR to have an assignment recorded by the Office.

Applicant 1 Remove

If the applicant is the inventor (or the remaining joint inventor or inventors under 37 CFR 1.45), this section should not be completed. The information to be provided in this section is the name and address of the legal representative who is the applicant under 37 CFR 1.43; or the name and address of the assignee, person to whom the inventor is under an obligation to assign the invention, or person who otherwise shows sufficient proprietary interest in the matter who is the applicant under 37 CFR 1.46. If the applicant is an applicant under 37 CFR 1.46 (assignee, person to whom the inventor is obligated to assign, or person who otherwise shows sufficient proprietary interest) together with one or more joint inventors, then the joint inventor or inventors who are also the applicant should be identified in this section.

Clear

● Assignee	Legal Representative under 35 U.S.C. 117	Joint Inventor
Person to whom the inventor is obligated to assign.	Person who shows sufficient proprietary interest	

If applicant is the legal representative, indicate the authority to file the patent application, the inventor is:

Name of the Deceased or Legally Incapacitated Inventor:

If the Applicant is an Organization check here. ☒

Organization Name	Takeda Vaccines, Inc.		
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Email Address			

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
	Application Number	
Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

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Clear

◉ Assignee	Legal Representative under 35 U.S.C. 117	Joint Inventor
Person to whom the inventor is obligated to assign.	Person who shows sufficient proprietary interest	

If applicant is the legal representative, indicate the authority to file the patent application, the inventor is:

Name of the Deceased or Legally Incapacitated Inventor:

If the Applicant is an Organization check here. ☒

Organization Name	Centers For Disease Control And Prevention		
Mailing Address Information For Applicant:			
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Address 2	Technology Transfer Office		
City	Atlanta	**State/Province**	GA
Country	US	Postal Code	30341
Phone Number		Fax Number	
Email Address			

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Application Data Sheet 37 CFR 1.76	Attorney Docket Number	606773001300
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Title of Invention	ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME	

Prefix	**Given Name**	Middle Name	**Family Name**	Suffix

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Address 1			
Address 2			
City		**State/Province**	
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Signature	/Roberto K. Rodriguez/			Date (YYYY-MM-DD)	2017-11-03
First Name	Roberto	Last Name	Rodriguez	Registration Number	66244

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ZIKA VACCINES AND IMMUNOGENIC COMPOSITIONS, AND METHODS OF USING THE SAME

STATEMENT REGARDING FEDERALLY-SPONSORED RESEARCH

[0001] This invention was made with government support under Grant No. HHSO100201600015C awarded by the Department of Health and Human Services, Office of the Assistant Secretary for Preparedness and Response, Biomedical Advanced Research and Development Authority. This invention was created in the performance of a Cooperative Research and Development Agreement with the Centers for Disease Control and Prevention, an Agency of the Department of Health and Human Services. The Government of the United States has certain rights in the invention.

FIELD OF THE INVENTION

[0002] The present disclosure relates to Zika virus vaccines and immunogenic compositions having one or more antigens from a non-human cell adapted Zika virus, and methods of manufacture, formulation, testing, and uses thereof.

BACKGROUND

[0003] Zika virus, a *flavivirus* classified with other mosquito-borne viruses (*e.g.*, yellow fever, dengue, West Nile, and Japanese encephalitis viruses) within the *Flaviviridae* family has spread rapidly in a hemispheric-wide epidemic since the virus was introduced into Brazil in 2013. The virus has reached the Central and North Americas, including territories of the United States, consequently now threatening the continental US. Indeed, Zika virus strain PRVABC59 was isolated from serum from a person who had traveled to Puerto Rico in 2015. The genome of this strain has been sequenced at least three times (*See* Lanciotti *et al.* Emerg. Infect. Dis. 2016 May;22(5):933-5 and GenBank Accession Number KU501215.1; GenBank Accession Number KX087101.3; and Yun *et al.* Genome Announc. 2016 Aug 18;4(4) and GenBank Accession Number ANK57897.1).

[0004] Initially isolated in 1947 in Uganda, the virus was first linked to human disease in 1952, and has been recognized sporadically as a cause of mild, self-limited febrile illness in Africa and Southeast Asia (Weaver *et al.* 2016; Faria *et al.* 2016). However, in 2007, an outbreak appeared in the North Pacific island of Yap, and then disseminated from island to island across the Pacific, leading to an extensive outbreak in 2013-2014 in French

Polynesia, spreading then to New Caledonia, the Cook Islands, and ultimately, to Easter Island. An Asian lineage virus was subsequently transferred to the Western Hemisphere by routes that remain undetermined (Faria *et al.* 2016). The virus may be transmitted zoonotically by *Aedes aegypti*, *A. albopictus*, and possibly by *A. hensilli* and *A. polynieseinsis* (Weaver *et al.* 2016). Additionally, it is thought that other vectors for transmitting the virus may exist, and the virus may be transmitted by blood transfusion, transplacentally, and/or through sexual transmission.

[0005] In late 2015, a significant increase in fetal abnormalities (*e.g.,* microcephaly) and Guillain-Barre syndrome (GBS) in areas of widespread Zika virus infection raised alarm that Zika virus might be much more virulent than originally thought, prompting the World Health Organization (WHO) to declare a Public Health Emergency of International Concern (PHEIC) (Heymann *et al.* 2016). While Zika virus poses a substantial public health threat, no FDA-approved vaccine or treatment currently exists, and the only preventative measures for controlling Zika virus involve managing mosquito populations.

[0006] In recent efforts to characterize a recombinant Zika virus for the development of a potential vaccine, a non-human cell adapted Zika virus was identified that harbors a mutation in the viral Envelope protein at position 330 (Weger-Lucarelli *et al.* 2017. Journal of Virology). The authors of this study found that full-length infectious cDNA clones of Zika virus strain PRVABC59 were genetically unstable when amplified during cloning, and opted to split the viral genome to address the observed instability, developing and applying a two plasmid system. However, a two plasmid system for the development of a Zika vaccine is less desirable.

BRIEF SUMMARY

[0007] Thus, there is a need to develop vaccines and immunogenic compositions for treating and/or preventing Zika virus infection that utilize a genetically stable Zika virus that does not require a two vector system. To meet the above and other needs, the present disclosure is directed to a genetically stable non-human cell adapted Zika virus harboring an adaptation in the Non-structural protein 1 (with a wild-type Envelope protein), allowing for the use a single virus/viral genome system for vaccine production. Accordingly, the present disclosure provides vaccines and immunogenic compositions useful for treating and/or preventing Zika virus infection (*e.g.,* in humans) that include one or more Zika virus

antigens (*e.g.*, one or more antigens from a whole inactivated Zika virus) from a Zika virus harboring at least one non-human cell adaptation mutation (*e.g.*, a mutation in Zika virus Non-structural protein 1).

[0008] The present disclosure is based, at least in part, on the surprising finding that both high and low dose vaccines comprising one or more antigens from separately derived clonal virus populations of non-human cell adapted Zika virus were able to induce robust immune responses and provide significant protection from Zika virus infection (*See* Example 2 below). Clonal isolation of the Zika virus strains also allowed for the successful purification of the virus away from contaminating agents (*e.g.*, adventitious agents that may be co-purified with the parental strain). Moreover, the present disclosure is based, at least in part, on the finding that clonal isolated Zika viruses harboring an adaptation mutation in protein NS1 grew well and predictably in Vero cells to high titer, and surprisingly, were genetically stable/genetically homogenous without any detectable mutations to the viral envelope protein (*See* Examples 1 and 2 below). While a similar mutation in Zika virus Non-structural protein 1 may have been observed in the genomic sequencing analysis of 1 out of 3 published sequences of Zika virus strain PRVABC59 (Yun *et al.* Genome Announc. 2016 Aug 18;4(4)), this reference fails to teach or suggest that a mutation in NS1 may improve stability of the virus; fails to teach or suggest that a virus harboring the mutation may be used in the development of an effective vaccine against Zika virus; and fails to teach or suggest that such a vaccine may be effective in inducing a robust immune response and providing significant protection from Zika virus infection when used at both low and high doses. Thus, without wishing to be bound by theory, the inventors of the present disclosure have determined that the adaptation mutation in protein NS1 appeared to enhance genetic stability within the Zika virus, resulting in increased/enhanced replication efficiency. Further, the Zika strain harboring an adaptation mutation in protein NS1 was able to be passaged multiple times without developing further mutations. Such a stable Zika virus strain is advantageous as a master virus seed (MVS), or subsequent seeds derived from the MVS, for vaccine production and manufacturing, as the risk of the master virus seed developing undesirable mutations is reduced. Moreover, without wishing to be bound by theory, the adaptation mutation in protein NS1 of the Zika strain of the present disclosure may also reduce or otherwise inhibit the occurrence of undesirable mutations, such as within the envelope protein E (Env) of the Zika virus strain.

[0009] Accordingly, certain aspects of the present disclosure relate to a vaccine or immunogenic composition containing one or more antigen from a Zika virus, where the Zika virus contains at least one non-human cell adaptation mutation. In some embodiments, the at least one non-human cell adaptation mutation is in Zika virus Non-structural protein 1 (NS1). In some embodiments, the at least one adaptation mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1. In some embodiments, the at least one adaptation mutation is a Trp98Gly mutation.

[0010] In some embodiments that may be combined with any of the preceding embodiments, the at least one adaptation mutation enhances genetic stability as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments that may be combined with any of the preceding embodiments, the at least one adaptation mutation enhances viral replication as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus does not comprise a mutation in Envelope protein E (Env).

[0011] In some embodiments that may be combined with any of the preceding embodiments, the non-human cell is a mammalian cell. In some embodiments that may be combined with any of the preceding embodiments, the non-human cell is a monkey cell. In some embodiments, the monkey cell is from a Vero cell line. In some embodiments, the Vero cell line is a WHO Vero 10-87 cell line.

[0012] In some embodiments that may be combined with any of the preceding embodiments, the Zika virus is an African lineage virus or an Asian lineage virus. In some embodiments, the Zika virus is an Asian lineage virus. In some embodiments, the Zika virus is from strain PRVABC59.

[0013] In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a purified antigen vaccine or immunogenic composition, a subunit vaccine or immunogenic composition, an inactivated whole virus vaccine or immunogenic composition, or an attenuated virus vaccine or immunogenic composition. In some embodiments, the vaccine or immunogenic composition is an inactivated whole virus vaccine or immunogenic composition. In some embodiments that may be combined with any of the preceding embodiments, the virus was chemically inactivated. In some embodiments, the virus was chemically inactivated with

one or more of a detergent, formalin, beta-propiolactone (BPL), binary ethylamine (BEI), acetyl ethyleneimine, methylene blue, and psoralen. In some embodiments, the virus was chemically inactivated with formalin.

[0014] In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition further contains an adjuvant. In some embodiments, the adjuvant is selected from aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and/or Incomplete Freund's Adjuvant (IFA). In some embodiments, the adjuvant is an aluminum salt. In some embodiments, the adjuvant is selected from the group consisting of alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85. In some embodiments, at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of the one or more antigens are adsorbed to the adjuvant.

[0015] In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a low dose vaccine or immunogenic composition (*e.g.,* containing from about 1μg to about 2μg antigen). In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a high dose vaccine or immunogenic composition (*e.g.,* containing about 10μg antigen). In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition contains from about 0.1 μg to about 25 μg of the one or more antigens. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition contains from about 0.1 μg Env to about 100 μg Env. In some embodiments, the vaccine or immunogenic composition is unadjuvanted. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus is a clonal isolate. In some embodiments, the clonal isolate is substantially free of one or more adventitious agents (*e.g.,* free of one or more adventitious agents that may be co-purified with the parental strain).

[0016] Other aspects of the present disclosure relate to a vaccine or immunogenic composition containing: a) an aluminum salt adjuvant; and b) a whole inactivated Zika virus, where the Zika virus contains a non-human cell adaptation mutation, and where the non-human cell adaptation mutation is a Trp98Gly mutation at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1.

[0017] Other aspects of the present disclosure relate to a method of treating or preventing Zika virus infection in a subject in need thereof, including administering to the subject a therapeutically effective amount of any of the vaccines or immunogenic compositions described herein.

[0018] Other aspects of the present disclosure relate to a method for inducing an immune response in a subject in need thereof, including administering to the subject an immunogenic amount of any of the vaccines or immunogenic compositions described herein.

[0019] In some embodiments that may be combined with nay of the preceding embodiments, the subject is a human. In some embodiments, the subject is pregnant or intends to become pregnant.

[0020] In some embodiments that may be combined with any of the preceding embodiments, administration of the vaccine or immunogenic composition induces a protective immune response in the subject. In some embodiments, the protective immune response induced in the subject is greater than a protective immune response induced in a corresponding subject administered a vaccine or immunogenic composition containing one or more antigens from a Zika virus lacking the at least one non-human cell adaptation mutation. In some embodiments that may be combined with any of the preceding embodiments, administration of the vaccine or immunogenic composition induces the generation of neutralizing antibodies to Zika virus in the subject. In some embodiments, the concentration of neutralizing antibodies generated in the subject is higher than a concentration of neutralizing antibodies generated in a corresponding subject administered a vaccine or immunogenic composition comprising one or more antigens from a Zika virus lacking the at least one non-human cell adaptation mutation.

[0021] In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is administered by a route selected

from subcutaneous administration, transcutaneous administration, intradermal administration, subdermal administration, intramuscular administration, peroral administration, intranasal administration, buccal administration, intraperitoneal administration, intravaginal administration, anal administration and/or intracranial administration. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is administered one or more times. In some embodiments, the vaccine or immunogenic composition is administered as a first (prime) and a second (boost) administration. In some embodiments, the second (boost) administration is administered at least 28 days after the first (prime) administration.

[0022] Other aspects of the present disclosure relate to a method for inactivating a Zika virus preparation, including (a) isolating the Zika virus preparation from one or more non-human cells, where the cells are used to produce the virus preparation, and where the Zika virus contains at least one non-human cell adaptation mutation; and (b) treating the virus preparation with an effective amount of formalin. In some embodiments, the method further includes (c) neutralizing the formalin-treated virus preparation with sodium metabisulfite. In some embodiments, the virus preparation is neutralized at least five, at least seven, at least nine, at least 11, or at least 14 days after formalin treatment. In some embodiments, the method further includes (d) purifying the neutralized virus preparation. In some embodiments, the neutralized virus preparation is purified by process selected from cross flow filtration (CFF), multimodal chromatography, size exclusion chromatography, cation exchange chromatography, and/or anion exchange chromatography.

[0023] In some embodiments that may be combined with any of the preceding embodiments, the virus preparation is mixed with an adjuvant. In some embodiments, the adjuvant is selected from aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and/or Incomplete Freund's Adjuvant (IFA). In some embodiments, the adjuvant is an aluminum salt. In some embodiments, the adjuvant is selected from alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and/or Alhydrogel 85. In some

embodiments, at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of one or more antigens in the virus preparation are adsorbed to the adjuvant.

[0024] In some embodiments that may be combined with any of the preceding embodiments, the at least one non-human cell adaptation mutation is in Zika virus NS1. In some embodiments, the at least one adaptation mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1. In some embodiments, the at least one adaptation mutation is a Trp98Gly mutation. In some embodiments that may be combined with any of the preceding embodiments, the at least one adaptation mutation enhances genetic stability as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments that may be combined with any of the preceding embodiments, the at least one adaptation mutation enhances viral replication as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus does not comprise a mutation in Envelope protein E (Env).

[0025] Other aspects of the present disclosure related to a genetically stable, clonally-derived Zika virus harboring a mutation in the Non-structural protein 1 (with a wild-type Envelope protein), allowing for the use a single virus/viral genome system for vaccine production. Accordingly, the present disclosure provides vaccines and immunogenic compositions useful for treating and/or preventing Zika virus infection (*e.g.,* in humans) that include one or more Zika virus antigens (*e.g.,* one or more antigens from a whole inactivated Zika virus) from a Zika virus harboring at least one mutation (*e.g.,* a mutation in Zika virus Non-structural protein 1). The present disclosure is based, at least in part, on the surprising finding that both high and low dose vaccines comprising one or more antigens from separately derived clonal virus populations of Zika virus having a mutation in Non-structural protein 1 (NS1) were able to induce robust immune responses and provide significant protection from Zika virus infection (*See* Example 2 below). Clonal isolation of the Zika virus strains also allowed for the successful purification of the virus away from contaminating agents (*e.g.,* adventitious agents that may be co-purified with the parental strain). Moreover, the present disclosure is based, at least in part, on the finding that clonal isolated Zika viruses harboring a mutation in protein NS1 grew well and predictably in Vero cells to high titer, and surprisingly, were genetically stable/genetically homogenous

without any detectable mutations to the viral envelope protein (*See* Examples 1 and 2 below). While a similar mutation in Zika virus NS1 may have been observed in the genomic sequencing analysis of 1 out of 3 published sequences of Zika virus strain PRVABC59 (Yun *et al.* Genome Announc. 2016 Aug 18;4(4)), this reference fails to teach or suggest that a mutation in NS1 may improve stability of the virus; fails to teach or suggest that a virus harboring the mutation may be used in the development of an effective vaccine against Zika virus; and fails to teach or suggest that such a vaccine may be effective in inducing a robust immune response and providing significant protection from Zika virus infection when used at both low and high doses. Thus, without wishing to be bound by theory, the inventors of the present disclosure have determined that the mutation in protein NS1 appeared to enhance genetic stability within the Zika virus, resulting in increased/enhanced replication efficiency. Further, the Zika strain harboring an adaptation mutation in protein NS1 was able to be passaged multiple times without developing further mutations. Such a stable Zika virus strain is advantageous as a master virus seed (MVS), or subsequent seeds derived from the MVS, for vaccine production and manufacturing, as the risk of the master virus seed developing undesirable mutations is reduced. Moreover, without wishing to be bound by theory, the mutation in protein NS1 of the Zika strain of the present disclosure may also reduce or otherwise inhibit the occurrence of undesirable mutations, such as within the envelope protein E (Env) of the Zika virus strain.

[0026] Accordingly, certain aspects of the present disclosure relate to a vaccine or immunogenic composition containing one or more antigen from a Zika virus, where the Zika virus contains at least one mutation. In some embodiments, the at least one mutation is in Zika virus Non-structural protein 1 (NS1). In some embodiments, the at least one mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1. In some embodiments, the at least one mutation is a Trp98Gly mutation. In some embodiments, the Zika virus does not contain a Tryptophan (Trp) at position 98 of the NS1 protein (SEQ ID NO: 1).

[0027] In some embodiments that may be combined with any of the preceding embodiments, the at least one mutation enhances genetic stability as compared to a Zika virus lacking the at least one mutation. In some embodiments that may be combined with any of the preceding embodiments, the at least one mutation enhances viral replication as compared to a Zika virus lacking the at least one mutation. In some embodiments that may

be combined with any of the preceding embodiments, the Zika virus does not comprise a mutation in Envelope protein E (Env). In some embodiments that may be combined with any of the preceding embodiments, the non-human cell is a mammalian cell. In some embodiments that may be combined with any of the preceding embodiments, the non-human cell is a monkey cell. In some embodiments, the monkey cell is from a Vero cell line. In some embodiments, the Vero cell line is a WHO Vero 10-87 cell line. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus is an African lineage virus or an Asian lineage virus. In some embodiments, the Zika virus is an Asian lineage virus. In some embodiments, the Zika virus is from strain PRVABC59. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a purified antigen vaccine or immunogenic composition, a subunit vaccine or immunogenic composition, an inactivated whole virus vaccine or immunogenic composition, or an attenuated virus vaccine or immunogenic composition. In some embodiments, the vaccine or immunogenic composition is an inactivated whole virus vaccine or immunogenic composition. In some embodiments that may be combined with any of the preceding embodiments, the virus was chemically inactivated. In some embodiments, the virus was chemically inactivated with one or more of a detergent, formalin, beta-propiolactone (BPL), binary ethylamine (BEI), acetyl ethyleneimine, methylene blue, and psoralen. In some embodiments, the virus was chemically inactivated with formalin. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition further contains an adjuvant. In some embodiments, the adjuvant is selected from aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and/or Incomplete Freund's Adjuvant (IFA). In some embodiments, the adjuvant is an aluminum salt. In some embodiments, the adjuvant is selected from the group consisting of alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85. In some embodiments, at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of the one or more antigens are adsorbed to the adjuvant. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a low

dose vaccine or immunogenic composition (*e.g.*, containing from about 1µg to about 2µg antigen). In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is a high dose vaccine or immunogenic composition (*e.g.*, containing about 10µg antigen). In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition contains from about 0.1 µg to about 25 µg of the one or more antigens. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition contains from about 0.1 µg Env to about 100 µg Env. In some embodiments, the vaccine or immunogenic composition is unadjuvanted. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus is a clonal isolate. In some embodiments, the clonal isolate is substantially free of one or more adventitious agents (*e.g.*, free of one or more adventitious agents that may be co-purified with the parental strain).

[0028] Other aspects of the present disclosure relate to a vaccine or immunogenic composition containing: a) an aluminum salt adjuvant; and b) a whole inactivated Zika virus, where the Zika virus contains a mutation, and where the mutation is a Trp98Gly mutation at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1. In some embodiments, the Zika virus does not contain a Tryptophan (Trp) at position 98 of the NS1 protein (SEQ ID NO: 1).

[0029] Other aspects of the present disclosure relate to a method of treating or preventing Zika virus infection in a subject in need thereof, including administering to the subject a therapeutically effective amount of any of the vaccines or immunogenic compositions described herein. Other aspects of the present disclosure relate to a method for inducing an immune response in a subject in need thereof, including administering to the subject an immunogenic amount of any of the vaccines or immunogenic compositions described herein.

[0030] In some embodiments that may be combined with nay of the preceding embodiments, the subject is a human. In some embodiments, the subject is pregnant or intends to become pregnant. In some embodiments that may be combined with any of the preceding embodiments, administration of the vaccine or immunogenic composition induces a protective immune response in the subject. In some embodiments, the protective immune response induced in the subject is greater than a protective immune response

induced in a corresponding subject administered a vaccine or immunogenic composition containing one or more antigens from a Zika virus lacking the at least one mutation. In some embodiments that may be combined with any of the preceding embodiments, administration of the vaccine or immunogenic composition induces the generation of neutralizing antibodies to Zika virus in the subject. In some embodiments, the concentration of neutralizing antibodies generated in the subject is higher than a concentration of neutralizing antibodies generated in a corresponding subject administered a vaccine or immunogenic composition comprising one or more antigens from a Zika virus lacking the at least one mutation. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is administered by a route selected from subcutaneous administration, transcutaneous administration, intradermal administration, subdermal administration, intramuscular administration, peroral administration, intranasal administration, buccal administration, intraperitoneal administration, intravaginal administration, anal administration and/or intracranial administration. In some embodiments that may be combined with any of the preceding embodiments, the vaccine or immunogenic composition is administered one or more times. In some embodiments, the vaccine or immunogenic composition is administered as a first (prime) and a second (boost) administration. In some embodiments, the second (boost) administration is administered at least 28 days after the first (prime) administration.

[0031] Other aspects of the present disclosure relate to a method for inactivating a Zika virus preparation, including (a) isolating the Zika virus preparation from one or more non-human cells, where the cells are used to produce the virus preparation, and where the Zika virus contains at least one mutation; and (b) treating the virus preparation with an effective amount of formalin. In some embodiments, the method further includes (c) neutralizing the formalin-treated virus preparation with sodium metabisulfite. In some embodiments, the virus preparation is neutralized at least five, at least seven, at least nine, at least 11, or at least 14 days after formalin treatment. In some embodiments, the method further includes (d) purifying the neutralized virus preparation. In some embodiments, the neutralized virus preparation is purified by process selected from cross flow filtration (CFF), multimodal chromatography, size exclusion chromatography, cation exchange chromatography, and/or anion exchange chromatography.

[0032] In some embodiments that may be combined with any of the preceding embodiments, the virus preparation is mixed with an adjuvant. In some embodiments, the adjuvant is selected from aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and/or Incomplete Freund's Adjuvant (IFA). In some embodiments, the adjuvant is an aluminum salt. In some embodiments, the adjuvant is selected from alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and/or Alhydrogel 85. In some embodiments, at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of one or more antigens in the virus preparation are adsorbed to the adjuvant. In some embodiments that may be combined with any of the preceding embodiments, the at least one mutation is in Zika virus NS1. In some embodiments, the at least one mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1. In some embodiments, the at least one mutation is a Trp98Gly mutation. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus does not contain a Tryptophan (Trp) at position 98 of the NS1 protein (SEQ ID NO: 1). In some embodiments that may be combined with any of the preceding embodiments, the at least one mutation enhances genetic stability as compared to a Zika virus lacking the at least one mutation. In some embodiments that may be combined with any of the preceding embodiments, the at least one mutation enhances viral replication as compared to a Zika virus lacking the at least one mutation. In some embodiments that may be combined with any of the preceding embodiments, the Zika virus does not comprise a mutation in Envelope protein E (Env).

[0033] It is to be understood that one, some, or all of the properties of the various embodiments described above and herein may be combined to form other embodiments of the present disclosure. These and other aspects of the present disclosure will become apparent to one of skill in the art. These and other embodiments of the present disclosure are further described by the detailed description that follows.

BRIEF DESCRIPTION OF THE DRAWINGS

[0034] **FIG. 1** shows bright field microscopy images of Vero cell monolayers mock infected (top) or infected with ZIKAV strain PRVABC59 (bottom).

[0035] **FIG. 2** shows growth kinetics of ZIKAV PRVABC59 P1 on Vero cell monolayers, as determined by $TCID_{50}$.

[0036] **FIG. 3** shows potency assay testing ($TCID_{50}$) of Zika virus PRVABC59 P5 clones a-f.

[0037] **FIG. 4** shows bright-field microscopy images depicting the cytopathic effect (CPE) of growth of Zika virus PRVABC59 P6 clones a-f on Vero cell monolayers.

[0038] **FIG. 5** shows potency assay testing ($TCID_{50}$) of Zika virus PRVABC59 P6 clones a-f

[0039] **FIG. 6** shows an amino acid sequence alignment comparing the envelope glycoprotein sequence of Zika virus near residue 330 from Zika virus strains PRVABC59 P6e (SEQ ID NO: 8) and PRVABC59 (SEQ ID NO: 9) with several other flaviviruses (WNV (SEQ ID NO: 10); JEV (SEQ ID NO: 11); SLEV (SEQ ID NO: 12); YFV (SEQ ID NO: 13); DENV 1 16007 (SEQ ID NO: 14); DENV 2 16681 (SEQ ID NO: 15); DENV 3 16562 (SEQ IDNO: 16); and DENV 4 1036 (SEQ ID NO: 17)).

[0040] **FIG. 7** shows an amino acid sequence alignment comparing the NS1 protein sequence of Zika virus near residue 98 from Zika virus strains PRVABC59 P6e (SEQ ID NO: 18) and PRVABC59 (SEQ ID NO: 19) with several other flaviviruses (WNV (SEQ ID NO: 20); JEV (SEQ ID NO: 21); SLEV (SEQ ID NO: 22); YFV (SEQ ID NO: 23); DENV 1 16007 (SEQ ID NO: 24); DENV 2 16681 (SEQ ID NO: 25); DENV 3 16562 (SEQ IDNO: 26); and DENV 4 1036 (SEQ ID NO: 27)).

[0041] **FIG. 8** shows the plaque phenotype of ZIKAV PRVABC59 P6 virus clones a-f compared to ZIKAV PRVABC59 P1 virus.

[0042] **FIG. 9** shows the mean plaque size of ZIKAV PRVABC59 P6 virus clones compared to ZIKAV PRVABC59 P1 virus.

[0043] **FIG. 10** shows the growth kinetics of ZIKAV PRVABC59 P6 clones a-f in Vero cells under serum-free growth conditions.

[0044] **FIG. 11** shows a schematic of the steps taken to prepare PRVABC59 P6b and P6e formulated drug product for the immunization experiments.

[0045] **FIG. 12A** shows the schedule of dosing of CD-1 mice with vaccine formulations derived from the ZIKAV PRVABC59 P6b and P6e clones. PBS was used as placebo.

[0046] **FIG. 12B** shows the serum ZIKAV neutralizing antibody titers of CD-1 mice immunized as described in FIG. 12A using vaccine formulations derived from ZIKAV PRVABC59 P6b and P6e clones. ZIKAV neutralizing antibody titers were determined by Reporter Virus Particle (RVP) neutralization assay. Solid lines represent the geometric mean of a group. The limit of detection (1.93 $\log_{10}$) is represented by a dashed line.

[0047] **FIG. 13A** shows the schedule of dosing of AG129 mice with vaccine formulations derived from the ZIKAV PRVABC59 P6b and P6e clines. PBS was used as a placebo.

[0048] **FIG. 13B** shows the serum ZIKAV neutralizing antibody titers of AG129 mice immunized as described in FIG. 13A using vaccine formulations derived from ZIKAV PRVABC59 P6b and P6e clones. Solid lines represent the geometric mean of a group. The limit of detection (1.30 $\log_{10}$) is represented by a dashed line. Animals with no detectable titer (<1.30) were assigned a titer of 0.5.

[0049] **FIG. 14** shows the mean weight of AG129 test groups post-challenge, represented as a percentage of starting weight. Error bars represent standard deviation.

[0050] **FIG. 15** shows the serum viremia of individual AG129 mice two days post-challenge, reported as PFU/mL. Solid lines represent the mean of a group. The limit of detection (2.0 $\log_{10}$) is represented by a dashed line.

[0051] **FIG. 16** shows the survival analysis of AG129 test groups post-challenge.

[0052] **FIG. 17** shows the pre-challenge serum circulating ZIKAV neutralizing antibody (Nab) titers following passive transfer of pooled sera from vaccinated and challenged AG129 mice.

[0053] **FIG. 18** shows the mean body weight of passive transfer and control mice challenged with Zika virus.

[0054] **FIG. 19** shows the serum viremia of individual AG129 mice three days post-challenge, reported as PFU/mL.

[0055] **FIG. 20** shows the survival analysis of passive transfer and control mice challenged with Zika virus.

[0056] **FIG. 21** shows the correlation between ZIKAV neutralizing antibody titers and viremia observed in passive transfer mice.

DETAILED DESCRIPTION

General Techniques

[0057] The techniques and procedures described or referenced herein are generally well understood and commonly employed using conventional methodology by those skilled in the art, such as, for example, the widely utilized methodologies described in Sambrook et al., *Molecular Cloning: A Laboratory Manual* 3d edition (2001) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.; *Current Protocols in Molecular Biology* (F.M. Ausubel, et al. eds., (2003)); the series *Methods in Enzymology* (Academic Press, Inc.): *PCR 2: A Practical Approach* (M.J. MacPherson, B.D. Hames and G.R. Taylor eds. (1995)), Harlow and Lane, eds. (1988) *Antibodies, A Laboratory Manual*, and *Animal Cell Culture* (R.I. Freshney, ed. (1987)); *Oligonucleotide Synthesis* (M.J. Gait, ed., 1984); *Methods in Molecular Biology*, Humana Press; *Cell Biology: A Laboratory Notebook* (J.E. Cellis, ed., 1998) Academic Press; *Animal Cell Culture* (R.I. Freshney), ed., 1987); *Introduction to Cell and Tissue Culture* (J.P. Mather and P.E. Roberts, 1998) Plenum Press; *Cell and Tissue Culture: Laboratory Procedures* (A. Doyle, J.B. Griffiths, and D.G. Newell, eds., 1993-8) J. Wiley and Sons; *Handbook of Experimental Immunology* (D.M. Weir and C.C. Blackwell, eds.); *Gene Transfer Vectors for Mammalian Cells* (J.M. Miller and M.P. Calos, eds., 1987); *PCR: The Polymerase Chain Reaction*, (Mullis et al., eds., 1994); *Current Protocols in Immunology* (J.E. Coligan et al., eds., 1991); *Short Protocols*

in Molecular Biology (Wiley and Sons, 1999); *Immunobiology* (C.A. Janeway and P. Travers, 1997); *Antibodies* (P. Finch, 1997); *Antibodies: A Practical Approach* (D. Catty., ed., IRL Press, 1988-1989); *Monoclonal Antibodies: A Practical Approach* (P. Shepherd and C. Dean, eds., Oxford University Press, 2000); *Using Antibodies: A Laboratory Manual* (E. Harlow and D. Lane (Cold Spring Harbor Laboratory Press, 1999); and *The Antibodies* (M. Zanetti and J. D. Capra, eds., Harwood Academic Publishers, 1995).

Zika virus

[0058] Certain aspects of the present disclosure relate to at least one Zika virus (*e.g.*, a Zika virus comprising one or more non-human cell adaptation mutations) that may be useful in vaccines and/or immunogenic compositions including, without limitation, purified viruses, inactivated viruses, attenuated viruses, recombinant viruses, or purified and/or recombinant viral proteins for subunit vaccines.

[0059] Zika virus (ZIKV) is a mosquito-borne flavivirus first isolated from a sentinel rhesus monkey in the Zika Forest in Uganda in 1947. Since that time, isolations have been made from humans in both Africa and Asia, and more recently, the Americas. ZIKV is found in two (possibly three) lineages: an African lineage (possibly separate East and West African lineages) and an Asian lineage. Accordingly, examples of suitable Zika viruses of the present disclosure include, without limitation, viruses from the African and/or Asian lineages. In some embodiments, the Zika virus is an African lineage virus. In some embodiments, the Zika virus is an Asian lineage virus. Additionally, multiple strains within the African and Asian lineages of Zika virus have been previously identified. Any one or more suitable strains of Zika virus known in the art may be used in the present disclosure, including, for examples, strains Mr 766, ArD 41519, IbH 30656, P6-740, EC Yap, FSS13025, ArD 7117, ArD 9957, ArD 30101, ArD 30156, ArD 30332, HD 78788, ArD 127707, ArD 127710, ArD 127984, ArD 127988, ArD 127994, ArD 128000, ArD 132912, 132915, ArD 141170, ArD 142623, ArD 149917, ArD 149810, ArD 149938, ArD 157995, ArD 158084, ArD 165522, ArD 165531, ArA 1465, ArA 27101, ArA 27290, ArA 27106, ArA 27096, ArA 27407, ArA 27433, ArA 506/96, ArA 975-99, Ara 982-99, ArA 986-99, ArA 2718, ArB 1362, Nigeria68, Malaysia66, Kedougou84, Suriname, MR1429, PRVABC59, ECMN2007, DakAr41524, H/PF/2013, R103451, 103344, 8375, JMB-185, ZIKV/H, sapiens/Brazil/Natal/2015, SPH2015, ZIKV/Hu/Chiba/S36/2016, and/or Cuba2017.

[0060] In some embodiments, an example of a Zika virus genome sequence is set forth below as SEQ ID NO: 2:

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   1 gttgttgatc tgtgtgaatc agactgcgac agttcgagtt tgaagcgaaa gctagcaaca
  61 gtatcaacag gttttatttt ggatttggaa acgagagttt ctggtcatga aaaacccaaa
 121 aaagaaaacc ggaggattcc ggattgtcaa tatgctaaaa cgcggagtag cccgtgtgag
 181 ccccctttggg ggcttgaaga ggctgccagc cggacttctg ctgggtcatg ggcccatcag
 241 gatggtcctg gcgattctag ccttttttgag attcacggca atcaagccat cactgggtct
 301 catcaataga tggggttcag tggggaaaaa agaggctatg gaaacaataa agaagttcaa
 361 gaaagatctg gctgccatgc tgagaataat caatgctagg aaggagaaga agagacgagg
 421 cgcagatact agtgtcggaa ttgttggcct cctgctgacc acagctatgg cagcggaggt
 481 cactagacgt gggagtgcat actatatgta cttggacaga aacgatgctg gggaggccat
 541 atcttttcca accacattgg ggatgaataa gtgttatata cagatcatgg acttggaca
 601 catgtgtgat gccaccatga gctatgaatg ccctatgctg gatgaggggg tggaaccaga
 661 tgacgtcgat tgttggtgca acacgacgtc aacttgggtt gtgtacggaa cctgccatca
 721 caaaaaaggt gaagcacgga gatctagaag agctgtgacg ctcccctccc acttccaccag
 781 gaagctgcaa acgcggtcgc aaacctggtt ggaatcaaga gaatacacaa agcacttgat
 841 tagagtcgaa aattggatat tcaggaaccc tggcttcgcg ttagcagcag ctgccatcgc
 901 ttggctttctg ggaagctcaa cgagccaaaa agtcatatac ttggtcatga tactgctgat
 961 tgccccggca tacagcatca ggtgcatagg agtcagcaat agggactttg tggaaggtat
1021 gtcaggtggg acttgggttg atgttgtctt ggaacatgga ggttgtgtca ccgtaatggc
1081 acaggacaaa ccgactgtcg acatagagct ggttacaaca acagtcagca acatggcgga
1141 ggtaagacCC tactgctatg aggcatcaat atcagacatg gcttctgaca gccgctgccc
1201 aacacaaggt gaagccCacc tTgacaagca atcagacact caatatgtct gcaaaagaac
1261 gttagtggac agaggctggg gaaatggatg tggacttttt ggcaaaggga gcctggtgac
1321 atgcgctaag tttgcatgct ccaagaaaat gaccgggaag agcatccagc cagagaatct
1381 ggagtaccgg ataatgctgt cagttcatgg ctcccagcac agtgggatga tcgttaatga
1441 cacaggacat gaaactgatg agaatagagc gaaagttgag ataacgccca attcaccgag
1501 agccgaagcc accctggggg gttttggaag cctaggactt gattgtgaac cgaggacagg
1561 ccttgacctt tcagatttgt attacttgac tatgaataac aagcactggt tggttcacaa
1621 ggagtggctc cacgacattc cattaccttg gcacgctggg gcagacaccg gaactccaca
1681 ctggaacaac aaagaagcac tggtagagtt caaggacgca catgccaaaa ggcaaactgt
1741 cgtggttcta gggagtcaag aaggagcagt tcacacggcc cttgctggag cactggaggc
1801 tgagatggat ggtgcaaagg gaaggctgtc ctctggccac ttgaaatgtc gcctgaaaat
1861 ggataaactt agattgaagg gcgtgtcata ctccttgtgt actgcagcgt tcacattcac
1921 caagatcccg gctgaaacac tgcacgggac agtcacagtg gaggtacagt acgcagggac
1981 agatggacct tgcaaggttc cagctcagat ggcggtggac atgcaaactc tgaccccagt
2041 tgggaggctg ataaccgcta accccgtaat cactgaaagc actgagaact ctaagatgat
2101 gctggaactt gatccaccat ttggggactc ttacattgtc ataggagtcg gggagaagaa
2161 gatcacccac cactggcaca ggagtggcag caccattgga aaagcatttg aagccactgt
2221 gagaggtgcc aagagaatgg cagtcttggg agacacagcc tgggactttg gatcagttgg
2281 aggcgctctc aactcattgg gcaagggcat ccatcaaatt tttggagcag ctttcaaatc
2341 attgtttgga ggaatgtcct ggttctcaca aattctcatt ggaacgttgc tgatgtggtt
2401 gggtctgaac acaaagaatg gatctatttc ccttatgtgc ttggccttag ggggagtgtt
2461 gatcttctta tccacagccg tctctgctga tgtggggtgc tcggtggact tctcaaagaa
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2521 ggagacgaga tgcggtacag gggtgttcgt ctataacgac gttgaagcct ggagggacag
2581 gtacaagcac catcctgact ccccccgtag attggcagca gcagtcaagc aagcctggga
2641 agatggtatc tgcgggatct cctctgtttc aagaatggaa aacatcatgt ggagatcagt
2701 agaaggggag ctcaacgcaa tcctggaaga gaatggagtt caactgacgg tcgttgtggg
2761 atctgtaaaa aaccccatgt ggagaggtcc acagagattg cccgtgcctg tgaacgagct
2821 gccccacggc tggaaggctt gggggaaatc gtatttcgtc agagcagcaa agacaaataa
2881 cagctttgtc gtggatggtg acacactgaa ggaatgccca ctcaaacata gagcatggaa
2941 cagctttctt gtggaggatc atgggttcgg ggtatttcac actagtgtct ggctcaaggt
3001 tagagaagat tattcattag agtgtgatcc agccgttatt ggaacagctg ttaagggaaa
3061 ggaggctgta cacagtgatc taggctactg gattgagagt gagaagaatg acacatggag
3121 gctgaagagg gcccatctga tcgagatgaa aacatgtgaa tggccaaagt cccacacatt
3181 gtggacagat ggaatagaag agagtgatct gatcataccc aagtctttag ctgggccact
3241 cagccatcac aataccagag agggctacag gacccaaatg aaagggccat ggcacagtga
3301 agagcttgaa attcggtttg aggaatgccc aggcactaag gtccacgtgg aggaaacatg
3361 tggaacaaga ggaccatctc tgagatcaac cactgcaagc ggaagggtga tcgaggaatg
3421 gtgctgcagg gagtgcacaa tgcccccact gtcgttccgg gctaaagatg gctgttggta
3481 tggaatggag ataaggccca ggaaagaacc agaaagcaac ttagtaaggt caatggtgac
3541 tgcaggatca actgatcaca tggaccactt ctcccttgga gtgcttgtga tcctgctcat
3601 ggtgcaggaa gggctgaaga agagaatgac cacaaagatc atcataagca catcaatggc
3661 agtgctggta gctatgatcc tgggaggatt ttcaatgagt gacctggcta agcttgcaat
3721 tttgatgggt gccaccttcg cggaaatgaa cactggagga gatgtagctc atctggcgct
3781 gatagcggca ttcaaagtca gaccagcgtt gctggtatct ttcatcttca gagctaattg
3841 gacaccccgt gaaagcatgc tgctggcctt ggcctcgtgt ctttlgcaaa ctgcgatctc
3901 cgccttggaa ggcgacctga tggttctcat caatggtttt gctttggcct ggttggcaat
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6361 acacggagag aaaagagtgc tcaaaccgag gtggatggac gccagagttt gttcagatca
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[0061] In some embodiments, the Zika virus may comprise the genome sequence of GenBank Accession number KU501215.1. In some embodiments, the Zika virus is from strain PRVABC59. In some embodiments the genome sequence of GenBank Accession number KU501215.1 comprised the sequence of SEQ ID NO: 2. In some embodiments, the Zika virus may comprise a genomic sequence that has at least 70%, at least 71%, at least 72%, at least 73%, at least 74%,at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%,at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%,at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity with the sequence of SEQ ID NO: 2.

[0062] In some embodiments, the Zika virus may comprise a polypeptide encoded by the sequence of SEQ ID NO: 2. In some embodiments, the Zika virus may comprise a polypeptide having an amino acid sequence that has at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%,at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity with an amino acid sequence encoded by the sequence of SEQ ID NO: 2.

[0063] Accordingly, in some embodiments, Zika viruses of the present disclosure may be used in any of the vaccines and/or immunogenic compositions disclosed herein. For example, Zika viruses of the present disclosure may be used to provide one or more antigens useful for treating or preventing Zika virus infection in a subject in need thereof and/or inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof.

Viral Antigens

[0064] In some embodiments, the present disclosure relates to one or more antigens from any Zika virus described herein useful in vaccines and/or immunogenic compositions including, without limitation, purified viruses, inactivated viruses, attenuated viruses, recombinant viruses, or purified and/or recombinant viral proteins for subunit vaccines. In

some embodiments, the vaccines and/or immunogenic compositions include inactivated whole viruses.

[0065] Antigens of the present disclosure may be any substance capable of eliciting an immune response. Examples of suitable antigens include, but are not limited to, whole virus, attenuated virus, inactivated virus, proteins, polypeptides (including active proteins and individual polypeptide epitopes within proteins), glycopolypeptides, lipopolypeptides, peptides, polysaccharides, polysaccharide conjugates, peptide and non-peptide mimics of polysaccharides and other molecules, small molecules, lipids, glycolipids, and carbohydrates.

[0066] Antigens of the present disclosure may include at least one non-human cell adaptation mutation. Adaptation mutations may be generated by adapting a virus to growth in a particular cell line. For example, a cell may be transfected or electroporated with a virus, RNA transcribed from a virus (*e.g.,* an infectious virus), and/or RNA purified from a whole virus and passaged such that the virus and/or viral RNA replicates and its nucleic acid mutates. Nucleic acid mutations may be point mutations, insertion mutations, or deletion mutations. Nucleic acid mutations may lead to amino acid changes within viral proteins that facilitate growth of the virus in a non-human cell. Adaptation mutations may facilitate phenotypic changes in the virus, including altered plaque size, growth kinetics, temperature sensitivity, drug resistance, virulence, and virus yield in cell culture. These adaptive mutations may be useful in vaccine manufacture by increasing the speed, yield, and consistency of virus cultured in a cell line. Adaptive mutations may change (*e.g.,* enhance) immunogenicity of viral antigens by altering the structure of immunogenic epitopes. In addition, adaptive mutations may also increase the genetic stability of the virus and/or reduce or otherwise inhibit the development of undesirable mutations in the virus through multiple (*e.g.*, at least five, at least six, at least seven, at least eight, at least nine, at least 10, at least 11, at least 12, at least 13, at least 14, at least 15, at least 16, at least 17, at least 18, at least 19, at least 20, or more) passages.

[0067] Accordingly, in certain embodiments, antigens of the present disclosure include at least one non-human cell adaptation mutation. In certain embodiments, the adaptation mutation is a mutation of a viral antigen to a non-human cell. In some embodiments, the non-human cell is a mammalian cell. Any suitable mammalian cell known in the art may be used, including, without limitation, VERO cells (from monkey kidneys), LLC-MK2 cells

(from monkey kidneys), MDBK cells, MDCK cells, ATCC CCL34 MDCK (NBL2) cells, MDCK 33016 (deposit number DSM ACC 2219 as described in WO97/37001) cells, BHK21-F cells, HKCC cells, or Chinese hamster ovary cells (CHO cells). In some embodiments, the non-human cell is a monkey cell. In some embodiments, the monkey cell is from a Vero cell line. Any suitable Vero cell line known in the art may be used, including, without limitation, WHO Vero 10-87, ATCC CCL-81, Vero 76 (ATCC Accession No. CRL-1587), or Vero C1008 (ATCC Accession No. CRL-1586). In some embodiments, the Vero cell line is WHO Vero 10-87.

[0068] Zika viruses possess a positive sense, single-stranded RNA genome encoding both structural and nonstructural polypeptides. The genome also contains non-coding sequences at both the 5'- and 3'- terminal regions that play a role in virus replication. Structural polypeptides encoded by these viruses include, without limitation, capsid (C), precursor membrane (prM), and envelope (E). Non-structural (NS) polypeptides encoded by these viruses include, without limitation, NS1, NS2A, NS2B, NS3, NS4A, NS4B, and NS5.

[0069] In certain embodiments, antigens of the present disclosure may contain at least one (*e.g.,* at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, at least 10, at least 11, at least 12, at least 13, at least 14, at least 15, at least 16, at least 17, at least 18, at least 19, at least 20, *etc.*) non-human cell adaptation mutations within one or more (*e.g.,* one or more, two or more, three or more, four or more, five or more, six or more, seven or more, eight or more, nine or more, or all ten) viral antigens/polypeptides, including, without limitation, C, prM, E, NS1, NS2A, NS2B, NS3, NS4A, NS4B, and NS5. In some embodiments, antigens of the present disclosure include at least one non-human cell adaptation mutation in Zika virus Non-structural protein 1 (NS1). In some embodiments, antigens of the present disclosure include whole, inactivated virus that may contain at least one (*e.g.,* at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, at least 10, *etc.*) non-human cell adaptation mutations. In some embodiments, antigens of the present disclosure include whole, inactivated virus that may contain at least one non-human cell adaptation mutation in Zika virus Non-structural protein 1 (NS1).

[0070] In some embodiments, the at least one non-human cell adaptation mutation is within the NS1 polypeptide. The amino acid sequence of a wild-type, non-cell adapted NS1 polypeptide from an exemplary Zika virus strain is set forth as:

DVGCSVDFSKKETRCGTGVFVYNDVEAWRDRYKYHPDSPRRLAAAVKQAWEDGI
CGISSVSRMENIMWRSVEGELNAILEENGVQLTVVVGSVKNPMWRGPQRLPVPVN
ELPHGWKAWGKSYFVRAAKTNNSFVVDGDTLKECPLKHRAWNSFLVEDHGFGVF
HTSVWLKVREDYSLECDPAVIGTAVKGKEAVHSDLGYWIESEKNDTWRLKRAHLI
EMKTCEWPKSHTLWTDGIEESDLIIPKSLAGPLSHHNTREGYRTQMKGPWHSEELE
IRFEECPGTKVHVEETCGTRGPSLRSTTASGRVIEEWCCRECTMPPLSFRAKDGCW
YGMEIRPRKEPESNLVRSMVT (SEQ ID NO: 1).

[0071] In some embodiments, the amino acid sequence of the NS1 polypeptide has at least 80%, at least 81%, at least 82%, at least 83%, at least 84%,at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%,at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity with the sequence of SEQ ID NO: 1. In some embodiments, the amino acid sequence of the NS1 polypeptide may be from the amino acid sequence encoded by the sequence of GenBank Accession number KU501215.1 (SEQ ID NO: 2). In some embodiments, the amino acid sequence of the NS1 polypeptide may be amino acid positions 795 to 1145 of the amino acid sequence encoded by the sequence of GenBank Accession number KU501215.1. In some embodiments, amino acid sequence of the NS1 polypeptide may be from Zika virus strain PRVABC59.

[0072] In some embodiments, the at least one non-human cell adaptation mutation occurs at one or more amino acid positions within the NS1 polypeptide. In some embodiments, the mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1 when aligned to SEQ ID NO: 1 using a pairwise alignment algorithm. In some embodiments, the mutation at position 98 is a tryptophan to glycine substitution.

[0073] In some embodiments, antigens of the present disclosure contain at least one non-human cell adaptation mutation within the NS1 protein, and contain at least one mutation (*e.g.,* at least one adaptation mutation) within one or more of the C, prM, E, NS1, NS2A, NS2B, NS3, NS4A, NS4B, and NS5 viral proteins. In some embodiments, antigens of the present disclosure contain one or more non-human cell adaptation mutations within the NS1 protein, and do not contain at least one mutation (*e.g.,* at least one non-human cell adaptation mutation) within one or more of the C, prM, E, NS1, NS2A, NS2B, NS3, NS4A,

NS4B, and NS5 viral proteins. In some embodiments, antigens of the present disclosure contain at least one non-human cell adaptation mutation within the NS1 protein and do not contain at least one mutation (*e.g.*, at least one non-human cell adaptation mutation) within the envelope protein E. In some embodiments, antigens of the present disclosure include whole, inactivated virus that contains at least one non-human cell adaptation mutation in Zika virus Non-structural protein 1 (NS1), and do not include a mutation in Zika virus envelope protein E (Env).

[0074] In some embodiments, antigens of the present disclosure, such as Zika virus, contain at least one non-human cell adaptation mutation that enhances genetic stability as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments, antigens of the present disclosure, such as Zika virus, contain at least one non-human cell adaptation mutation that enhances viral replication as compared to a Zika virus lacking the at least one adaptation mutation. In some embodiments, antigens of the present disclosure, such as Zika virus, contain at least one non-human cell adaptation mutation reduces or otherwise inhibits the occurrence of undesirable mutations, such as within the envelope protein E (Env) of the Zika virus.

[0075] In the above embodiments of the present disclosure, an exemplary pairwise alignment algorithm is the Needleman-Wunsch global alignment algorithm, using default parameters (e.g. with Gap opening penalty=10.0, and with Gap extension penalty=0.5, using the EBLOSUM62 scoring matrix). This algorithm is conveniently implemented in the needle tool in the EMBOSS package.

[0076] In some embodiments, antigens of the present disclosure from a Zika virus may be used in any of the vaccines and immunogenic compositions of the present disclosure. For example, the antigens of the present disclosure may be useful for treating or preventing Zika virus infection in a subject in need thereof and/or inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof.

Production of Vaccines and Immunogenic Compositions

[0077] Other aspects of the present disclosure relate to Zika virus vaccines and immunogenic compositions containing one or more antigens of the present disclosure from at least one Zika virus. Such vaccines and immunogenic compositions may be useful, for example, for treating or preventing Zika virus infection in a subject in need thereof and/or

inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof. Vaccines and/or immunogenic compositions of the present disclosure may include, without limitation, purified viruses, inactivated viruses, attenuated viruses, recombinant viruses, purified and/or recombinant viral proteins for subunit vaccines. Vaccines and/or immunogenic compositions of the present disclosure may further include a purified antigen vaccine or immunogenic composition, a subunit vaccine or immunogenic composition, an inactivated whole virus vaccine or immunogenic composition, or an attenuated virus vaccine or immunogenic composition.

[0078] Production of vaccines and/or immunogenic compositions of the present disclosure includes growth of Zika virus, with antigens being prepared from the grown virus. Growth in cell culture is a method for preparing vaccines and/or immunogenic compositions of the present disclosure. Cells for viral growth may be cultured in suspension or in adherent conditions.

[0079] Cell lines suitable for growth of the at least one virus of the present disclosure are preferably of mammalian origin, and include but are not limited to: VERO cells (from monkey kidneys), horse, cow (e.g. MDBK cells), sheep, dog (e.g. MDCK cells from dog kidneys, ATCC CCL34 MDCK (NBL2) or MDCK 33016, deposit number DSM ACC 2219 as described in WO97/37001), cat, and rodent (e.g. hamster cells such as BHK21-F, HKCC cells, or Chinese hamster ovary cells (CHO cells)), and may be obtained from a wide variety of developmental stages, including for example, adult, neonatal, fetal, and embryo. In certain embodiments, the cells are immortalized (e.g. PERC.6 cells, as described in WO01/38362 and WO02/40665, and as deposited under ECACC deposit number 96022940). In preferred embodiments, mammalian cells are utilized, and may be selected from and/or derived from one or more of the following non-limiting cell types: fibroblast cells (*e.g.* dermal, lung), endothelial cells (*e.g.* aortic, coronary, pulmonary, vascular, dermal microvascular, umbilical), hepatocytes, keratinocytes, immune cells (*e.g.* T cell, B cell, macrophage, NK, dendritic), mammary cells (*e.g.* epithelial), smooth muscle cells (*e.g.* vascular, aortic, coronary, arterial, uterine, bronchial, cervical, retinal pericytes), melanocytes, neural cells (*e.g.* astrocytes), prostate cells (*e.g.* epithelial, smooth muscle), renal cells (*e.g.* epithelial, mesangial, proximal tubule), skeletal cells (*e.g.* chondrocyte, osteoclast, osteoblast), muscle cells (*e.g.* myoblast, skeletal, smooth, bronchial), liver cells, retinoblasts, and stromal cells. WO97/37000 and WO97/37001 describe production of

animal cells and cell lines that capable of growth in suspension and in serum free media and are useful in the production and replication of viruses.

[0080] Culture conditions for the above cell types are known and described in a variety of publications, or alternatively culture medium, supplements, and conditions may be purchased commercially, such as for example, as described in the catalog and additional literature of Cambrex Bioproducts (East Rutherford, N.J.).

[0081] In certain embodiments, the cells used in the methods described herein are cultured in serum free and/or protein free media. A medium is referred to as a serum-free medium in the context of the present disclosure in which there are no additives from serum of human or animal origin. Protein-free is understood to mean cultures in which multiplication of the cells occurs with exclusion of proteins, growth factors, other protein additives and non-serum proteins, but can optionally include proteins such as trypsin or other proteases that may be necessary for viral growth. The cells growing in such cultures naturally contain proteins themselves.

[0082] Known serum-free media include Iscove's medium, Ultra-CHO medium (BioWhittaker) or EX-CELL (JRH Bioscience). Ordinary serum-containing media include Eagle's Basal Medium (BME) or Minimum Essential Medium (MEM) (Eagle, Science, 130, 432 (1959)) or Dulbecco's Modified Eagle Medium (DMEM or EDM), which are ordinarily used with up to 10% fetal calf serum or similar additives. Optionally, Minimum Essential Medium (MEM) (Eagle, Science, 130, 432 (1959)) or Dulbecco's Modified Eagle Medium (DMEM or EDM) may be used without any serum containing supplement. Protein-free media like PF-CHO (JHR Bioscience), chemically-defined media like ProCHO 4CDM (BioWhittaker) or SMIF 7 (Gibco/BRL Life Technologies) and mitogenic peptides like Primactone, Pepticase or HyPep.TM. (all from Quest International) or lactalbumin hydrolysate (Gibco and other manufacturers) are also adequately known in the prior art. The media additives based on plant hydrolysates have the special advantage that contamination with viruses, mycoplasma or unknown infectious agents can be ruled out.

[0083] Cell culture conditions (temperature, cell density, pH value, etc.) are variable over a very wide range owing to the suitability of the cell line employed according to the present disclosure and can be adapted to the requirements of particular viral strains.

[0084] The method for propagating virus in cultured cells generally includes the steps of inoculating the cultured cells with the strain to be cultured, cultivating the infected cells for a desired time period for virus propagation, such as for example as determined by virus titer or antigen expression (e.g. between 24 and 168 hours after inoculation) and collecting the propagated virus. The cultured cells are inoculated with a virus (measured by PFU or TCID50) to cell ratio of 1:500 to 1:1, preferably 1:100 to 1:5. The virus is added to a suspension of the cells or is applied to a monolayer of the cells, and the virus is absorbed on the cells for at least 10 minutes, at least 20 minutes, at least 30 minutes, at least 40 minutes, at least 50 minutes, at least 60 minutes but usually less than 300 minutes at 25°C to 40°C, preferably 28°C to 38°C. The infected cell culture (*e.g.* monolayers) may be removed either by freeze-thawing or by enzymatic action to increase the viral content of the harvested culture supernatants. The harvested fluids are then either inactivated or stored frozen. Cultured cells may be infected at a multiplicity of infection ("MOI") of about 0.0001 to 10, preferably 0.002 to 5, more preferably to 0.001 to 2. Still more preferably, the cells are infected at an MOI of about 0.01. Infected cells may be harvested from 30 to 60 hours post infection, or 3 to 10 days post infection. In certain preferred embodiments, the cells are harvested 3 to 7 days post infection. More preferably, the cells are harvested 3 to 5 days post infection. In some embodiments, proteases (*e.g.,* trypsin) may be added during cell culture to allow viral release, and the proteases may be added at any suitable stage during the culture. Alternatively, in certain embodiments, the supernatant of infected cell cultures may be harvested and the virus may be isolated or otherwise purified from the supernatant.

[0085] The viral inoculum and the viral culture are preferably free from (*i.e.* will have been tested for and given a negative result for contamination by) herpes simplex virus, respiratory syncytial virus, parainfluenza virus 3, SARS coronavirus, adenovirus, rhinovirus, reoviruses, polyomaviruses, birnaviruses, circoviruses, and/or parvoviruses [WO2006/027698].

[0086] Where virus has been grown on a cell line then it is standard practice to minimize the amount of residual cell line DNA in the final vaccine, in order to minimize any oncogenic activity of the host cell DNA. Contaminating DNA can be removed during vaccine preparation using standard purification procedures *e.g.* chromatography, *etc.* Removal of residual host cell DNA can be enhanced by nuclease treatment *e.g.* by using a DNase. A convenient method for reducing host cell DNA contamination is disclosed in

references (Lundblad (2001) Biotechnology and Applied Biochemistry 34:195-197, *Guidance for Industry: Bioanalytical Method Validation.* U.S. Department of Health and Human Services Food and Drug Administration Center for Drug Evaluation and Research (CDER) Center for Veterinary Medicine (CVM). May 2001.) involving a two-step treatment, first using a DNase (*e.g.* Benzonase), which may be used during viral growth, and then a cationic detergent (*e.g.* CTAB), which may be used during virion disruption. Removal by β-propiolactone treatment can also be used.

Production of Antigens

[0087] Antigens of the present disclosure for use in vaccines and/or immunogenic compositions including, without limitation, purified viruses, inactivated viruses, attenuated viruses, recombinant viruses, or purified and/or recombinant viral proteins for subunit vaccines to treat and/or prevent Zika virus infection and/or induce an immune response, such as a protective immune response, against Zika virus, may be produced and/or purified or otherwise isolated by any suitable method known in the art. Antigens of the present disclosure may include, without limitation, whole virus, attenuated virus, inactivated virus, proteins, polypeptides (including active proteins and individual polypeptide epitopes within proteins), glycopolypeptides, lipopolypeptides, peptides, polysaccharides, polysaccharide conjugates, peptide and non-peptide mimics of polysaccharides and other molecules, small molecules, lipids, glycolipids, and carbohydrates produced, derived, purified, and/or otherwise isolated from a Zika virus. For example, suitable antigens may include, without limitation, structural polypeptides such as C, prM, and/or E, and non-structural polypeptides, such as NS1, NS2A, NS2B, NS3, NS4A, NS4B, and/or NS5 from Zika virus.

[0088] Antigen of the present disclosure may be synthesized chemically or enzymatically, produced recombinantly, isolated from a natural source, or a combination of the foregoing. In certain embodiments, antigens of the present disclosure are produced, purified, isolated, and/or derived from at least one Zika virus of the present disclosure. Antigens of the present disclosure may be purified, partially purified, or a crude extract. In some embodiments, antigens of the present disclosure are viruses, such as inactivated viruses, produced as described in the above section entitled "Production of Vaccines and Immunogenic Compositions."

[0089] In certain embodiments, one or more antigens of the present disclosure may be produced by culturing a non-human cell. Cell lines suitable for production of the one or more antigens of the present disclosure are preferably of mammalian origin, and include but are not limited to: VERO cells (from monkey kidneys), horse, cow (e.g. MDBK cells), sheep, dog (e.g. MDCK cells from dog kidneys, ATCC CCL34 MDCK (NBL2) or MDCK 33016, deposit number DSM ACC 2219 as described in WO97/37001), cat, and rodent (e.g. hamster cells such as BHK21-F, HKCC cells, or Chinese hamster ovary cells (CHO cells)), and may be obtained from a wide variety of developmental stages, including for example, adult, neonatal, fetal, and embryo. In certain embodiments, the cells are immortalized (e.g. PERC.6 cells, as described in WO01/38362 and WO02/40665, and as deposited under ECACC deposit number 96022940). In preferred embodiments, mammalian cells are utilized, and may be selected from and/or derived from one or more of the following non-limiting cell types: fibroblast cells (*e.g.* dermal, lung), endothelial cells (*e.g.* aortic, coronary, pulmonary, vascular, dermal microvascular, umbilical), hepatocytes, keratinocytes, immune cells (*e.g.* T cell, B cell, macrophage, NK, dendritic), mammary cells (*e.g.* epithelial), smooth muscle cells (*e.g.* vascular, aortic, coronary, arterial, uterine, bronchial, cervical, retinal pericytes), melanocytes, neural cells (*e.g.* astrocytes), prostate cells (*e.g.* epithelial, smooth muscle), renal cells (*e.g.* epithelial, mesangial, proximal tubule), skeletal cells (*e.g.* chondrocyte, osteoclast, osteoblast), muscle cells (*e.g.* myoblast, skeletal, smooth, bronchial), liver cells, retinoblasts, and stromal cells. WO97/37000 and WO97/37001 describe production of animal cells and cell lines that capable of growth in suspension and in serum free media and are useful in the production of viral antigens. In certain embodiments, the non-human cell is cultured in serum-free media.

[0090] Polypeptide antigens may be isolated from natural sources using standard methods of protein purification known in the art, including, but not limited to, liquid chromatography (*e.g.*, high performance liquid chromatography, fast protein liquid chromatography, *etc.*), size exclusion chromatography, gel electrophoresis (including one-dimensional gel electrophoresis, two-dimensional gel electrophoresis), affinity chromatography, or other purification technique. In many embodiments, the antigen is a purified antigen, *e.g.*, from about 50% to about 75% pure, from about 75% to about 85% pure, from about 85% to about 90% pure, from about 90% to about 95% pure, from about 95% to about 98% pure, from about 98% to about 99% pure, or greater than 99% pure.

[0091] One may employ solid phase peptide synthesis techniques, where such techniques are known to those of skill in the art. See Jones, The Chemical Synthesis of Peptides (Clarendon Press, Oxford) (1994). Generally, in such methods a peptide is produced through the sequential additional of activated monomeric units to a solid phase bound growing peptide chain.

[0092] Well-established recombinant DNA techniques can be employed for production of polypeptides, where, *e.g.*, an expression construct comprising a nucleotide sequence encoding a polypeptide is introduced into an appropriate host cell (*e.g.,* a eukaryotic host cell grown as a unicellular entity in *in vitro* cell culture, *e.g.,* a yeast cell, an insect cell, a mammalian cell, *etc.*) or a prokaryotic cell (*e.g.,* grown in *in vitro* cell culture), generating a genetically modified host cell; under appropriate culture conditions, the protein is produced by the genetically modified host cell.

[0093] Besides killed and attenuated virus immunogenic compositions, one can use a subunit immunogenic composition or other type of immunogenic composition which presents to the animal the antigenic components of Zika virus. The antigenic component may be a protein, glycoprotein, lipid-conjugated protein or glycoprotein, a modified lipid moiety, or other viral component which, when injected into a human, stimulates an immune response in the human such that the human develops protective immunity against Zika virus. For a subunit immunogenic composition, the virus can be cultured on mammalian cells, as described above. The cell culture can be homogenized and an immunogenic composition can be isolated by passage of the cell culture homogenate over the appropriate column or through the appropriate pore size filter or via centrifugation of the cell culture homogenate.

[0094] If the antigenic component is a protein, then one can isolate the nucleic acid which encodes that protein and generate an immunogenic composition that contains that isolated nucleic acid. The nucleic acid encoding the antigenic component can be placed on a plasmid downstream of a signal sequence of a eukaryotic promoter. That plasmid can contain one or more selectable markers and be transfected into an attenuated prokaryotic organism, such as Salmonella spp., Shigella spp., or other suitable bacteria. The bacteria can then be administered to the human so that the human can generate a protective immune response to the antigenic component. Alternatively, the nucleic acid encoding the antigenic component can be placed downstream of a prokaryotic promoter, have one or more

selectable markers, and be transfected into an attenuated prokaryotic organism such as Salmonella spp., Shigella spp., or other suitable bacteria. The bacteria can then be administered to the eukaryotic subject for which immune response to the antigen of interest is desired. See, for example, U.S. Pat. No. 6,500,419.

[0095] For a subunit immunogenic composition, the nucleic acid encoding a proteinaceous antigenic component of a Zika virus can be cloned into a plasmid such as those described in International Patent Application Publication Number WO 00/32047 (Galen) and International Patent Application Publication Number WO 02/083890 (Galen). Then the plasmid can be transfected into bacteria and the bacteria can produce the desired antigenic protein. One can isolate and purify the desired antigenic protein by a variety of methods described in both patent applications.

Virus Inactivation

[0096] Certain aspects of the present disclosure relate to Zika virus vaccines and immunogenic compositions containing one or more antigens from a Zika virus. Vaccines and/or immunogenic compositions of the present disclosure may include a purified virus, a whole virus, a recombinant virus, a live attenuated whole virus or, preferably, an inactivated whole virus, or subunits, polypeptides, and/or antigens from an inactivated virus. As such, certain embodiments of the present disclosure relate to Zika virus vaccines and/or immunogenic compositions containing one or more antigens from at least one inactivated Zika virus. Methods of inactivating or killing viruses to destroy their ability to infect mammalian cells are known in the art. Such methods include both chemical and physical means. Suitable means for inactivating a virus include, without limitation, treatment with an effective amount of one or more agents selected from detergents, formalin (also referred to herein as "formaldehyde"), beta-propiolactone (BPL), binary ethylamine (BEI), acetyl ethyleneimine, heat, electromagnetic radiation, x-ray radiation, gamma radiation, ultraviolet radiation (UV radiation),UV-A radiation, UV-B radiation, UV-C radiation, methylene blue, psoralen, carboxyfullerene (C60) and any combination of any thereof.

[0097] In certain embodiments of the present disclosure the at least one virus is chemically inactivated. Agents for chemical inactivation and methods of chemical inactivation are well-known in the art and described herein. In some embodiments, the at

least one virus is chemically inactivated with one or more of BPL, formalin, or BEI. In certain embodiments where the at least one virus is chemically inactivated with BPL, the virus may contain one or more modifications. In some embodiments, the one or more modifications may include a modified nucleic acid. In some embodiments, the modified nucleic acid is an alkylated nucleic acid. In other embodiments, the one or more modifications may include a modified polypeptide. In some embodiments, the modified polypeptide contains a modified amino acid residue including one or more of a modified cysteine, methionine, histidine, aspartic acid, glutamic acid, tyrosine, lysine, serine, and threonine. In certain embodiments where the at least one virus is chemically inactivated with formalin, the virus may contain one or more modifications. In some embodiments, the one or more modifications may include a modified polypeptide. In some embodiments, the one or more modifications may include a cross-linked polypeptide. In some embodiments where the at least one virus is chemically inactivated with formalin, the vaccine or immunogenic composition further includes formalin. In certain embodiments where the at least one virus is chemically inactivated with BEI, the virus may contain one or more modifications. In some embodiments, the one or more modifications may include a modified nucleic acid. In some embodiments, the modified nucleic acid is an alkylated nucleic acid.

[0098] In some embodiments where the at least one virus is chemically inactivated with formalin, any residual unreacted formalin may be neutralized with sodium metabisulfite, may be dialyzed out, and/or may be buffer exchanged to remove the residual unreacted formalin. In some embodiments, the sodium metabisulfite is added in excess. In some embodiments, the solutions may be mixed using a mixer, such as an in-line static mixer, and subsequently filtered or further purified (*e.g.,* using a cross flow filtrations system).

[0099] Certain embodiments of the present disclosure relate to a method for inactivating a Zika virus preparation. In some embodiments, the method involves (a) isolating the Zika virus preparation from one or more non-human cells that are used to produce the virus preparation and (b) treating the virus preparation with an effective amount of formalin. In certain embodiments, treating with an effective amount of formalin includes, without limitation, treating with formalin in an amount that ranges from about 0.001% v/v to about 3.0% v/v. For example, treating with an effective amount of formalin may include treating with formalin in an amount that ranges from about 0.001% to about

3.0% v/v, about 0.005% to about 2.0% v/v, or about 0.01% to about 1.0% v/v, or in an amount of about 0.001%, about 0.0025%, about 0.005%, about 0.0075%, about 0.01%, about 0.02%, about 0.03%, about 0.04%, about 0.05%, about 0.06%, about 0.07%, about 0.08%, about 0.09%, about 0.1%, about 0.2%, about 0.3%, about 0.4%, about 0.5%, about 0.6%, about 0.7%, about 0.8%, about 0.9%, about 1.0%, about 1.25%, about 1.5%, about 1.75%, about 2.0%, about 2.25%, about 2.5%, about 2.75%, or about 3.0% v/v.

[0100] In certain embodiments of the method, the Zika virus preparation is treated with formalin at a temperature that ranges from about 2°C to about 42°C. For example, the Zika virus preparation may be treated with formalin at a temperature that ranges from about 2°C to about 42°C, about 2°C to about 8°C, about 15°C to about 37°C, about 17°C to about 27°C, about 20°C to about 25°C, or at a temperature of about 2°C, about 4°C, about 8°C, about 10°C, about 15°C, about 17°C, about 18°C, about 19°C, about 20°C, about 21°C, about 22°C, about 23°C, about 24°C, about 25°C, about 26°C, about 27°C, about 28°C, about 29°C, about 30°C, about 37°C, or about 42°C. In some embodiments, the Zika virus preparation is treated with formalin at room temperature.

[0101] In some embodiments, the Zika virus preparation is treated with formalin for at last about 1 day. For example, the Zika virus preparation may be treated with formalin for at least about 1 day, at least about 2 days, at least about 3 days, at least about 4 days, at least about 5 days, at least about 6 days, at least about 7 days, at least about 8 days, at least about 9 das, at least about 10 days, at least about 11 days, at least about 12 days, at least about 13 days, at least about 14 days, at least about 15 days, at least about 16 days, at least about 17 days, at least about 18 days, at least about 19 days, at least about 20 days, at least about 21 days, at least about 22 days, at least about 23 days, at least about 24 days, at least about 25 days, at least about 26 days, at least about 27 days, at least about 28 days, at least about 29 days, at least about 30 days, or more. In some embodiments, the Zika virus preparation is treated with formalin for at least about 9 days. In some embodiments, the Zika virus preparation is treated with formalin for at least about 11 days. In some embodiments, the Zika virus preparation is treated with formalin for at least about 14 days. In some embodiments, the Zika virus preparation is treated with formalin for at least about 20 days. In some embodiments, the Zika virus preparation is treated with formalin for at least about 30 days.

[0102] In some embodiments, the method further involves neutralizing unreacted formalin with an effective amount of sodium metabisulfite. In some embodiments, the

effective amount of sodium metabisulfite ranges from about 0.01 mM to about 100 mM. For example, the sodium metabisulfite may be added at an effective concentration of from about 0.01 mM to about 100 mM, from about 0.1 mM to about 50 mM, from about 0.5 mM to about 20mM, or from about 1 mM to about 10 mM, or at a concentration of about 0.01mM, about 0.05mM, about 0.1mM, about 0.25mM, about 0.5mM, about 0.75mM, about 1mM, about 2mM, about 3mM, about 4mM, about 5mM, about 6mM, about 7mM, about 8mM, about 9mM, about 10mM, about 20mM, about 30mM about 40mM, about 50mM, about 75mM or about 100mM. In some embodiments, the formalin is neutralized with about 2mM sodium metabisulfite.

[0103] In some embodiments, the method involves (a) isolating the Zika virus preparation from one or more non-human cells that are used to produce the virus preparation; (b) treating the virus preparation with an effective amount of formalin; (c) neutralizing the virus preparation with an effective amount of sodium metabisulfite; and (d) purifying the neutralized virus preparation. Any method of purifying a virus preparation known in the art may be employed, including, without limitation, using cross flow filtration (CFF), multimodal chromatography, size exclusion chromatography, cation exchange chromatography, and/or anion exchange chromatography. In some embodiments, the neutralized virus preparation is purified by cross flow filtration (CFF). In some embodiments, the virus preparation is purified to a high degree in an amount that is about 70%, about 75%, about 80%, about 85%, about 90%, about 91%, about 92%, about 93%, about 94%, about 95% about 96%, about 97%, about 98%, about 99%, or more.

[0104] The vaccines and/or immunogenic compositions of the present disclosure containing one or more antigens from at least one inactivated Zika virus may be useful for treating or preventing Zika virus infection in a subject in need thereof and/or inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof.

Adjuvants

[0105] Other aspects of the present disclosure relate to Zika virus vaccines and/or immunogenic compositions containing one or more antigens from at least one Zika virus described herein in combination with one or more adjuvants. Such adjuvanted vaccines and/or immunogenic compositions of the present disclosure may be useful for treating or

preventing Zika virus infection in a subject in need thereof and/or inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof.

[0106] Various methods of achieving an adjuvant effect for vaccines are known and may be used in conjunction with the Zika virus vaccines and/or immunogenic compositions disclosed herein. General principles and methods are detailed in "The Theory and Practical Application of Adjuvants", 1995, Duncan E. S. Stewart-Tull (ed.), John Wiley & Sons Ltd, ISBN 0-471-95170-6, and also in "Vaccines: New Generation Immunological Adjuvants", 1995, Gregoriadis G et al. (eds.), Plenum Press, New York, ISBN 0-306-45283-9.

[0107] In some embodiments, a Zika virus vaccine or immunogenic composition includes the antigens and an adjuvant. Antigens may be in a mixture with at least one adjuvant, at a weight-based ratio of from about 10:1 to about 10^{10}:1 antigen:adjuvant, e.g., from about 10:1 to about 100:1, from about 100:1 to about 10^3:1, from about 10^3:1 to about 10^4:1, from about 10^4:1 to about 10^5:1, from about 10^5:1 to about 10^6:1, from about 10^6:1 to about 10^7:1, from about 10^7:1 to about 10^8:1, from about 10^8:1 to about 10^9:1, or from about 10^9:1 to about 10^{10}:1 antigen:adjuvant. One of skill in the art can readily determine the appropriate ratio through information regarding the adjuvant and routine experimentation to determine optimal ratios.

[0108] Exemplary adjuvants may include, but are not limited to, aluminum salts, calcium phosphate, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), MLA derivatives, synthetic lipid A, lipid A mimetics or analogs, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions (oil emulsions), chitosan, vitamin D, stearyl or octadecyl tyrosine, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and Incomplete Freund's Adjuvant (IFA). In some embodiments, the adjuvant is an aluminum salt.

[0109] In some embodiments, the adjuvant includes at least one of alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85. In some embodiments, aluminum salt adjuvants of the present disclosure have been found to increase adsorption of the antigens of the Zika virus vaccines and/or immunogenic compositions of

the present disclosure. Accordingly, in some embodiments, at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 91%, at least about 92%, at least about 93%, at least about 94%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% of the antigen is adsorbed to the aluminum salt adjuvant.

[0110] In some embodiments, the vaccine or immunogenic composition includes an aluminum salt adjuvant (*e.g.,* alum) from about 1 μg to about 500 μg, about 5 μg to about 500 μg, about 10 μg to about 500 μg, about 15 μg to about 500 μg, about 25 μg to about 500 μg, about 50 μg to about 500 μg, about 75 μg to about 500 μg, about 100 μg to about 500 μg, about 125 μg to about 500 μg, about 150 μg to about 500 μg, about 175 μg to about 500 μg, about 200 μg to about 500 μg, about 225 μg to about 500 μg, about 250 μg to about 500 μg, about 275 μg to about 500 μg, about 300 μg to about 500 μg, about 325 μg to about 500 μg, about 350 μg to about 500 μg, about 375 μg to about 500 μg, about 400 μg to about 500 μg, about 425 μg to about 500 μg, about 450 μg to about 500 μg, about 475 μg to about 500 μg, about 1 μg to about 475 μg, about 5 μg to about 475 μg, about 10 μg to about 475 μg, about 15 μg to about 475 μg, about 25 μg to about 475 μg, about 50 μg to about 475 μg, about 75 μg to about 475 μg, about 100 μg to about 475 μg, about 125 μg to about 475 μg, about 150 μg to about 475 μg, about 175 μg to about 475 μg, about 200 μg to about 475 μg, about 225 μg to about 475 μg, about 250 μg to about 475 μg, about 275 μg to about 475 μg, about 300 μg to about 475 μg, about 325 μg to about 475 μg, about 350 μg to about 475 μg, about 375 μg to about 475 μg, about 400 μg to about 475 μg, about 425 μg to about 475 μg, about 450 μg to about 475 μg, about 1 μg to about 450 μg, about 5 μg to about 450 μg, about 10 μg to about 450 μg, about 15 μg to about 450 μg, about 25 μg to about 450 μg, about 50 μg to about 450 μg, about 75 μg to about 450 μg, about 100 μg to about 450 μg, about 125 μg to about 450 μg, about 150 μg to about 450 μg, about 175 μg to about 450 μg, about 200 μg to about 450 μg, about 225 μg to about 450 μg, about 250 μg to about 450 μg, about 275 μg to about 450 μg, about 300 μg to about 450 μg, about 325 μg to about 450 μg, about 350 μg to about 450 μg, about 375 μg to about 450 μg, about 400 μg to about 450 μg, about 425 μg to about 450 μg, about 1 μg to about 425 μg, about 5 μg to about 425 μg, about 10 μg to about 425 μg, about 15 μg to about 425 μg, about 25 μg to about 425 μg, about 50 μg to about 425 μg, about 75 μg to about 425 μg, about 100 μg to about 425 μg, about 125 μg to about 425 μg, about 150 μg to about 425 μg, about 175 μg to about 425 μg, about 200 μg to about 425 μg, about 225 μg to about 425 μg, about 250 μg to about 425 μg, about 275 μg to about 425

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300 µg, about 325 µg, about 350 µg, about 375 µg, about 400 µg, about 425 µg, about 450 µg, about 475 µg, or about 500 µg per dose.

[0111] Certain embodiments of the present disclosure include a method for preparing an adjuvanted Zika virus vaccine or immunogenic composition, which involves (a) mixing the vaccine or immunogenic composition with an aluminum salt adjuvant, with the vaccine or immunogenic composition including one or more antigens from at least one Zika virus described herein and (b) incubating the mixture under suitable conditions for a period of time that ranges from about 1 hour to about 24 hours (*e.g.*, about 16 hours to about 24 hours), with at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 91%, at least about 92%, at least about 93%, at least about 94%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% of the antigen adsorbed to the aluminum salt adjuvant. In certain embodiments of the method, the at least one Zika virus is a Zika virus comprising a non-human cell adaptation mutation (*e.g.*, a non-human cell adaptation mutation in protein NS1 such as a Trp98Gly mutation). In some embodiments of the method, the mixture is incubated at a temperature that ranges from about 2°C to about 8°C. In some embodiments of the method, the mixture is incubated under constant mixing using any suitable mixer known in the art. In some embodiments of the method, the mixture is incubated at pH that ranges in value from about 6.5 to about 8.5, from about 6.5 to about 8, from about 6.8 to about 7.8, from about 6.9 to about 7.6, from about 7 to about 7.5, from about 6.8 to about 8.5, from about 6.9 to about 8.5, or from about 7 to about 8.5. In certain preferred embodiments, the mixture is incubated at a neutral pH. In some embodiments of the method, the aluminum salt adjuvant is selected from alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85.

[0112] Monophosphoryl Lipid A (MLA), a non-toxic derivative of lipid A from Salmonella, is a potent TLR-4 agonist that has been developed as a vaccine adjuvant (Evans *et al.* 2003). In pre-clinical murine studies intranasal MLA has been shown to enhance secretory, as well as systemic, humoral responses (Baldridge *et al.* 2000; Yang *et al.* 2002). It has also been proven to be safe and effective as a vaccine adjuvant in clinical studies of greater than 120,000 patients (Baldrick *et al.*, 2002; 2004). MLA stimulates the induction of innate immunity through the TLR-4 receptor and is thus capable of eliciting nonspecific immune responses against a wide range of infectious pathogens, including both gram negative and gram positive bacteria, viruses, and parasites (Baldrick *et al.* 2004; Persing *et al.*

2002). Inclusion of MLA in intranasal formulations should provide rapid induction of innate responses, eliciting nonspecific immune responses from viral challenge while enhancing the specific responses generated by the antigenic components of the vaccine.

[0113] Accordingly, in one embodiment, the present disclosure provides a composition comprising monophosphoryl lipid A (MLA), 3 De-O-acylated monophosphoryl lipid A (3D-MLA), or a derivative thereof as an enhancer of adaptive and innate immunity. Chemically 3D-MLA is a mixture of 3 De-O-acylated monophosphoryl lipid A with 4, 5 or 6 acylated chains. A preferred form of 3 De-O-acylated monophosphoryl lipid A is disclosed in European Patent 0 689 454 B1 (SmithKline Beecham Biologicals SA). In another embodiment, the present disclosure provides a composition comprising synthetic lipid A, lipid A mimetics or analogs, such as BioMira's PET Lipid A, or synthetic derivatives designed to function like TLR-4 agonists.

[0114] Additional exemplary adjuvants include, without limitation, polypeptide adjuvants that may be readily added to the antigens described herein by co-expression with the polypeptide components or fusion with the polypeptide components to produce chimeric polypeptides. Bacterial flagellin, the major protein constituent of flagella, is an adjuvant which has received increasing attention as an adjuvant protein because of its recognition by the innate immune system by the toll-like receptor TLR5 (65). Flagellin signaling through TLR5 has effects on both innate and adaptive immune functions by inducing DC maturation and migration as well as activation of macrophages, neutrophils, and intestinal epithelial cells resulting in production of pro-inflammatory mediators (66-72).

[0115] TLR5 recognizes a conserved structure within flagellin monomers that is unique to this protein and is required for flagellar function, precluding its mutation in response to immunological pressure (73). The receptor is sensitive to a 100 fM concentration but does not recognize intact filaments. Flagellar disassembly into monomers is required for binding and stimulation.

[0116] As an adjuvant, flagellin has potent activity for induction of protective responses for heterologous antigens administered either parenterally or intranasally and adjuvant effects for DNA vaccines have also been reported. A Th2 bias is observed when flagellin is employed which would be appropriate for a respiratory virus such as influenza but no evidence for IgE induction in mice or monkeys has been observed. In addition, no local or

systemic inflammatory responses have been reported following intranasal or systemic administration in monkeys. The Th2 character of responses elicited following use of flagellin is somewhat surprising since flagellin signals through TLR5 in a MyD88-dependent manner and all other MyD88-dependent signals through TLRs have been shown to result in a Th1 bias. Importantly, pre-existing antibodies to flagellin have no appreciable effect on adjuvant efficacy making it attractive as a multi-use adjuvant.

[0117] A common theme in many recent intranasal vaccine trials is the use of adjuvants and/or delivery systems to improve vaccine efficacy. In one such study an influenza H3 vaccine containing a genetically detoxified *E. coli* heat-labile enterotoxin adjuvant (LT R192G) resulted in heterosubtypic protection against H5 challenge but only following intranasal delivery. Protection was based on the induction of cross neutralizing antibodies and demonstrated important implications for the intranasal route in development of new vaccines.

[0118] Cytokines, colony-stimulating factors (e.g., GM-CSF, CSF, and the like); tumor necrosis factor; interleukin-2, -7, -12, interferons and other like growth factors, may also be used as adjuvants as they may be readily included in the Zika virus vaccines or immunogenic compositions by admixing or fusion with the polypeptide component.

[0119] In some embodiments, the Zika virus vaccine and/or immunogenic compositions disclosed herein may include other adjuvants that act through a Toll-like receptor such as a nucleic acid TLR9 ligand comprising a 5'-TCG-3' sequence; an imidazoquinoline TLR7 ligand; a substituted guanine TLR7/8 ligand; other TLR7 ligands such as Loxoribine, 7-deazadeoxyguanosine, 7-thia-8-oxodeoxyguanosine, Imiquimod (R-837), and Resiquimod (R-848).

[0120] Certain adjuvants facilitate uptake of the vaccine molecules by APCs, such as dendritic cells, and activate these. Non-limiting examples are selected from the group consisting of an immune targeting adjuvant; an immune modulating adjuvant such as a toxin, a cytokine, and a mycobacterial derivative; an oil formulation; a polymer; a micelle forming adjuvant; a saponin; an immunostimulating complex matrix (ISCOM matrix); a particle; DDA; aluminum adjuvants; DNA adjuvants; MLA; and an encapsulating adjuvant.

[0121] Additional examples of adjuvants include agents such as aluminum salts such as hydroxide or phosphate (alum), commonly used as 0.05 to 0.1 percent solution in buffered saline (see, e.g., Nicklas (1992) Res. Immunol. 143:489-493), admixture with synthetic

polymers of sugars (e.g. Carbopol®) used as 0.25 percent solution, aggregation of the protein in the vaccine by heat treatment with temperatures ranging between 70° to 101°C for 30 second to 2 minute periods respectively and also aggregation by means of cross-linking agents are possible. Aggregation by reactivation with pepsin treated antibodies (Fab fragments) to albumin, mixture with bacterial cells such as C. parvum or endotoxins or lipopolysaccharide components of gram-negative bacteria, emulsion in physiologically acceptable oil vehicles such as mannide mono-oleate (Aracel A) or emulsion with 20 percent solution of a perfluorocarbon (Fluosol-DA) used as a block substitute may also be employed. Admixture with oils such as squalene and IFA may also be used.

[0122] DDA (dimethyldioctadecylammonium bromide) is an interesting candidate for an adjuvant, but also Freund's complete and incomplete adjuvants as well as quillaja saponins such as QuilA and QS21 are interesting. Further possibilities include poly[di(earboxylatophenoxy)phosphazene (PCPP) derivatives of lipopolysaccharides such as monophosphoryl lipid A (MLA), muramyl dipeptide (MDP) and threonyl muramyl dipeptide (tMDP). The lipopolysaccharide based adjuvants may also be used for producing a predominantly Th1-type response including, for example, a combination of monophosphoryl lipid A, such as 3-de-O-acylated monophosphoryl lipid A, together with an aluminum salt.

[0123] Liposome formulations are also known to confer adjuvant effects, and therefore liposome adjuvants may be used in conjunction with the Zika virus vaccines and/or immunogenic compositions.

[0124] Immunostimulating complex matrix type (ISCOM® matrix) adjuvants may also be used with the Zika virus vaccine antigens and immunogenic compositions, especially since it has been shown that this type of adjuvants are capable of up-regulating MHC Class II expression by APCs. An ISCOM matrix consists of (optionally fractionated) saponins (triterpenoids) from Quillaja saponaria, cholesterol, and phospholipid. When admixed with the immunogenic protein such as the Zika virus vaccine or immunogenic composition antigens, the resulting particulate formulation is what is known as an ISCOM particle where the saponin may constitute 60-70% w/w, the cholesterol and phospholipid 10-15% w/w, and the protein 10-15% w/w. Details relating to composition and use of immunostimulating complexes can for example be found in the above-mentioned text-books dealing with adjuvants, but also Morein B et al., 1995, Clin. Immunother. 3: 461-475 as well as Barr I G

and Mitchell G F, 1996, Immunol. and Cell Biol. 74: 8-25 provide useful instructions for the preparation of complete immunostimulating complexes.

[0125] The saponins, whether or not in the form of iscoms, that may be used in the adjuvant combinations with the Zika virus vaccines and immunogenic compositions disclosed herein include those derived from the bark of Quillaja Saponaria Molina, termed Quil A, and fractions thereof, described in U.S. Pat. No. 5,057,540 and "Saponins as vaccine adjuvants", Kensil, C. R., Crit Rev Ther Drug Carrier Syst, 1996, 12 (1-2):1-55; and EP 0 362 279 B1. Exemplary fractions of Quil A are QS21, QS7, and QS17.

[0126] β-Escin is another hemolytic saponins for use in the adjuvant compositions of the Zika virus vaccines and/or immunogenic compositions. Escin is described in the Merck index (12th ed: entry 3737) as a mixture of saponins occurring in the seed of the horse chestnut tree, Lat: Aesculus hippocastanum. Its isolation is described by chromatography and purification (Fiedler, Arzneimittel-Forsch. 4, 213 (1953)), and by ion-exchange resins (Erbring et al., U.S. Pat. No. 3,238,190). Fractions of escin have been purified and shown to be biologically active (Yoshikawa M, et al. (Chem Pharm Bull (Tokyo) 1996 August; 44(8):1454-1464)). β-escin is also known as aescin.

[0127] Another hemolytic saponin for use in the Zika virus vaccines and/or immunogenic compositions is Digitonin. Digitonin is described in the Merck index (12^{th} Edition, entry 3204) as a saponin, being derived from the seeds of Digitalis purpurea and purified according to the procedure described Gisvold et al., J.Am.Pharm.Assoc., 1934, 23, 664; and Ruhenstroth-Bauer, Physiol.Chem., 1955, 301, 621. Its use is described as being a clinical reagent for cholesterol determination.

[0128] Another interesting possibility of achieving adjuvant effect is to employ the technique described in Gosselin et al., 1992. In brief, the presentation of a relevant antigen such as an antigen in a Zika virus vaccine and/or immunogenic composition of the present disclosure can be enhanced by conjugating the antigen to antibodies (or antigen binding antibody fragments) against the F_C receptors on monocytes/macrophages. Especially conjugates between antigen and anti-F_CRI have been demonstrated to enhance immunogenicity for the purposes of vaccination. The antibody may be conjugated to the Zika virus vaccine or immunogenic composition antigens after generation or as a part of the generation including by expressing as a fusion to any one of the polypeptide components of

the Zika virus vaccine and/or immunogenic composition antigens. Other possibilities involve the use of the targeting and immune modulating substances (*e.g.,* cytokines). In addition, synthetic inducers of cytokines such as poly I:C may also be used.

[0129] Suitable mycobacterial derivatives may be selected from the group consisting of muramyl dipeptide, complete Freund's adjuvant, RIBI, (Ribi ImmunoChem Research Inc., Hamilton, Mont.) and a diester of trehalose such as TDM and TDE.

[0130] Examples of suitable immune targeting adjuvants include CD40 ligand and CD40 antibodies or specifically binding fragments thereof (cf. the discussion above), mannose, a Fab fragment, and CTLA-4.

[0131] Examples of suitable polymer adjuvants include a carbohydrate such as dextran, PEG, starch, mannan, and mannose; a plastic polymer; and latex such as latex beads.

[0132] Yet another interesting way of modulating an immune response is to include the immunogen (optionally together with adjuvants and pharmaceutically acceptable carriers and vehicles) in a "virtual lymph node" (VLN) (a proprietary medical device developed by ImmunoTherapy, Inc., 360 Lexington Avenue, New York, N.Y. 10017-6501). The VLN (a thin tubular device) mimics the structure and function of a lymph node. Insertion of a VLN under the skin creates a site of sterile inflammation with an upsurge of cytokines and chemokines. T- and B-cells as well as APCs rapidly respond to the danger signals, home to the inflamed site and accumulate inside the porous matrix of the VLN. It has been shown that the necessary antigen dose required to mount an immune response to an antigen is reduced when using the VLN, and that immune protection conferred by vaccination using a VLN surpassed conventional immunization using Ribi as an adjuvant. The technology is described briefly in Gelber C et al., 1998, "Elicitation of Robust Cellular and Humoral Immune Responses to Small Amounts of Immunogens Using a Novel Medical Device Designated the Virtual Lymph Node", in: "From the Laboratory to the Clinic, Book of Abstracts, Oct. 12-15, 1998, Seascape Resort, Aptos, Calif."

[0133] Oligonucleotides may be used as adjuvants in conjunction with the Zika virus vaccine and/or immunogenic composition antigens and may contain two or more dinucleotide CpG motifs separated by at least three or more or even at least six or more nucleotides. CpG-containing oligonucleotides (in which the CpG dinucleotide is unmethylated) induce a

predominantly Th1 response. Such oligonucleotides are well known and are described, for example, in WO 96/02555, WO 99/33488 and U.S. Pat. Nos. 6,008,200 and 5,856,462.

[0134] Such oligonucleotide adjuvants may be deoxynucleotides. In certain embodiments, the nucleotide backbone in the oligonucleotide is phosphorodithioate, or a phosphorothioate bond, although phosphodiester and other nucleotide backbones such as PNA including oligonucleotides with mixed backbone linkages may also be used. Methods for producing phosphorothioate oligonucleotides or phosphorodithioate are described in U.S. Pat. No. 5,666,153, U.S. Pat. No. 5,278,302 and W095/26204.

[0135] Exemplary oligonucleotides have the following sequences. The sequences may contain phosphorothioate modified nucleotide backbones:

(SEQ ID NO: 3) OLIGO 1: TCC ATG ACG TTC CTG ACG TT (CpG 1826);
(SEQ ID NO: 4) OLIGO 2: TCT CCC AGC GTG CGC CAT (CpG 1758);
(SEQ ID NO: 5) OLIGO 3: ACC GAT GAC GTC GCC GGT GAC GGC ACC ACG;
(SEQ ID NO: 6) OLIGO 4: TCG TCG TTT TGT CGT TTT GTC GTT (CpG 2006); and
(SEQ ID NO: 7) OLIGO 5: TCC ATG ACG TTC CTG ATG CT (CpG 1668)

[0136] Alternative CpG oligonucleotides include the above sequences with inconsequential deletions or additions thereto. The CpG oligonucleotides as adjuvants may be synthesized by any method known in the art (e.g., EP 468520). For example, such oligonucleotides may be synthesized utilizing an automated synthesizer. Such oligonucleotide adjuvants may be between 10-50 bases in length. Another adjuvant system involves the combination of a CpG-containing oligonucleotide and a saponin derivative particularly the combination of CpG and QS21 is disclosed in WO 00/09159.

[0137] Many single or multiphase emulsion systems have been described. One of skill in the art may readily adapt such emulsion systems for use with a Zika virus vaccine and/or immunogenic composition antigens so that the emulsion does not disrupt the antigen's structure. Oil in water emulsion adjuvants per se have been suggested to be useful as adjuvant compositions (EPO 399 843B), also combinations of oil in water emulsions and other active agents have been described as adjuvants for vaccines (WO 95/17210; WO 98/56414; WO 99/12565; WO 99/11241). Other oil emulsion adjuvants have been described, such as water

in oil emulsions (U.S. Pat. No. 5,422,109; EP 0 480 982 B2) and water in oil in water emulsions (U.S. Pat. No. 5,424,067; EP 0 480 981 B).

[0138] The oil emulsion adjuvants for use with the Zika virus vaccines and/or immunogenic compositions described herein may be natural or synthetic, and may be mineral or organic. Examples of mineral and organic oils will be readily apparent to one skilled in the art.

[0139] In order for any oil in water composition to be suitable for human administration, the oil phase of the emulsion system may include a metabolizable oil. The meaning of the term metabolizable oil is well known in the art. Metabolizable can be defined as "being capable of being transformed by metabolism" (Dorland's Illustrated Medical Dictionary, W.B. Sanders Company, 25th edition (1974)). The oil may be any vegetable oil, fish oil, animal oil or synthetic oil, which is not toxic to the recipient and is capable of being transformed by metabolism. Nuts (such as peanut oil), seeds, and grains are common sources of vegetable oils. Synthetic oils may also be used and can include commercially available oils such as NEOBEE® and others. Squalene (2,6,10,15,19,23-Hexamethyl-2,6,10,14,18,22-tetracosahexaene) is an unsaturated oil which is found in large quantities in shark-liver oil, and in lower quantities in olive oil, wheat germ oil, rice bran oil, and yeast, and may be used with the Zika virus vaccine and/or immunogenic compositions. Squalene is a metabolizable oil virtue of the fact that it is an intermediate in the biosynthesis of cholesterol (Merck index, 10th Edition, entry no.8619).

[0140] Exemplary oil emulsions are oil in water emulsions, and in particular squalene in water emulsions.

[0141] In addition, the oil emulsion adjuvants for use with the Zika virus vaccine and/or immunogenic compositions may include an antioxidant, such as the oil α-tocopherol (vitamin E, EP 0 382 271 B1).

[0142] WO 95/17210 and WO 99/11241 disclose emulsion adjuvants based on squalene, α-tocopherol, and TWEEN 80 (TM), optionally formulated with the immunostimulants QS21 and/or 3D-MLA. WO 99/12565 discloses an improvement to these squalene emulsions with the addition of a sterol into the oil phase. Additionally, a triglyceride, such as tricaprylin (C27H50O6), may be added to the oil phase in order to stabilize the emulsion (WO 98/56414).

[0143] The size of the oil droplets found within the stable oil in water emulsion may be less than 1 micron, may be in the range of substantially 30-600 nm, substantially around 30-500 nm in diameter, or substantially 150-500 nm in diameter, and in particular about 150 nm in diameter as measured by photon correlation spectroscopy. In this regard, 80% of the oil droplets by number may be within these ranges, more than 90% or more than 95% of the oil droplets by number are within the defined size ranges. The amounts of the components present in oil emulsions are conventionally in the range of from 2 to 10% oil, such as squalene; and when present, from 2 to 10% alpha tocopherol; and from 0.3 to 3% surfactant, such as polyoxyethylene sorbitan monooleate. The ratio of oil: alpha tocopherol may be equal or less than 1 as this provides a more stable emulsion. SPAN 85 (TM) may also be present at a level of about 1%. In some cases it may be advantageous that the Zika virus vaccines and/or immunogenic compositions disclosed herein will further contain a stabilizer.

[0144] The method of producing oil in water emulsions is well known to one skilled in the art. Commonly, the method includes the step of mixing the oil phase with a surfactant such as a PBS/TWEEN80® solution, followed by homogenization using a homogenizer, it would be clear to one skilled in the art that a method comprising passing the mixture twice through a syringe needle would be suitable for homogenizing small volumes of liquid. Equally, the emulsification process in microfluidizer (M110S microfluidics machine, maximum of 50 passes, for a period of 2 minutes at maximum pressure input of 6 bar (output pressure of about 850 bar)) could be adapted by one skilled in the art to produce smaller or larger volumes of emulsion. This adaptation could be achieved by routine experimentation comprising the measurement of the resultant emulsion until a preparation was achieved with oil droplets of the required diameter.

[0145] Alternatively the Zika virus vaccines and/or immunogenic compositions may be combined with vaccine vehicles composed of chitosan (as described above) or other polycationic polymers, polylactide and polylactide-coglycolide particles, poly-N-acetyl glucosamine-based polymer matrix, particles composed of polysaccharides or chemically modified polysaccharides, liposomes and lipid-based particles, particles composed of glycerol monoesters, etc. The saponins may also be formulated in the presence of cholesterol to form particulate structures such as liposomes or ISCOMs. Furthermore, the saponins may be formulated together with a polyoxyethylene ether or ester, in either a non-particulate

solution or suspension, or in a particulate structure such as a paucilamelar liposome or ISCOM.

[0146] Additional illustrative adjuvants for use in the Zika virus vaccines and/or immunogenic compositions as described herein include SAF (Chiron, Calif., United States), MF-59 (Chiron, see, e.g., Granoff et al. (1997) Infect Immun. 65 (5):1710-1715), the SBAS series of adjuvants (e.g., SB-AS2 (an oil-in-water emulsion containing MLA and QS21); SBAS-4 (adjuvant system containing alum and MLA), available from SmithKline Beecham, Rixensart, Belgium), Detox (Enhanzyn®) (GlaxoSmithKline), RC-512, RC-522, RC-527, RC-529, RC-544, and RC-560 (GlaxoSmithKline) and other aminoalkyl glucosaminide 4-phosphates (AGPs), such as those described in pending U.S. patent application Ser. Nos. 08/853,826 and 09/074,720.

[0147] Other examples of adjuvants include, but are not limited to, Hunter's TiterMax® adjuvants (CytRx Corp., Norcross, Ga.); Gerbu adjuvants (Gerbu Biotechnik GmbH, Gaiberg, Germany); nitrocellulose (Nilsson and Larsson (1992) Res. Immunol. 143:553-557); alum (e.g., aluminum hydroxide, aluminum phosphate) emulsion based formulations including mineral oil, non-mineral oil, water-in-oil or oil-in-water emulsions, such as the Seppic ISA series of Montamide adjuvants (e.g., ISA-51, ISA-57, ISA-720, ISA-151, etc.; Seppic, Paris, France); and PROVAX® (IDEC Pharmaceuticals); OM-174 (a glucosamine disaccharide related to lipid A); Leishmania elongation factor; non-ionic block copolymers that form micelles such as CRL 1005; and Syntex Adjuvant Formulation. See, e.g., O'Hagan et al. (2001) Biomol Eng. 18(3):69-85; and "Vaccine Adjuvants: Preparation Methods and Research Protocols" D. O'Hagan, ed. (2000) Humana Press.

[0148] Other exemplary adjuvants include adjuvant molecules of the general formula:

$$HO(CH_2CH_2O)_n\text{-}A\text{-}R, \quad (I)$$

where, n is 1-50, A is a bond or --C(O)--, R is $C_{1\text{-}50}$ alkyl or Phenyl $C_{1\text{-}50}$ alkyl.

[0149] One embodiment consists of a vaccine formulation comprising a polyoxyethylene ether of general formula (I), where n is between 1 and 50, 4-24, or 9; the R component is $C_{1\text{-}50}$, C_4-C_{20} alkyl, or C_{12} alkyl, and A is a bond. The concentration of the polyoxyethylene ethers should be in the range 0.1-20%, from 0.1-10%, or in the range 0.1-1%. Exemplary polyoxyethylene ethers are selected from the following group: polyoxyethylene-9-lauryl

ether, polyoxyethylene-9-steoryl ether, polyoxyethylene-8-steoryl ether, polyoxyethylene-4-lauryl ether, polyoxyethylene-35-lauryl ether, and polyoxyethylene-23-lauryl ether. Polyoxyethylene ethers such as polyoxyethylene lauryl ether are described in the Merck index (12th edition: entry 7717). These adjuvant molecules are described in WO 99/52549.

[0150] The polyoxyethylene ether according to the general formula (I) above may, if desired, be combined with another adjuvant. For example, an adjuvant combination may include the CpG as described above.

[0151] Further examples of suitable pharmaceutically acceptable excipients for use with the Zika virus vaccines and/or immunogenic compositions disclosed herein include water, phosphate buffered saline, isotonic buffer solutions.

Formulations of Vaccines and/or immunogenic compositions

[0152] Further aspects of the present disclosure relate to formulations of vaccines and/or immunogenic compositions of the present disclosure containing one or more antigens from a Zika virus described herein. Such vaccines and/or immunogenic compositions of the present disclosure containing one or more antigens from a Zika virus described herein may be useful for treating or preventing Zika virus infection in a subject in need thereof and/or inducing an immune response, such as a protective immune response, against Zika virus in a subject in need thereof.

[0153] Typically, vaccines and/or immunogenic compositions of the present disclosure are prepared as injectables either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid prior to injection may also be prepared. Such preparations may also be emulsified or produced as a dry powder. The active immunogenic ingredient is often mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredient. Suitable excipients are, for example, water, saline, dextrose, sucrose, glycerol, ethanol, or the like, and combinations thereof. In addition, if desired, the vaccine or immunogenic composition may contain auxiliary substances such as wetting or emulsifying agents, pH buffering agents, or adjuvants which enhance the effectiveness of the vaccine or immunogenic composition.

[0154] Vaccines or immunogenic compositions may be conventionally administered parenterally, by injection, for example, either subcutaneously, transcutaneously,

intradermally, subdermally or intramuscularly. Additional formulations which are suitable for other modes of administration include suppositories and, in some cases, oral, peroral, intranasal, buccal, sublingual, intraperitoneal, intravaginal, anal and intracranial formulations. For suppositories, traditional binders and carriers may include, for example, polyalkalene glycols or triglycerides; such suppositories may be formed from mixtures containing the active ingredient in the range of 0.5% to 10%, or even 1-2%.In certain embodiments, a low melting wax, such as a mixture of fatty acid glycerides or cocoa butter is first melted and the Zika virus vaccine or/or immunogenic composition described herein are dispersed homogeneously, for example, by stirring. The molten homogeneous mixture is then poured into conveniently sized molds, allowed to cool, and to solidify.

[0155] Formulations suitable for intranasal delivery include liquids (*e.g.*, aqueous solution for administration as an aerosol or nasal drops) and dry powders (*e.g.* for rapid deposition within the nasal passage). Formulations include such normally employed excipients as, for example, pharmaceutical grades of mannitol, lactose, sucrose, trehalose, xylitol, and chitosan. Mucosadhesive agents such as chitosan can be used in either liquid or powder formulations to delay mucociliary clearance of intranasally-administered formulations. Sugars such as mannitol, sorbitol, trehalose, and/or sucrose can be used as stability agents in liquid formulations and as stability, bulking, or powder flow and size agents in dry powder formulations. In addition, adjuvants such as monophosphoryl lipid A (MLA), or derivatives thereof, or CpG oligonucleotides can be used in both liquid and dry powder formulations as an immunostimulatory adjuvant.

[0156] Formulations suitable for oral delivery include liquids, solids, semi-solids, gels, tablets, capsules, lozenges, and the like. Formulations suitable for oral delivery include tablets, lozenges, capsules, gels, liquids, food products, beverages, nutraceuticals, and the like. Formulations include such normally employed excipients as, for example, pharmaceutical grades of mannitol, sorbitol, trehalose, sucrose, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, and the like. Other Zika virus vaccines and immunogenic compositions may take the form of solutions, suspensions, pills, sustained release formulations or powders and contain 10-95% of active ingredient, or 25-70%.For oral formulations, cholera toxin is an interesting formulation partner (and also a possible conjugation partner).

[0157] The Zika virus vaccines and/or immunogenic compositions when formulated for vaginal administration may be in the form of pessaries, tampons, creams, gels, pastes, foams or sprays. Any of the foregoing formulations may contain agents in addition to Zika virus vaccine and/or immunogenic compositions, such as carriers, known in the art to be appropriate.

[0158] In some embodiments, the Zika virus vaccines and/or immunogenic compositions of the present disclosure may be formulated for systemic or localized delivery. Such formulations are well known in the art. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's or fixed oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers (such as those based on Ringer's dextrose), and the like. Systemic and localized routes of administration include, e.g., intradermal, topical application, intravenous, intramuscular, etc.

[0159] The vaccines and/or immunogenic compositions of the present disclosure may be administered in a manner compatible with the dosage formulation, and in such amount as will be therapeutically effective and immunogenic. The quantity to be administered depends on the subject to be treated, including, e.g., the capacity of the individual's immune system to mount an immune response, and the degree of protection desired. Suitable dosage ranges may include, for example, from about 0.1 μg to about 100 μg, about 1 μg to about 100 μg, about 2 μg to about 100 μg, about 5 μg to about 100 μg, about 10 μg to about 100 μg, about 15 μg to about 100 μg, about 20 μg to about 100 μg, about 25 μg to about 100 μg, about 30 μg to about 100 μg, about 35 μg to about 100 μg, about 40 μg to about 100 μg, about 45 μg to about 100 μg, about 50 μg to about 100 μg, about 55 μg to about 100 μg, about 60 μg to about 100 μg, about 65 μg to about 100 μg, 70 μg to about 100 μg, about 75 μg to about 100 μg, about 80 μg to about 100 μg, about 85 μg to about 100 μg, about 90 μg to about 100 μg, about 95 μg to about 100 μg, about 0.1 μg to about 95 μg, about 1 μg to about 95 μg, about 2 μg to about 95 μg, about 5 μg to about 95 μg, about 10 μg to about 95 μg, about 15 μg to about 95 μg, about 20 μg to about 95 μg, about 25 μg to about 95 μg, about 30 μg to about 95 μg, about 35 μg to about 95 μg, about 40 μg to about 95 μg, about 45 μg to about 95 μg, about 50 μg to about 95 μg, about 55 μg to about 95 μg, about 60 μg to about 95 μg, about 65 μg to about 95 μg, 70 μg to about 95 μg, about 75 μg to about 95 μg, about 80 μg to about 95 μg, about 85 μg to about 95 μg, about 90 μg to about 95 μg, about 0.1 μg to about 90 μg, about 1 μg to about 90 μg, about 2 μg to about 90 μg, about 5 μg to about 90 μg, about 10 μg

to about 90 µg, about 15 µg to about 90 µg, about 20 µg to about 90 µg, about 25 µg to about 90 µg, about 30 µg to about 90 µg, about 35 µg to about 90 µg, about 40 µg to about 90 µg, about 45 µg to about 90 µg, about 50 µg to about 90 µg, about 55 µg to about 90 µg, about 60 µg to about 90 µg, about 65 µg to about 90 µg, 70 µg to about 90 µg, about 75 µg to about 90 µg, about 80 µg to about 90 µg, about 85 µg to about 90 µg, about 0.1 µg to about 85 µg, about 1 µg to about 85 µg, about 2 µg to about 85 µg, about 5 µg to about 85 µg, about 10 µg to about 85 µg, about 15 µg to about 85 µg, about 20 µg to about 85 µg, about 25 µg to about 85 µg, about 30 µg to about 85 µg, about 35 µg to about 85 µg, about 40 µg to about 85 µg, about 45 µg to about 85 µg, about 50 µg to about 85 µg, about 55 µg to about 85 µg, about 60 µg to about 85 µg, about 65 µg to about 85 µg, 70 µg to about 85 µg, about 75 µg to about 85 µg, about 80 µg to about 85 µg, about 0.1 µg to about 80 µg, about 1 µg to about 80 µg, about 2 µg to about 80 µg, about 5 µg to about 80 µg, about 10 µg to about 80 µg, about 15 µg to about 80 µg, about 20 µg to about 80 µg, about 25 µg to about 80 µg, about 30 µg to about 80 µg, about 35 µg to about 80 µg, about 40 µg to about 80 µg, about 45 µg to about 80 µg, about 50 µg to about 80 µg, about 55 µg to about 80 µg, about 60 µg to about 80 µg, about 65 µg to about 80 µg, 70 µg to about 80 µg, about 75 µg to about 80 µg, about 0.1 µg to about 75 µg, about 1 µg to about 75 µg, about 2 µg to about 75 µg, about 5 µg to about 75 µg, about 10 µg to about 75 µg, about 15 µg to about 75 µg, about 20 µg to about 75 µg, about 25 µg to about 75 µg, about 30 µg to about 75 µg, about 35 µg to about 75 µg, about 40 µg to about 75 µg, about 45 µg to about 75 µg, about 50 µg to about 75 µg, about 55 µg to about 75 µg, about 60 µg to about 75 µg, about 65 µg to about 75 µg, 70 µg to about 75 µg, about 0.1 µg to about 70 µg, about 1 µg to about 70 µg, about 2 µg to about 70 µg, about 5 µg to about 70 µg, about 10 µg to about 70 µg, about 15 µg to about 70 µg, about 20 µg to about 70 µg, about 25 µg to about 70 µg, about 30 µg to about 70 µg, about 35 µg to about 70 µg, about 40 µg to about 70 µg, about 45 µg to about 70 µg, about 50 µg to about 70 µg, about 55 µg to about 70 µg, about 60 µg to about 70 µg, about 65 µg to about 70 µg, about 0.1 µg to about 65 µg, about 1 µg to about 65 µg, about 2 µg to about 65 µg, about 5 µg to about 65 µg, about 10 µg to about 65 µg, about 15 µg to about 65 µg, about 20 µg to about 65 µg, about 25 µg to about 65 µg, about 30 µg to about 65 µg, about 35 µg to about 65 µg, about 40 µg to about 65 µg, about 45 µg to about 65 µg, about 50 µg to about 65 µg, about 55 µg to about 65 µg, about 60 µg to about 65 µg, about 0.1 µg to about 60 µg, about 1 µg to about 60 µg, about 2 µg to about 60 µg, about 5 µg to about 60 µg, about 10 µg to about 60 µg, about 15 µg to about 60 µg, about 20 µg to about 60 µg, about 25 µg to about 60 µg, about 30 µg to about 60 µg, about 35 µg to about 60 µg, about 40 µg to about 60 µg, about 45

μg to about 60 μg, about 50 μg to about 60 μg, about 55 μg to about 60 μg, about 0.1 μg to about 55 μg, about 1 μg to about 55 μg, about 2 μg to about 55 μg, about 5 μg to about 55 μg, about 10 μg to about 55 μg, about 15 μg to about 55 μg, about 20 μg to about 55 μg, about 25 μg to about 55 μg, about 30 μg to about 55 μg, about 35 μg to about 55 μg, about 40 μg to about 55 μg, about 45 μg to about 55 μg, about 50 μg to about 55 μg, about 0.1 μg to about 50 μg, about 1 μg to about 50 μg, about 2 μg to about 50 μg, about 5 μg to about 50 μg, about 10 μg to about 50 μg, about 15 μg to about 50 μg, about 20 μg to about 50 μg, about 25 μg to about 50 μg, about 30 μg to about 50 μg, about 35 μg to about 50 μg, about 40 μg to about 50 μg, about 45 μg to about 50 μg, about 0.1 μg to about 45 μg, about 1 μg to about 45 μg, about 2 μg to about 45 μg, about 5 μg to about 45 μg, about 10 μg to about 45 μg, about 15 μg to about 45 μg, about 20 μg to about 45 μg, about 25 μg to about 45 μg, about 30 μg to about 45 μg, about 35 μg to about 45 μg, about 40 μg to about 45 μg, about 0.1 μg to about 40 μg, about 1 μg to about 40 μg, about 2 μg to about 40 μg, about 5 μg to about 40 μg, about 10 μg to about 40 μg, about 15 μg to about 40 μg, about 20 μg to about 40 μg, about 25 μg to about 40 μg, about 30 μg to about 40 μg, about 35 μg to about 40 μg, about 0.1 μg to about 35 μg, about 1 μg to about 35 μg, about 2 μg to about 35 μg, about 5 μg to about 35 μg, about 10 μg to about 35 μg, about 15 μg to about 35 μg, about 20 μg to about 35 μg, about 25 μg to about 35 μg, about 30 μg to about 35 μg, about 0.1 μg to about 30 μg, about 1 μg to about 30 μg, about 2 μg to about 30 μg, about 5 μg to about 30 μg, about 10 μg to about 30 μg, about 15 μg to about 30 μg, about 20 μg to about 30 μg, about 25 μg to about 30 μg, about 0.1 μg to about 25 μg, about 1 μg to about 25 μg, about 2 μg to about 25 μg, about 5 μg to about 25 μg, about 10 μg to about 25 μg, about 15 μg to about 25 μg, about 20 μg to about 25 μg, about 0.1 μg to about 20 μg, about 1 μg to about 20 μg, about 2 μg to about 20 μg, about 5 μg to about 20 μg, about 10 μg to about 20 μg, about 15 μg to about 20 μg, about 0.1 μg to about 15 μg, about 1 μg to about 15 μg, about 2 μg to about 15 μg, about 5 μg to about 15 μg, about 10 μg to about 15 μg, about 0.1 μg to about 10 μg, about 1 μg to about 10 μg, about 2 μg to about 10 μg, about 5 μg to about 10 μg, about 0.1 μg to about 5 μg, about 1 μg to about 5 μg, about 2 μg to about 5 μg, about 0.1 μg to about 2 μg, about 1 μg to about 2 μg, or about 0.1 μg to about 1 μg per dose. In certain embodiments, the dosage can be about 0.1 μg, about 0.2 μg, about 0.3 μg, about 0.4 μg, about 0.5 μg, about 0.6 μg, about 0.7 μg, about 0.8 μg, about 0.9 μg, about 1 μg, about 1.1 μg, about 1.2 μg, about 1.3 μg, about 1.4 μg, about 1.5 μg, about 1.6 μg, about 1.7 μg, about 1.8 μg, about 1.9 μg, about 2 μg, about 2.1 μg, about 2.2 μg, about 2.3 μg, about 2.4 μg, about 2.5 μg, about 2.6 μg, about 2.7 μg, about 2.8 μg, about 2.9 μg, about 3 μg, about 4 μg,

about 5 μg, about 6 μg, about 7 μg, about 8 μg, about 9 μg, about 10 μg, about 11 μg, about 12 μg, about 13 μg, about 14 μg, about 15 μg, about 16 μg, about 17 μg, about 18 μg, about 19 μg, about 20 μg, about 21 μg, about 22 μg, about 23 μg, about 24 μg, about 25 μg, about 26 μg, about 27 μg, about 28 μg, about 29 μg, about 30 μg, about 31 μg, about 32 μg, about 33 μg, about 34 μg, about 35 μg, about 36 μg, about 37 μg, about 38 μg, about 39 μg, about 40 μg, about 41 μg, about 42 μg, about 43 μg, about 44 μg, about 45 μg, about 46 μg, about 47 μg, about 48 μg, about 49 μg, about 50 μg, about 51 μg, about 52 μg, about 53 μg, about 54 μg, about 55 μg, about 56 μg, about 57 μg, about 58 μg, about 59 μg, about 60 μg, about 61 μg, about 62 μg, about 63 μg, about 64 μg, about 65 μg, about 66 μg, about 67 μg, about 68 μg, about 69 μg, about 70 μg, about 71 μg, about 72 μg, about 73 μg, about 74 μg, about 75 μg, about 76 μg, about 77 μg, about 78 μg, about 79 μg, about 80 μg, about 81 μg, about 82 μg, about 83 μg, about 84 μg, about 85 μg, about 86 μg, about 87 μg, about 88 μg, about 89 μg, about 90 μg, about 91 μg, about 92 μg, about 93 μg, about 94 μg, about 95 μg, about 96 μg, about 97 μg, about 98 μg, about 99 μg, or about 100 μg per dose.

[0160] In some embodiments, the dosage may be measured in micrograms (μg) of Zika Envelope protein E (Env). Accordingly, in some embodiments, the dosage ranges may include, for example, from about 0.1 μg Env to about 100 μg Env, about 1 μg Env to about 100 μg Env, about 2 μg Env to about 100 μg Env, about 5 μg Env to about 100 μg Env, about 10 μg Env to about 100 μg Env, about 15 μg Env to about 100 μg Env, about 20 μg Env to about 100 μg Env, about 25 μg Env to about 100 μg Env, about 30 μg Env to about 100 μg Env, about 35 μg Env to about 100 μg Env, about 40 μg Env to about 100 μg Env, about 45 μg Env to about 100 μg Env, about 50 μg Env to about 100 μg Env, about 55 μg Env to about 100 μg Env, about 60 μg Env to about 100 μg Env, about 65 μg Env to about 100 μg Env, 70 μg Env to about 100 μg Env, about 75 μg Env to about 100 μg Env, about 80 μg Env to about 100 μg Env, about 85 μg Env to about 100 μg Env, about 90 μg Env to about 100 μg Env, about 95 μg Env to about 100 μg Env, about 0.1 μg Env to about 95 μg Env, about 1 μg Env to about 95 μg Env, about 2 μg Env to about 95 μg Env, about 5 μg Env to about 95 μg Env, about 10 μg Env to about 95 μg Env, about 15 μg Env to about 95 μg Env, about 20 μg Env to about 95 μg Env, about 25 μg Env to about 95 μg Env, about 30 μg Env to about 95 μg Env, about 35 μg Env to about 95 μg Env, about 40 μg Env to about 95 μg Env, about 45 μg Env to about 95 μg Env, about 50 μg Env to about 95 μg Env, about 55 μg Env to about 95 μg Env, about 60 μg Env to about 95 μg Env, about 65 μg Env to about 95 μg Env, 70 μg Env to about 95 μg Env, about 75 μg Env to about 95 μg Env, about 80 μg Env to about 95

μg Env, about 85 μg Env to about 95 μg Env, about 90 μg Env to about 95 μg Env, about 0.1 μg Env to about 90 μg Env, about 1 μg Env to about 90 μg Env, about 2 μg Env to about 90 μg Env, about 5 μg Env to about 90 μg Env, about 10 μg Env to about 90 μg Env, about 15 μg Env to about 90 μg Env, about 20 μg Env to about 90 μg Env, about 25 μg Env to about 90 μg Env, about 30 μg Env to about 90 μg Env, about 35 μg Env to about 90 μg Env, about 40 μg Env to about 90 μg Env, about 45 μg Env to about 90 μg Env, about 50 μg Env to about 90 μg Env, about 55 μg Env to about 90 μg Env, about 60 μg Env to about 90 μg Env, about 65 μg Env to about 90 μg Env, 70 μg Env to about 90 μg Env, about 75 μg Env to about 90 μg Env, about 80 μg Env to about 90 μg Env, about 85 μg Env to about 90 μg Env, about 0.1 μg Env to about 85 μg Env, about 1 μg Env to about 85 μg Env, about 2 μg Env to about 85 μg Env, about 5 μg Env to about 85 μg Env, about 10 μg Env to about 85 μg Env, about 15 μg Env to about 85 μg Env, about 20 μg Env to about 85 μg Env, about 25 μg Env to about 85 μg Env, about 30 μg Env to about 85 μg Env, about 35 μg Env to about 85 μg Env, about 40 μg Env to about 85 μg Env, about 45 μg Env to about 85 μg Env, about 50 μg Env to about 85 μg Env, about 55 μg Env to about 85 μg Env, about 60 μg Env to about 85 μg Env, about 65 μg Env to about 85 μg Env, 70 μg Env to about 85 μg Env, about 75 μg Env to about 85 μg Env, about 80 μg Env to about 85 μg Env, about 0.1 μg Env to about 80 μg Env, about 1 μg Env to about 80 μg Env, about 2 μg Env to about 80 μg Env, about 5 μg Env to about 80 μg Env, about 10 μg Env to about 80 μg Env, about 15 μg Env to about 80 μg Env, about 20 μg Env to about 80 μg Env, about 25 μg Env to about 80 μg Env, about 30 μg Env to about 80 μg Env, about 35 μg Env to about 80 μg Env, about 40 μg Env to about 80 μg Env, about 45 μg Env to about 80 μg Env, about 50 μg Env to about 80 μg Env, about 55 μg Env to about 80 μg Env, about 60 μg Env to about 80 μg Env, about 65 μg Env to about 80 μg Env, 70 μg Env to about 80 μg Env, about 75 μg Env to about 80 μg Env, about 0.1 μg Env to about 75 μg Env, about 1 μg Env to about 75 μg Env, about 2 μg Env to about 75 μg Env, about 5 μg Env to about 75 μg Env, about 10 μg Env to about 75 μg Env, about 15 μg Env to about 75 μg Env, about 20 μg Env to about 75 μg Env, about 25 μg Env to about 75 μg Env, about 30 μg Env to about 75 μg Env, about 35 μg Env to about 75 μg Env, about 40 μg Env to about 75 μg Env, about 45 μg Env to about 75 μg Env, about 50 μg Env to about 75 μg Env, about 55 μg Env to about 75 μg Env, about 60 μg Env to about 75 μg Env, about 65 μg Env to about 75 μg Env, 70 μg Env to about 75 μg Env, about 0.1 μg Env to about 70 μg Env, about 1 μg Env to about 70 μg Env, about 2 μg Env to about 70 μg Env, about 5 μg Env to about 70 μg Env, about 10 μg Env to about 70 μg Env, about 15 μg Env to about 70 μg Env, about 20 μg Env to about 70 μg Env, about 25 μg Env to about 70 μg Env,

about 30 µg Env to about 70 µg Env, about 35 µg Env to about 70 µg Env, about 40 µg Env to about 70 µg Env, about 45 µg Env to about 70 µg Env, about 50 µg Env to about 70 µg Env, about 55 µg Env to about 70 µg Env, about 60 µg Env to about 70 µg Env, about 65 µg Env to about 70 µg Env, about 0.1 µg Env to about 65 µg Env, about 1 µg Env to about 65 µg Env, about 2 µg Env to about 65 µg Env, about 5 µg Env to about 65 µg Env, about 10 µg Env to about 65 µg Env, about 15 µg Env to about 65 µg Env, about 20 µg Env to about 65 µg Env, about 25 µg Env to about 65 µg Env, about 30 µg Env to about 65 µg Env, about 35 µg Env to about 65 µg Env, about 40 µg Env to about 65 µg Env, about 45 µg Env to about 65 µg Env, about 50 µg Env to about 65 µg Env, about 55 µg Env to about 65 µg Env, about 60 µg Env to about 65 µg Env, about 0.1 µg Env to about 60 µg Env, about 1 µg Env to about 60 µg Env, about 2 µg Env to about 60 µg Env, about 5 µg Env to about 60 µg Env, about 10 µg Env to about 60 µg Env, about 15 µg Env to about 60 µg Env, about 20 µg Env to about 60 µg Env, about 25 µg Env to about 60 µg Env, about 30 µg Env to about 60 µg Env, about 35 µg Env to about 60 µg Env, about 40 µg Env to about 60 µg Env, about 45 µg Env to about 60 µg Env, about 50 µg Env to about 60 µg Env, about 55 µg Env to about 60 µg Env, about 0.1 µg Env to about 55 µg Env, about 1 µg Env to about 55 µg Env, about 2 µg Env to about 55 µg Env, about 5 µg Env to about 55 µg Env, about 10 µg Env to about 55 µg Env, about 15 µg Env to about 55 µg Env, about 20 µg Env to about 55 µg Env, about 25 µg Env to about 55 µg Env, about 30 µg Env to about 55 µg Env, about 35 µg Env to about 55 µg Env, about 40 µg Env to about 55 µg Env, about 45 µg Env to about 55 µg Env, about 50 µg Env to about 55 µg Env, about 0.1 µg Env to about 50 µg Env, about 1 µg Env to about 50 µg Env, about 2 µg Env to about 50 µg Env, about 5 µg Env to about 50 µg Env, about 10 µg Env to about 50 µg Env, about 15 µg Env to about 50 µg Env, about 20 µg Env to about 50 µg Env, about 25 µg Env to about 50 µg Env, about 30 µg Env to about 50 µg Env, about 35 µg Env to about 50 µg Env, about 40 µg Env to about 50 µg Env, about 45 µg Env to about 50 µg Env, about 0.1 µg Env to about 45 µg Env, about 1 µg Env to about 45 µg Env, about 2 µg Env to about 45 µg Env, about 5 µg Env to about 45 µg Env, about 10 µg Env to about 45 µg Env, about 15 µg Env to about 45 µg Env, about 20 µg Env to about 45 µg Env, about 25 µg Env to about 45 µg Env, about 30 µg Env to about 45 µg Env, about 35 µg Env to about 45 µg Env, about 40 µg Env to about 45 µg Env, about 0.1 µg Env to about 40 µg Env, about 1 µg Env to about 40 µg Env, about 2 µg Env to about 40 µg Env, about 5 µg Env to about 40 µg Env, about 10 µg Env to about 40 µg Env, about 15 µg Env to about 40 µg Env, about 20 µg Env to about 40 µg Env, about 25 µg Env to about 40 µg Env, about 30 µg Env to about 40 µg Env, about 35 µg Env to about 40 µg Env, about 0.1 µg Env

to about 35 μg Env, about 1 μg Env to about 35 μg Env, about 2 μg Env to about 35 μg Env, about 5 μg Env to about 35 μg Env, about 10 μg Env to about 35 μg Env, about 15 μg Env to about 35 μg Env, about 20 μg Env to about 35 μg Env, about 25 μg Env to about 35 μg Env, about 30 μg Env to about 35 μg Env, about 0.1 μg Env to about 30 μg Env, about 1 μg Env to about 30 μg Env, about 2 μg Env to about 30 μg Env, about 5 μg Env to about 30 μg Env, about 10 μg Env to about 30 μg Env, about 15 μg Env to about 30 μg Env, about 20 μg Env to about 30 μg Env, about 25 μg Env to about 30 μg Env, about 0.1 μg Env to about 25 μg Env, about 1 μg Env to about 25 μg Env, about 2 μg Env to about 25 μg Env, about 5 μg Env to about 25 μg Env, about 10 μg Env to about 25 μg Env, about 15 μg Env to about 25 μg Env, about 20 μg Env to about 25 μg Env, about 0.1 μg Env to about 20 μg Env, about 1 μg Env to about 20 μg Env, about 2 μg Env to about 20 μg Env, about 5 μg Env to about 20 μg Env, about 10 μg Env to about 20 μg Env, about 15 μg Env to about 20 μg Env, about 0.1 μg Env to about 15 μg Env, about 1 μg Env to about 15 μg Env, about 2 μg Env to about 15 μg Env, about 5 μg Env to about 15 μg Env, about 10 μg Env to about 15 μg Env, about 0.1 μg Env to about 10 μg Env, about 1 μg Env to about 10 μg Env, about 2 μg Env to about 10 μg Env, about 5 μg Env to about 10 μg Env, about 0.1 μg Env to about 5 μg Env, about 1 μg Env to about 5 μg Env, about 2 μg Env to about 5 μg Env, about 0.1 μg Env to about 2 μg Env, about 1 μg Env to about 2 μg Env, or about 0.1 μg Env to about 1 μg Env per dose. In certain embodiments, the dosage can be about 0.1 μg Env, about 0.2 μg Env, about 0.3 μg Env, about 0.4 μg Env, about 0.5 μg Env, about 0.6 μg Env, about 0.7 μg Env, about 0.8 μg Env, about 0.9 μg Env, about 1 μg Env, about 1.1 μg Env, about 1.2 μg Env, about 1.3 μg Env, about 1.4 μg Env, about 1.5 μg Env, about 1.6 μg Env, about 1.7 μg Env, about 1.8 μg Env, about 1.9 μg Env, about 2 μg Env, about 2.1 μg Env, about 2.2 μg Env, about 2.3 μg Env, about 2.4 μg Env, about 2.5 μg Env, about 2.6 μg Env, about 2.7 μg Env, about 2.8 μg Env, about 2.9 μg Env, about 3 μg Env, about 4 μg Env, about 5 μg Env, about 6 μg Env, about 7 μg Env, about 8 μg Env, about 9 μg Env, about 10 μg Env, about 11 μg Env, about 12 μg Env, about 13 μg Env, about 14 μg Env, about 15 μg Env, about 16 μg Env, about 17 μg Env, about 18 μg Env, about 19 μg Env, about 20 μg Env, about 21 μg Env, about 22 μg Env, about 23 μg Env, about 24 μg Env, about 25 μg Env, about 26 μg Env, about 27 μg Env, about 28 μg Env, about 29 μg Env, about 30 μg Env, about 31 μg Env, about 32 μg Env, about 33 μg Env, about 34 μg Env, about 35 μg Env, about 36 μg Env, about 37 μg Env, about 38 μg Env, about 39 μg Env, about 40 μg Env, about 41 μg Env, about 42 μg Env, about 43 μg Env, about 44 μg Env, about 45 μg Env, about 46 μg Env, about 47 μg Env, about 48 μg Env, about 49 μg Env, about 50 μg Env, about 51 μg

Env, about 52 μg Env, about 53 μg Env, about 54 μg Env, about 55 μg Env, about 56 μg Env, about 57 μg Env, about 58 μg Env, about 59 μg Env, about 60 μg Env, about 61 μg Env, about 62 μg Env, about 63 μg Env, about 64 μg Env, about 65 μg Env, about 66 μg Env, about 67 μg Env, about 68 μg Env, about 69 μg Env, about 70 μg Env, about 71 μg Env, about 72 μg Env, about 73 μg Env, about 74 μg Env, about 75 μg Env, about 76 μg Env, about 77 μg Env, about 78 μg Env, about 79 μg Env, about 80 μg Env, about 81 μg Env, about 82 μg Env, about 83 μg Env, about 84 μg Env, about 85 μg Env, about 86 μg Env, about 87 μg Env, about 88 μg Env, about 89 μg Env, about 90 μg Env, about 91 μg Env, about 92 μg Env, about 93 μg Env, about 94 μg Env, about 95 μg Env, about 96 μg Env, about 97 μg Env, about 98 μg Env, about 99 μg Env, or about 100 μg Env per dose.

[0161] Suitable regimens for initial administration and booster shots are also variable but are typified by an initial administration followed by subsequent inoculations or other administrations.

[0162] The manner of application may be varied widely. Any of the conventional methods for administration of a vaccine or immunogenic composition are applicable. These include oral application on a solid physiologically acceptable base or in a physiologically acceptable dispersion, parenterally, by injection or the like. The dosage of the vaccine or immunogenic composition will depend on the route of administration and may vary according to the age of the person to be vaccinated and the formulation of the antigen. The vaccine or immunogenic composition can have a unit dosage volume of more than 0.5mL, of 0.5mL or of less than 0.5mL, as described herein. For instance, it can be administered at a volume of 0.25mL.

[0163] Delivery agents that improve mucoadhesion can also be used to improve delivery and immunogenicity especially for intranasal, oral or lung based delivery formulations. One such compound, chitosan, the N-deacetylated form of chitin, is used in many pharmaceutical formulations. It is an attractive mucoadhesive agent for intranasal vaccine delivery due to its ability to delay mucociliary clearance and allow more time for mucosal antigen uptake and processing. In addition, it can transiently open tight junctions which may enhance transepithelial transport of antigen to the NALT. In a recent human trial, a trivalent inactivated influenza vaccine administered intranasally with chitosan but without any additional adjuvant yielded seroconversion and HI titers that were only marginally lower than those obtained following intramuscular inoculation.

[0164] Chitosan can also be formulated with adjuvants that function well intranasally such as the genetically detoxified *E. coli* heat-labile enterotoxin mutant LTK63. This adds an immunostimulatory effect on top of the delivery and adhesion benefits imparted by chitosan resulting in enhanced mucosal and systemic responses.

[0165] Finally, it should be noted that chitosan formulations can also be prepared in a dry powder format that has been shown to improve vaccine stability and result in a further delay in mucociliary clearance over liquid formulations. This was seen in a recent human clinical trial involving an intranasal dry powder diphtheria toxoid vaccine formulated with chitosan in which the intranasal route was as effective as the traditional intramuscular route with the added benefit of secretory IgA responses. The vaccine was also very well tolerated. Intranasal dry powdered vaccines for anthrax containing chitosan and MLA, or derivatives thereof, induce stronger responses in rabbits than intramuscular inoculation and are also protective against aerosol spore challenge.

[0166] Intranasal vaccines represent an exemplary formulation as they can affect the upper and lower respiratory tracts in contrast to parenterally administered vaccines which are better at affecting the lower respiratory tract. This can be beneficial for inducing tolerance to allergen-based vaccines and inducing immunity for pathogen-based vaccines.

[0167] In addition to providing protection in both the upper and lower respiratory tracts, intranasal vaccines avoid the complications of needle inoculations and provide a means of inducing both mucosal and systemic humoral and cellular responses via interaction of particulate and/or soluble antigens with nasopharyngeal-associated lymphoid tissues (NALT).

[0168] Vaccines and/or immunogenic compositions of the present disclosure are pharmaceutically acceptable. They may include components in addition to the antigen and adjuvant *e.g.* they will typically include one or more pharmaceutical carrier(s) and/or excipient(s). A thorough discussion of such components is available in Gennaro (2000) Remington: The Science and Practice of Pharmacy. 20th edition, ISBN: 0683306472.

[0169] To control tonicity, it is preferred to include a physiological salt, such as a sodium salt. Sodium chloride (NaCl) is preferred, which may be present at between 1 and 20 mg/ml. Other salts that may be present include potassium chloride, potassium dihydrogen phosphate, disodium phosphate dehydrate, magnesium chloride, calcium chloride, etc.

[0170] Vaccines and/or immunogenic compositions of the present disclosure may include one or more buffers. Typical buffers include: a phosphate buffer; a Tris buffer; a borate buffer; a succinate buffer; a histidine buffer (particularly with an aluminum hydroxide adjuvant); or a citrate buffer. Buffers will typically be included in the 5-20mM range.

[0171] The pH of a vaccine or immunogenic composition will generally be between 5.0 and 8.5 or 5.0 and 8.1, and more typically between 6.0 and 8.5 *e.g.* 6.0 and 8.0, 6.5 and 8.0, 6.5 and 7.5, 7.0 and 8.5, 7.0 and 8.0, or between 7.0 and 7.8. A manufacturing process of the present disclosure may therefore include a step of adjusting the pH of the bulk vaccine prior to packaging.

[0172] The vaccine or immunogenic composition is preferably sterile. It is preferably non pyrogenic e.g. containing <1 EU (endotoxin unit, a standard measure) per dose, and preferably <0.1 EU per dose. It is preferably gluten free.

[0173] In certain embodiments, the vaccines and/or immunogenic compositions of the present disclosure may include a detergent in an effective concentration. In some embodiments, an effective amount of detergent may include without limitation, about 0.00005% v/v to about 5% v/v or about 0.0001% v/v to about 1% v/v. In certain embodiments, an effective amount of detergent is about 0.001% v/v, about 0.002% v/v, about 0.003% v/v, about 0.004% v/v, about 0.005% v/v, about 0.006% v/v, about 0.007% v/v, about 0.008% v/v, about 0.009% v/v, or about 0.01% v/v. Without wishing to be bound by theory, detergents help maintain the vaccines and/or immunogenic compositions of the present disclosure in solution and helps to prevent the vaccines and/or immunogenic compositions from aggregating.

[0174] Suitable detergents include, for example, polyoxyethylene sorbitan ester surfactant (known as 'Tweens'), octoxynol (such as octoxynol-9 (Triton X 100) or t octylphenoxypolyethoxyethanol), cetyl trimethyl ammonium bromide ('CTAB'), and sodium deoxycholate, particularly for a split or surface antigen vaccine. The detergent may be present only at trace amounts. Other residual components in trace amounts could be antibiotics (e.g. neomycin, kanamycin, polymyxin B). In some embodiments, the detergent contains polysorbate. In some embodiments, the effective concentration of detergent includes ranges from about 0.00005% v/v to about 5% v/v.

[0175] The vaccines and/or immunogenic compositions are preferably stored at between 2°C and 8°C. They should ideally be kept out of direct light. The antigen and emulsion will typically be in admixture, although they may initially be presented in the form of a kit of separate components for extemporaneous admixing. Vaccines and/or immunogenic compositions will generally be in aqueous form when administered to a subject.

Methods of the Present Disclosure

[0176] Further aspects of the present disclosure relate to methods for using vaccines and/or or immunogenic compositions described herein containing one or more antigens from at least one Zika virus (*e.g.,* a Zika virus comprising a non-human cell adaptation mutation such as a non-human cell adaptation mutation in protein NS1) to treat or prevent Zika virus in a subject in need thereof and/or to induce an immune response to Zika virus in a subject in need thereof. In some embodiments, the present disclosure relates to methods for treating or preventing Zika virus infection in a subject in need thereof by administering to the subject a therapeutically effective amount of a vaccine and/or or immunogenic composition of the present disclosure containing one or more antigens from at least one Zika virus (*e.g.,* a Zika virus comprising a non-human cell adaptation mutation such as a non-human cell adaptation mutation in protein NS1). In some embodiments, the present disclosure relates to methods for inducing an immune response to Zika virus in a subject in need thereof by administering to the subject a therapeutically effective amount of a vaccine and/or or immunogenic composition of the present disclosure containing one or more antigens from at least one Zika (*e.g.,* a Zika virus comprising a non-human cell adaptation mutation such as a non-human cell adaptation mutation in protein NS1). In some embodiments, the administering step induces a protective immune response against Zika virus in the subject. In some embodiments, the subject is a human. In some embodiments, the subject is pregnant or intends to become pregnant.

[0177] The Zika virus vaccines and/or immunogenic compositions disclosed herein may be used to protect or treat a subject (*e.g.,* a mammal such as a human) susceptible to, or suffering from a viral infection, by means of administering the vaccine by intranasal, peroral, oral, buccal, sublingual, intramuscular, intraperitoneal, intradermal, transdermal, subdermal, intravaginal, anal, intracranial, intravenous, transcutaneous, or subcutaneous administration. Methods of systemic administration of the vaccines and/or immunogenic compositions of the present disclosure may include conventional syringes and needles, or devices designed for

ballistic delivery of solid vaccines (WO 99/27961), or needleless pressure liquid jet device (U.S. Pat. No. 4,596,556; U.S. Pat. No. 5,993,412), or transdermal patches (WO 97/48440; WO 98/28037). The Zika virus vaccines and/or immunogenic compositions of the present disclosure may also be applied to the skin (transdermal or transcutaneous delivery WO 98/20734; WO 98/28037). The Zika virus vaccines and/or immunogenic compositions of the present disclosure therefore may include a delivery device for systemic administration, pre-filled with the Zika virus vaccine or immunogenic compositions. Accordingly there is provided methods for treating or preventing Zika virus infection and/or for inducing an immune response in a subject (*e.g.*, a mammal such as a human), including the step of administering a vaccine or immunogenic composition of the present disclosure and optionally including an adjuvant and/or a carrier, to the subject, where the vaccine or immunogenic composition is administered via the parenteral or systemic route.

[0178] The vaccines and/or immunogenic compositions of the present disclosure may be used to protect or treat a subject (*e.g.*, a mammal such as a human) susceptible to, or suffering from a viral infection, by means of administering the vaccine or immunogenic composition via a mucosal route, such as the oral/alimentary or nasal route. Alternative mucosal routes are intravaginal and intra-rectal. The mucosal route of administration may be via the nasal route, termed intranasal vaccination. Methods of intranasal vaccination are well known in the art, including the administration of a droplet, spray, or dry powdered form of the vaccine into the nasopharynx of the individual to be immunized. Nebulized or aerosolized vaccine formulations are potential forms of the Zika virus vaccines and/or immunogenic compositions disclosed herein. Enteric formulations such as gastro resistant capsules and granules for oral administration, suppositories for rectal or vaginal administration are also formulations of the vaccines and/or immunogenic compositions of the present disclosure.

[0179] The Zika virus vaccines and/or immunogenic compositions of the present disclosure may also be administered via the oral route. In such cases the pharmaceutically acceptable excipient may also include alkaline buffers, or enteric capsules or microgranules. The Zika virus vaccines and/or immunogenic compositions of the present disclosure may also be administered by the vaginal route. In such cases, the pharmaceutically acceptable excipients may also include emulsifiers, polymers such as CARBOPOL®, and other known stabilizers of vaginal creams and suppositories. The Zika virus vaccines and/or immunogenic

compositions may also be administered by the rectal route. In such cases the excipients may also include waxes and polymers known in the art for forming rectal suppositories.

[0180] In some embodiments, the administering step includes one or more administrations. Administration can be by a single dose schedule or a multiple dose (prime-boost) schedule. In a multiple dose schedule the various doses may be given by the same or different routes *e.g.* a parenteral prime and mucosal boost, a mucosal prime and parenteral boost, *etc.* Typically they will be given by the same route. Multiple doses will typically be administered at least 1 week apart (*e.g.* about 2 weeks, about 3 weeks, about 4 weeks, about 5 weeks, about 6 weeks, about 7 weeks, about 8 weeks, about 9 weeks, about 10 weeks, about 11 weeks, about 12 weeks, about 16 weeks, *etc.*). Giving two doses separated by from 25-30 days (*e.g.* 28 days) is particularly useful.

[0181] The methods of the present disclosure include administration of a therapeutically effective amount or an immunogenic amount of the Zika virus vaccines and/or immunogenic compositions of the present disclosure. A therapeutically effective amount or an immunogenic amount may be an amount of the vaccines and/or immunogenic compositions of the present disclosure that will induce a protective immunological response in the uninfected, infected or unexposed subject to which it is administered. Such a response will generally result in the development in the subject of a secretory, cellular and/or antibody-mediated immune response to the vaccine. Usually, such a response includes but is not limited to one or more of the following effects; the production of antibodies from any of the immunological classes, such as immunoglobulins A, D, E, G or M; the proliferation of B and T lymphocytes; the provision of activation, growth and differentiation signals to immunological cells; expansion of helper T cell, suppressor T cell, and/or cytotoxic T cell.

[0182] In some embodiments, the protective immunological response induced in the subject after administration of a vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure is greater than the immunological response induced in a corresponding subject administered a vaccine and/or immunogenic composition containing a Zika virus that is not adapted for non-human cell growth and/or comprises a different non-human cell adaptation mutation. In some embodiments, the protective immunological response induced in the subject after administration of the vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure is at least about 5%, at least about 10%, at least about 15%,at least about

20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 95%, or at least about 99% greater than the immunological response induced in a corresponding subject administered a vaccine and/or immunogenic composition containing a Zika virus that is not adapted for non-human cell growth and/or comprises a different non-human cell adaptation mutation. Methods of measuring protective immunological responses are generally known to one of ordinary skill in the art.

[0183] In some embodiments, administration of a vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure induces generation of neutralizing antibodies to Zika virus in the subject. In some embodiments, administration of a vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure induces generation of neutralizing antibodies to Zika virus in the subject in an amount that is greater than the amount of neutralizing antibodies induced in a corresponding subject administered a vaccine and/or immunogenic composition containing a Zika virus that is not adapted for non-human cell growth and/or comprises a different non-human cell adaptation mutation. In some embodiments, administration of a vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure induces generation of neutralizing antibodies to Zika virus in the subject in an amount that is at least about 5%, at least about 10%, at least about 15%,at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 95%, or at least about 99% greater than the than the amount of neutralizing antibodies induced in a corresponding subject administered a vaccine and/or immunogenic composition containing a Zika virus that is not adapted for non-human cell growth and/or comprises a different non-human cell adaptation mutation. In some embodiments, administration of a vaccine and/or immunogenic composition containing a non-human cell adapted Zika virus of the present disclosure induces generation of neutralizing antibodies to Zika virus in the subject in an amount that is at least about 1-fold, at least about 2-fold, at least about 3-fold, at least about 4-fold, at least about 5-fold, at least about 6-fold, at least about 7-fold, at least about 8-fold, at least about 9-fold, at least about 10-fold, at least about 100-fold, or at least about 1000-fold greater than the than the amount

of neutralizing antibodies induced in a corresponding subject administered a vaccine and/or immunogenic composition containing a Zika virus that is not adapted for non-human cell growth and/or comprises a different non-human cell adaptation mutation. Methods of measuring neutralizing antibodies in a subject are generally known to one of ordinary skill in the art.

[0184] Preferably, the therapeutically effective amount or immunogenic amount is sufficient to bring about treatment or prevention of disease symptoms. The exact amount necessary will vary depending on the subject being treated; the age and general condition of the subject to be treated; the capacity of the subject's immune system to synthesize antibodies; the degree of protection desired; the severity of the condition being treated; the particular Zika virus antigen selected and its mode of administration, among other factors. An appropriate therapeutically effective amount or immunogenic amount can be readily determined by one of skill in the art. A therapeutically effective amount or immunogenic amount will fall in a relatively broad range that can be determined through routine trials.

[0185] The present disclosure will be more fully understood by reference to the following Examples. They should not, however, be construed as limiting any aspect or scope of the present disclosure in any way.

EXAMPLES

Example 1: Clonal Zika Virus Strain Generation

[0186] This example describes the production of Zika virus (ZIKAV) strains with a known research history.

Materials and Methods

Vero Cell Maintenance

[0187] One vial of WHO Vero 10-87 cells was rapidly thawed in a water bath and directly inoculated into 19mL pre-warmed DMEM (Dulbecco's modified minimal essential medium) containing penicillin-streptomycin, L-glutamine 40mM, and 10% FBS in a T-75cm^2 flask at 36°C+/2°C, at 5% CO_2. Cells were allowed to grow to confluency and subcultured using TryplE. This flask was expanded to two T-185cm^2 flasks, grown to confluency and subcultured to 31 xT-185cm^2 flasks and grown until the cells reached 100% confluency. Cells

were harvested by trypsinization, centrifuged at 800 x g for 10 minutes, and resuspended in DMEM containing 10% FBS and 10% DMSO at a concentration of $1.9x10^7$ cells/mL. One vial of the Vero cells was rapidly thawed and resuscitated as described above into a T-75cm^2 flask. These were subcultured twice to produce the a cell bank in 13 x T-185cm^2 flasks. After trypsinization, the cells were centrifuged at 800 x g and resuspended in freezing media (DMEM containing 10% FBS, and 10% DMSO) at a concentration of $4.68x10^5$ cells/mL. This cell bank was aliquoted into cryovials.

[0188] The Vero cells were grown and maintained in DMEM containing penicillin-streptomycin, L-glutamine and 10% FBS (cDMEM-10%-FBS). TryplExpress was used to maintain and trypsinize cells. Two days before viral adsorption, 6-well plates were seeded with 4-5 x 10^5 cells/well in 3 mL of cDMEM-10%-FBS or 7 x 10^5 cells in T-25cm^2 flasks in 5 mL cDMEM-10%-FBS, or 1 x 10^4 cells/well in 96-well plates in 0.1mL cDMEM-10%-FBS. Incubators were monitored daily to maintain indicated temperatures. The Vero cell lines were stored in liquid nitrogen.

Plaque Assay

[0189] Viral titers were determined by plaque titration in freshly confluent monolayers of Vero cells grown in 6-well plates. Frozen aliquots were thawed and ten-fold dilution series of the aliquots were made in cDMEM-0%-FBS in 96-well plates. The diluted viruses were maintained on ice prior to inoculation of the Vero cell monolayers. At the time of assay, the growth medium was aspirated from the 6-well plate, and 100 µL of each virus dilution was added to the wells. Virus was adsorbed for 60 min at 36°C±2°C, at 5% CO_2, with frequent (every 10 min) rocking of the plates to prevent drying of the cell sheets. Following viral adsorption, 4 mL of a first agarose overlay (1X cDMEM-2%-FBS + 0.8% agarose) maintained at 40-41°C was added to each well. The agarose was allowed to solidify for 30 min at room temperature, and the plates were then incubated upside down for 4-6 days at 36°C+/2°C, at 5% CO_2. Two mL of a second agarose overlay containing 160 µg/mL of neutral red vital dye was added on day 4. Plaques were visualized on days 5 and 6.

Virus Quantification by TCID50 Assay

[0190] Viral titers were also determined by titration in freshly confluent monolayers of Vero cells grown in 96-well plates. Frozen aliquots were thawed and ten-fold dilution series of the aliquots were made in cDMEM-2%-FBS diluent in 96-well plates. The diluted viruses

were maintained on ice prior to inoculation of the Vero cell monolayers. At the time of assay, the growth medium was aspirated from the 96-well plate, and 100 µL of each virus dilution was added to the wells. The plates were incubated for 5 days at 36°C+/2°C, at 5% CO_2. The 50% Tissue Culture Infective Dose (TCID50) titer was calculated using the Reed/Muench calculator.

Test Articles

[0191] Zika virus strain PRVABC59 (one 0.5 mL vial on dry ice) was received from the Centers for Disease Control and Prevention (CDC) Zika virus identification was confirmed through RT-PCR. The strain tested negative for Alphavirus and mycoplasma contamination by PCR. This information is summarized in **Table 1**.

Table 1: PRVABC59 strain information

Strain	Isolation Information	Patient information	Prep info	Analyses	PFU
PRVABC59 (Asian)	Human serum; travel to Puerto Rico in 2015	None provided	Passage: Vero(2)C6/36(1) Prep: 29Jan2016 Host: C6/36	▪ Sequencing by ion torrent: gene accession #KU501215 ▪ PFU by plaque assay ▪ Identity by RT-PCR ▪ (-) For alphaviruses by PCR ▪ (-) for mycoplasma by ATCC and ABM PCR	6.7 log pfu/mL

Sequencing

[0192] A QIAampViral RNA Mini Spin kit was used to extract RNA from stabilized virus harvests of each isolate according to manufacturer protocols. Extracted RNA from each isolate was used to create and amplify 6 cDNA fragments encompassing the entire Zika viral genome. Amplified cDNA fragments were analyzed for size and purity on a 1% Agarose/TBE gel and subsequently gel purified using a Qiagen Quick Gel Extraction Kit. An ABI 3130XL Genetic Analyzer sequencer was used to conduct automatic sequencing reactions. Lasergene SeqMan software was used to analyze sequencing data.

Results

[0193] A ZIKAV strain with a known research history that was relevant to the current ZIKAV outbreak in the America's was sought. For this reason, ZIKAV strain PRVABC59 was chosen. To generate a well-characterized virus adapted for growth in Vero cells, the ZIKAV PRVABC59 was first amplified in Vero cells (P1).

[0194] Flasks of Vero cells (T-175cm^2), 100% confluent, were infected at an MOI of 0.01 in 4mL of cDMEM-0%-FBS. Virus was adsorbed to the monolayer for 60 minutes at 36°C±2°C, at 5% CO_2, then 20 mL of cDMEM-0%-FBS was applied for viral amplification at 36°C±2°C, at 5% CO_2. The cell layer was monitored daily for cytopathic effect (CPE) following inoculation (**FIG. 1**). The supernatant was harvested after 96 hours by collecting the media and clarifying by centrifugation (600 x g, 4°C, 10 min). The harvest was stabilized by adding trehalose to a final concentration of 18% w/v. The bulk was aliquoted into 0.5mL cryovials and stored at -80 °C.

[0195] The stabilized P1 harvest was analyzed for the presence of infectious virus on Vero cell monoloayers by a TCID50 assay. Growth kinetics were monitored by taking daily aliquots beginning on hour 0. Peak titer was reached by hour 72 (**FIG. 2**).

[0196] P1 material was plaque-purified by titrating the harvest from day 3 on 6-well monolayers of Vero cells. Plaques were visualized on day 6, and 10 plaques to be isolated were identified by drawing a circle around a distinct and separate plaque on the bottom of the plastic plate. Plaques were picked by extracting the plug of agarose using a sterile wide bore pipette while scraping the bottom of the well and rinsing with cDMEM-10%-FBS. The agarose plug was added to 0.5 mL of cDMEM-10%-FBS, vortexed, labeled as PRVABC59 P2a-j and placed in an incubator overnight at 36°C±2°C, at 5% CO_2.

[0197] Three plaques (PRVABC59 P2a-c) were carried forward for additional purification. Each isolate was plated neat in duplicate onto a fresh 6-well monolayer of Vero cells. This P2/P3 transition was plaque purified, and labeled PRVABC59 P3a-j.

[0198] Six plaques (PRVABC59 P3a-f) were carried forward for a final round of purification. Each isolate was plated neat in duplicate onto a fresh 6-well monolayer of Vero cells. This P3/P4 transition was plaque purified, and labeled PRVABC59 P4a-j.

[0199] Six plaques (PRVABC59 P4a-f) from the P4 plaque purification were blind passaged on monolayers of Vero cells in T-25 cm^2 flasks. Each plaque pick was diluted in 2 mL cDMEM-0%-FBS – 1 mL was adsorbed for 1 hour at 36°C±2°C, at 5% CO_2; the other 1 mL was stabilized with trehalose (18% v/v final) and stored at <-60°C. Following virus adsorption, cDMEM-0%-FBS was added to each flask and allowed to grow at 36°C±2°C, at 5% CO_2 for 4 days. Virus supernatants were harvested, clarified by centrifugation (600 x g, 4C, 10 min), stabilized in 18% trehalose and aliquoted and stored at <-60°C. This P5 seed was tested by TCID50 for Zika virus potency (**FIG. 3**).

[0200] Confluent monolayers of T-175cm^2 flasks of Vero cells were infected with each of the six clones of PRVABC59 (P5a-f) at an MOI of 0.01 in 4mL cDMEM-0%-FBS. The virus was allowed to adsorb for 60 minutes at 36°C+/2°C, at 5% CO_2, after which 20 mL of cDMEM-0%-FBS was added to each flask and allowed to grow at 36°C+/2°C, at 5% CO_2. Vero cell monolayer health and CPE was monitored daily. Virus was harvested on days 3 and 5 as indicated (**FIG. 4**). The P6 strain harvests from days 3 and 5 were pooled, stabilized with 18% trehalose, aliquoted and stored <-60°C.

[0201] Each of the six clones of PRVABC59 (P6a-f) were tested for Zika virus potency (**FIG. 5**). The potency was determined by two different methods, TCID50 and plaque titration. The TCID50 was calculated by visual inspection of CPE (microscope) and by measuring the difference in absorbance (A_{560}-A_{420}) of the wells displaying CPE (yellow in color) compared with red (no CPE). The plates were read on a plate reader, and applied to the same calculator as the microscopically read-plates (absorbance). The values in TCID50 between the two scoring techniques are quite similar, while the values obtained by plaque titration are lower.

[0202] A summary of the generation of the P6 virus is shown in **Table 2** below.

Table 2: Summary of virus passage and characterization for the generation of clonal ZIKAV strains

Passage	Seed production/purification	Characterization
P1	Virus amplification in Vero	TCID50 titer
P2	Amplify P1 by plaque titration	plaque purification
P3	Pick and passage plaques from P2 plaque assay	plaque purification
P4	Pick and passage plaques from P3 plaque assay	plaque purification
P5	Amplify P4 plaques (a-f)	TCID50 titer

Passage	Seed production/purification	Characterization
P6	Amplify P5 (a-f) virus	TCID50 titer, plaque phenotype, genotype, growth kinetics

[0203] An isolated Zika virus clone that closely resembled the envelope glycoprotein sequence of the original isolate was sought, since the envelope protein of flaviviruses is the dominant immunogenic portion of the virus. PRVABC59 clones P6a, P6c, P6d and P6f contained a G→T mutation at nucleotide 990 in the envelope region (G990T), resulting in an amino acid mutation of Val→Leu at envelope residue 330, whereas the envelope gene of PRVABC59 clones P6b and P6e were identical relative to the reference strain (GenBank ref KU501215.1) (**Table 3** and **FIG. 6**).

Table 3: Sequencing of PRVABC59 P6 clones

Envelope sequencing (reference gene from PRVABC59; accession #KU501215)				
Strain	**Nucleotide**	**Amino Acid**	**Mutation**	**Comments**
PRVABC59 P6a	Env-990: G→T	Env-330: Val330→Leu	**Val/Leu**	Mutation in 3 of 4 reads.
PRVABC59 P6b	Env-1404: T→G silent	Wild type	Wild type	Wild type relative to reference.
PRVABC59 P6c	Env-990: G→T	Env-330: Val330→Leu	**Val/Leu**	Mutation in 3 of 4 reads.
PRVABC59 P6d	Env-990: G→T	Env-330: Val330→Leu	**Val/Leu**	Mutation in 2 of 2 reads.
PRVABC59 P6e	Wild type	Wild type	Wild type	Wild type relative to reference.
PRVABC59 P6f	Env-990: G→T	Env-330: Val330→Leu	**Val/Leu**	Mutation in 2 of 2 reads. 190 bp not sequenced (aa 421 – 484).
Full genome sequencing (reference gene from PRVABC59;accession #KU501215)				
Strain	**Nucleotide**	**Amino Acid**	**Mutation**	**Comments**
PRVABC59 P6b	Env-1404 T→G	Wild-type	Silent	Mutation in 2 of 2 reads
	NS1-292 T→G	NS1-98 Trp98→Gly	**Trp/Gly**	Mutation in 2 of 2 reads
PRVABC59 P6e	NS1-292 T→G	NS1-98 Trp98→Gly	**Trp/Gly**	Mutation in 2 of 2 reads

[0204] The two clones lacking mutations in the wild-type envelope sequence were then subjected to full genome sequencing. Sequencing results are summarized in **Table 3** above. Sequence analysis revealed a T→G substitution at nucleotide 292 in the NS1 region for both

clones, resulting in a Trp→Gly mutation at NS1 residue 98. This mutation was also later confirmed through deep sequencing. The NS1 W98G mutation is located in the intertwined loop of the wing domain of ZIKAV NS1, which has been implicated in membrane association, interaction with envelope protein and potentially hexameric NS1 formation. While other tryptophan residues (W115, W118), are highly conserved across flaviviruses, W98 is not (**FIG. 7**). Interestingly, however, 100% conservation of the W98 residue is observed across 11 different ZIKAV strains, including those from the African and Asian lineages. The identified mutations in each strain are summarized in **Table 4**.

Table 4: Summary of mutations identified in PRVABC59 P6 clones

Mutations identified in envelope		
Clone	**Nucleotide**	**Amino Acid**
P6a	G990T	V330L
P6b	T1404G	(silent)
P6c	G990T	V330L
P6d	G990T	V330L
P6e	none	none
P6f	G990T	V330L
Additional mutations identified in genome		
Clone	**Nucleotide**	**Amino Acid**
P6b	NS1-T292G	NS1-W98G
P6e	NS1-T292G	NS1-W98G
Ref sequence: KU501215.1 (PRVABC59)		

[0205] Phenotypic analysis of the ZIKAV PRVABC59 P6 stocks was conducted to characterize the ZIKAV clones. As illustrated in **FIG. 8** and quantified in **FIG. 9**, each clonal isolate consisted of a relatively homogeneous population of large-sized plaques as compared to the P1 virus which had a mixed population of large and small plaques. These data suggest the successful isolation of single ZIKAV clones.

[0206] Next, growth kinetics analyses in Vero cells of the ZIKAV PRVABC59 P6 clones were analyzed. Vero cells were infected with 0.01 $TCID_{50}$/cell of each ZIKAV P6 clones in serum free growth medium. Viral supernatant samples were taken daily and simultaneously assayed for infectious titer by $TCID_{50}$ assay. For all P6 clones, peak titer occurred between

day 3 and 4 (~9.0 $\log_{10}$ $TCID_{50}$/mL). There was no significant difference in growth kinetics of the various P6 clones (**FIG. 10**).

[0207] Taken together, the results indicate that a Zika virus seed was successfully generated. This seed selection required understanding of growth history, kinetics, yield, genotype, and phenotype of the virus. Importantly, clonal isolation of the Zika virus strains allowed for the successful purification of the virus away from contaminating agents (*e.g.*, adventitious agents that may be in the parental human isolate). Interestingly, three sequential plaque purifications succeeded in quickly selecting Vero-cell adapted virus (strains P6a-f), where these strains were able to replicate well in serum-free Vero cell cultures, with strain P6a, c, d, and f harboring a mutation in the viral envelope protein, while strains p6b and p6e obtained a mutation in the viral NS1 protein (with no modification to the viral envelope). Additionally, the Vero-adapted strains enabled efficient and reproducible growth and manufacture of subsequent viral passages propagated from these strains. Without wishing to be bound by theory, the Env-V330L mutation observed in strains P6a, c, d, and f may potentially be a result of *in vitro* adaptation, as a mutation at Env 330 was also observed upon passaging in Vero cells (Weger-Lucarelli *et al.* 2017. Journal of Virology). Because the envelope protein is the dominant immunogenic epitope of Zika virus, strains containing a Vero adaptive mutation in Env may negatively impact vaccine immunogenicity. Without wishing to be bound by theory, the adaptation mutation in protein NS1 appears not only to enhance viral replication, but may also reduce or otherwise inhibit the occurrence of undesirable mutations, such as in the envelope protein E (Env) of the Zika virus. In addition, NS1 may be known to bind to the Envelope protein during the life cycle of the virus. This mutation (NS1 W98G) may be implicated in changing the ability of the NS1 to associate, and possibly co-purify, with the virus during downstream processing. NS1 may also be immunogenic, and could be implicated in the immune response to the vaccine.

Example 2: Preclinical immunogenicity and efficacy of a purified inactivated Zika virus vaccine (PIZV) derived from the P6b and P6e strains

[0208] The following example describes the preclinical immunogenicity and efficacy in CD1 and AG129 mice of an inactivated Zika virus vaccine (PIZV) derived from the P6b and P6e strains. As described in Example 1, six clones were generated from the epidemically relevant PRVABC59 strain, and two (P6b and P6e) were chosen for further preclinical immunogenicity and efficacy studies.

Materials and Methods

Purification, inactivation and formulation of a Zika virus vaccine

[0209] A lot of inactivated ZIKAV vaccine, suitable for use in preclinical immunogenicity and efficacy studies, was generated and characterized. Virus was amplified from the P6b and P6e strains by infecting flasks of confluent Vero cells at a MOI of 0.01. Virus was adsorbed for 1 hour at 36°C ± 2°C / 5% CO_2. Following adsorption, 20 mL of cDMEM-0%-FBS was added to each flask, and incubated at 36°C ± 2°C / 5% CO_2 for five days. Cell supernatants were harvested on day 3 and 5 post-infection, and cell debris was clarified by centrifugation.

[0210] For each isolate, clarified supernatants were pooled, stabilized in DMEM containing 18% trehalose and stored at <-60°C. Pooled, clarified virus supernatants were thawed in a 37°C water bath and treated with benzonase overnight at 4°C. Following benzonase treatment, each sample was applied to a Sartorius PP3 depth filter. Following depth filtration, each sample was applied to a Centricon Plus-70 tangential flow filtration (TFF) device. Retentate was buffer exchanged, diluted, and applied to a Sartorius SartobindQ IEXNano. Each sample was applied to a second Sartorius SartobindQ IEXNano and eluted using a 3 step-elution process with 250 mM, 500 mM, and 750 mM NaCl. Following MonoQ chromatography and dilution, each 250 mM eluate was applied to a Centricon Plus-70 cross flow filtration (CFF) device for buffer exchange, diluted to 35 mL with PBS, and stored at 2-8°C.

[0211] For formalin inactivation, freshly prepared 1% formaldehyde was added dropwise to each purified sample with gentle swirling to obtain a final formaldehyde concentration of 0.02%.Samples were incubated at room temperature (~22°C) for 14 days with daily inversion. Formaldehyde was neutralized with sodium metabisulfite for 15' at room temperature before being applied to a Centricon Plus-70 tangential flow filtration (TFF) device. Buffer exchange was performed four times by the addition of 50 mL Drug Substance Buffer (10 mM NaH2PO4 , 50 mM NaCl, 6% sucrose, pH 7.4) . Each sample was then diluted to 15 mL with Drug Substance Buffer, sterilized using a 0.2m syringe filter, aliquoted into sterile stoppered glass vials (0.5 mL per vial) and frozen at <-60°C.

[0212] Virus inactivation was confirmed by TCID50 assay and double infectivity assay. Briefly drug substance sample was applied to C6/36 cells and allowed to amplify for 6 days.

Supernatant from C6/36 cells was applied to Vero cells and CPE was monitored for 8 days. For drug product formulation, vials of PIZV drug substance were thawed, pooled according to sample type, and diluted to 1 µg/mL or 10 µg/mL in PBS with or without Alhydrogel (Brenntag ; 0.5 mg/mL final, 0.050 mg/dose) and incubated overnight at 2-8°C with gentle agitation. The resulting drug product lots were then aliquoted into sterile stoppered glass vials and stored at 2-8°C until use. **FIG. 11** provides a summary of the steps used to prepare drug product.

Mouse immunization and challenge

[0213] For the immunogenicity study, six-week old male and female Swiss-ICR (CD-1) mice were divided into 6 groups (n = 10/group). On Day 0, mice in groups 1-5 were inoculated with 0.1 mL of vaccine by the intramuscular (i.m.) route (2 x 0.05 mL injections). Mice in group 6 were inoculated with PBS as a placebo control. Mice were boosted on day 28 and 56 using the same dosage and vaccine type as day 0. Blood samples were collected on day -1 (pre-immune), day 27 (prime), day 42 (boost 1) and day 70 (boost 2).

[0214] For the immunogenicity and efficacy study, four-week old male and female AG129 mice were divided into 7 groups (n = 5/group). On Day 0, mice in groups 1-6 were inoculated with 0.1 mL of vaccine by the intramuscular (i.m.) route (2 x 0.05 mL injections). Mice in group 7 were inoculated with PBS as a placebo control. Mice were boosted on day 28 using the same dosage and vaccine type as on day 0. Blood samples were collected from the tail vein on day -1 (pre-immune), day 27 (prime) and day 55 (boost). At the time of euthanization, mice were bled via cardiac puncture under deep anesthesia with isofluorane (terminal). On day 56, mice were intraperitoneally challenged with 10^4 plaque forming units (PFU) of ZIKAV PRVABC59.

Serum transfer

[0215] Serum was collected from PIZV-vaccinated and challenged AG129 mice, and were frozen after pooling (groups 1, 2, 4, and 5 of Table 6). The serum pool was thawed, and the test articles were generated by three-fold dilutions of the serum pool in PBS. A placebo was generated using 3-fold dilutions of AG129 normal mouse serum in PBS.

[0216] The test articles were administered as 0.1 mL intraperitoneal injections into AG129 mice (an equivalent volume of the placebo article was administered to control mice).

Animals were then challenged intraperitoneally with 10^4 plaque forming units of Zika virus strain PRVABC59 in 100µL.

[0217] Allowable blood volume by weight was collected as whole blood by tail bleeding from ten mice on day -11 (pre-immunization). Whole blood was collected from each mouse on day 1 (primary, circulating Nab) and day 4 (viremia) by tail bleeding. Terminal bleeding after lethal challenge was performed by heart puncture under deep anesthesia for larger volume before euthanization by cervical dislocation. Blood samples were collected in microtainer SST serum separation gel tubes and allowed to clot for at least 30 min before separation of serum by centrifugation (10,000 x g for 2 min) and frozen at -80°C.

Plaque reduction neutralization test

[0218] Neutralizing antibody titers were determined by a plaque reduction neutralization test (PRNT) as described previously (*See e.g.,* Osorio *et al.* Lancet Infect Dis. 2014 Sep;14(9):830-8).

Reporter virus particle (RVP) neutralization assay

[0219] Neutralizing antibody titers were analyzed by titration of serum samples with a constant amount of Zika RVPs in Vero cells grown in 96-well plates. RVPs contained the prME proteins of Zika (strain SPH2012) and a Dengue-based Renilla luciferase reporter. Briefly, sera were heat inactivated at 56°C for 30 min, diluted, and then incubated at 37°C with RVPs. The serum/RVP mixture was then mixed with Vero cells and incubated for 72 hours at 37°C ± 2°C/ 5% CO_2 before detection with luciferase substrate. Data was analyzed using JMP11 non-linear 4 parameter analysis, normalized to a positive tracking control and effective dose 50% (EC_{50}) was reported.

[0220] Unless indicated to the contrary, all additional experimental methods were carried out as described in Example 1 above.

Results

[0221] To assess the immunogenicity of the PIZV candidates in 6 week old male and female CD-1 mice, groups of CD-1 mice (N=10/group) were immunized by the i.m. route with either a 0.1 µg (+ alum), 1.0 µg (+ alum) dose of a vaccine derived from either ZIKAV PRVABC69 P6b or P6e virus strains. To assess the need for adjuvant, a group of animals was

vaccinated with 0.1 μg of vaccine derived from P6e and lacking alum adjuvant. Vaccinations occurred on days 0, 28, and 56, with group 6 receiving PBS as a placebo control (**FIG. 12A** and **Table 5**).

Table 5: PIZV formulations and challenges in CD-1 mice

Group	Strain	Dose (μg)	Alum (μg)	N
1	P6b	0.1	0.50	10
2	P6b	1.0	0.50	10
3	P6e	0.1	0.50	10
4	P6e	1.0	0.50	10
5	P6e	0.1	-	10
6	Placebo (PBS)	-	-	10

[0222] Following vaccination, serum samples collected after primary (day 27), secondary (day 40) and tertiary (day 70) immunizations were tested for ZIKAV-specific neutralizing antibodies by RVP neutralization assay (**FIG. 12B**). Twenty-seven days after receiving the first dose, a slight neutralizing antibody response was observed in mice vaccinated with PIZV derived from either clone containing alum, as compared to the PBS placebo control group. Importantly, this response increased significantly upon a second immunization (day 40), but was not additionally enhanced upon immunization with a third dose (day 70). No neutralizing antibody response was observed in mice vaccinated with non-adjuvanted vaccine (**FIG. 12B**).

[0223] To assess the immunogenicity and protective efficacy of the PIZV candidates, groups of 4 week old AG129 mice (n=5/group) were immunized by the i.m. route with either a 0.1 μg dose (+ alum), 1.0 μg dose (+ alum) or 0.1 μg dose (- alum) of a vaccine derived from either the ZIKAV PRVABC59 P6b or P6e stocks on days 1 and 28 (**FIG. 13A** and **Table 6**).

Table 6: PIZV formulations and challenges in AG129 mice

Group	Sex	Strain	Dose (μg)	Alum (μg)	N
1	F	P6b	0.1	0.50	5
2	F	P6b	1.0	0.50	5
3	F	P6b	0.1	-	5
4	M	P6e	0.1	0.50	5
5	M	P6e	1.0	0.50	5
6	M	P6e	0.1	-	5
7	M	Placebo (PBS)	-	-	5

[0224] Following vaccination, vaccinated and control mice were intraperitoneally challenged at day 56 with 10^4 PFU of ZIKAV PRVABC59 (low passage). Serum samples collected after primary (D27) and secondary (D55) immunizations were tested for ZIKAV-specific neutralizing antibody response (**FIG. 13B** and **Table 7**). Only groups receiving the high dose of alum-adjuvanted vaccine (groups 2 and 5) elicited a neutralizing antibody response after a single immunization, which increased dramatically after boosting. In contrast, groups receiving either the low or high dose of alum-adjuvanted vaccine produced a high neutralizing antibody response after a second dose. Upon receiving two doses of vaccine, there was no statistical difference between groups of mice receiving alum-adjuvanted vaccine, regardless of the dosage or the derivation from the P6 clone.

Table 7: ZIKAV-specific neutralizing antibody response

		Serum neutralizing antibody titers			
		D27 (prime)		D55 (boost)	
Group	**Formulation**	**GMT**	**% sc**	**GMT**	**% sc**
1	P6b 0.1 μg + alum	<20	40	1280	100
2	P6b 1.0 μg + alum	135	80	2229	100
3	P6b 0.1 μg - alum	<20	0	<20	0
4	P6e 0.1 μg + alum	<20	20	640	100
5	P6e 1.0 μg + alum	30	100	905	100
6	P6e 0.1 μg - alum	<20	0	<20	20
7	PBS	<20	0	<20	0

[0225] All groups were also monitored for mortality, morbidity and weight loss for 21 days post challenge. Viremia following challenge was detected and quantitated by plaque titration. Mice vaccinated with a low or high dose of PIZV candidates formulated with alum (groups 1, 2, 4 and 5) were fully protected from lethal ZIKAV challenge, as assessed by the plaque reduction neutralization test (PRNT) assay, as well as a comparable secondary neutralization assay (**Table 8**). No weight loss or clinical signs of illness were observed in vaccinated mice, none had detectable infectious viremia three days post challenge, and all mice vaccinated with either low or high dose antigen + alum adjuvant survived to 21 days post-challenge (**FIGS. 14-16**). In contrast, challenge of all naïve mice resulted in high

viremia on day 2 post challenge and morbidity/mortality between day 10 and 18 post challenge (median survival = D13). Additionally, challenge of mice vaccinated with a non-alum-adjuvanted low dose vaccine derived from strain P6b resulted in high viremia on day 2 post challenge and a median survival day similar to the placebo control group, while mice vaccinated with a non-alum-adjuvanted low dose derived from clone e remained partially protected with a median survival of 19 days. These results indicate immunization is more effective with alum, secondary immunization may be a requirement, and that low dose was as effective as high dose.

Table 8: Serum neutralizing antibody titers

Pool	Serum neutralizing antibody titers	
	Terminal (post challenge)	
	$PRNT_{50}$	Secondary assay
Alum (1,2,4,5)	10240	20480
No alum (3,6)	2560	2560
PBS (7)	1280	1280

[0226] Additionally, the presence of NS1 in the vaccine drug substance (DS) produced from whole inactivated P7b and P7e virus (one additional passage from the P6b and P6e strains, respectively) was tested. A sandwich ELISA was performed using plates pre-coated with a monoclonal antibody reactive to both Asian and African lineages of Zika virus NS1, but non-cross-reactive to Dengue NS1. Duplicate 2-, 4-, 8-, 16-, and 32-fold dilutions of DS were prepared, and were compared to a standard curve using recombinant purified NS1 in duplicate at a concentration of 0-8 ng/mL. Duplicate dilutions of DS buffer alone were prepared as negative controls. Bound NS1 was detected with anti-NS1 HRP-conjugate, and absorbance (A_{450}-A_{630}) of the wells with DS buffer alone was subtracted from the absorbance measured in the wells containing the matching DS samples. Results of the sandwich ELISA are shown in **Table 9** below. Interestingly, NS1 was observed to co-purify with the vaccine drug substance preparations, suggesting that viral NS1 may be an immunogenic component of the whole inactivated virus vaccine.

Table 9: NS1 ELISA

Strain in vaccine preparation	Sample OD	Predicted log ng/mL	Std Error	Lower 95%	Upper 95%	Dilution Factor	Predicted concentration (ng/mL)
P7b	3.61	0.951	0.018	0.915	0.986	32	~285
P7e	3.79	0.980	0.023	0.935	1.024	32	~306

[0227] The threshold of neutralizing antibody (Nab) needed to confer protection from wild-type Zika virus challenge after passive transfer of antibodies was next tested. (**Tables 10A and B**).

Table 10A: design of passive transfer study in AG129 mice

Group	Test Article	Serum dilution	Predicted Nab titer before IP
1	100 µL	1/3	6827 / 3.83
2	100 µL	1/9	2276 / 3.36
3	100 µL	1/27	759 / 2.88
4	100 µL	1/81	253 / 2.40
5	100 µL	1/243	84 / 1.93
6	100 µL	1/729	28 / 1.45
7	100 µL	1/2187	9 / 0.97
8	100 µL	PBS	-

Table 10B: Timing of passive transfer study in AG129 mice

Description	Study Day
Passive transfer	Day 0
Primary Bleed (AM)	Day 1
Challenge (PM)	Day 1
Viremia Bleed	Day 4
Terminal Bleed	Day 29 for survivors

[0228] Pooled serum from vaccinated and challenged AG129 mice was serially diluted 3-fold in PBS and intraperitoneally injected into 7 groups (N=5/group) of 5-6 week old AG129 mice. Pre-immune AG129 mouse serum was used as placebo control (group 8). Following passive transfer (~16-19 hours later), whole blood was collected and serum was separated by centrifugation from each mouse prior to virus challenge for determination of circulating neutralizing antibody titer (**FIG. 17**). Just prior to virus challenge, groups of mice (designated groups 1, 2, 3, 4, 5, 6, 7, 8) had mean log10 neutralizing antibody titers of 2.69, 2.26, 1.72, 1.30, <1.30, <1.30, <1.30, <1.30, respectively.

[0229] Twenty four hours following passive transfer of ZIKV nAbs, mice were intraperitoneally challenged with 10^4 pfu of ZIKV PRVABC59. Following challenge, animals were weighed daily and monitored 1-3 times a day for 28 days for signs of illness. A clinical score was given to each animal based on the symptoms (**Table 11**). Animals that were moribund and/or showed clear neurological signs (clinical score ≥2) were humanely euthanized and counted as non-survivors.

Table 11: Description of clinical scores given while monitoring for morbidity and mortality

Score	Description
0	Normal appearance and behavior
1	Slightly ruffled fur and/or general loss of condition
2	Increases in above behavior/appearance, breathing changes, twitching, anti-social behavior
3	First signs of neuropathy – Severely hunched posture, partial paralysis (immobility, unsteady gait, flaccid hind legs, severe twitching), or full paralysis
4	Found dead without showing signs of score of 2 or 3 first

[0230] Signs of disease began appearing nine days after challenge in the control group (group 8) and groups 5-7, with a corresponding loss in weight (**FIG. 18**). Whole blood was collected and serum was separated by centrifugation from each animal three days post challenge. Serum samples were analyzed for the presence of infectious ZIKV using a plaque titration assay (**FIG. 19**). The mean infectious titer (log_{10} pfu/mL) for mice in groups 1-8 were: 1.66, 2.74, 4.70, 4.92, 7.24, 7.54, 7.54 and 7.46, respectively. Importantly, mice in groups 1-4 with detectable levels of ZIKV neutralizing antibodies ($\geq$1.30 log_{10}) had statistically significant lower levels (102.5- to 106.0- fold lower titers) of viremia (p = 0.0001, 0.0003, 0.0007 and 0.0374) than control mice. These results suggested that detectable levels of ZIKV neutralizing antibodies ($\geq$1.30 log10) reduced viremia in a dose-dependent manner.

[0231] The median survival day of mice in groups 1-8 were: not determined, day 17, day 17, day 13, day 11, day 11, day 11, and day 10, respectively (**FIG. 20**). Importantly, the survival curves for groups of mice with detectable ZIKV neutralizing antibody titers (groups 1-4) were statistically different compared to the control group (group 8) (p = 0.0019, 0.0019, 0.0019, 0.0153, respectively). These results suggested that detectable levels ($\geq$1.30 log_{10}) of ZIKV neutralizing antibodies delayed onset of disease in a dose-dependent manner.

[0232] Finally, the ZIKV neutralizing antibody titer of each animal was graphed against its corresponding viremia titer and linear regression analysis was performed. A highly inversely correlated relationship between ZIKV neutralizing antibody titers and viremia levels at day 3 post-challenge was observed (**FIG. 21**). A summary of the results from the passive transfer studies is shown in **Table 12** below.

Table 12: Summary of passive transfer results

Group	Serum dilution	Circulating ZIKV nAb GMT	Viremia (D3) log10 pfu/mL	% survival (D28)	Median survival day
1	1/3	2.69 ± 0.17	1.66 ± 0.62	20	24
2	1/9	2.26 ± 0.13	2.73 ± 0.68	0	17
3	127	1.72 ± 0.16	4.69 ± 0.77	0	17
4	1/81	1.30 ± 0.16	4.94 ± 1.29	0	13
5	1/243	<1.30	7.25 ± 0.10	0	11
6	1/729	<1.30	7.54 ± 0.31	0	11
7	1/2187	<1.30	7.52 ± 0.39	0	11
8	PBS	<1.30	7.47 ± 0.37	0	10

[0233] While no groups of mice receiving ZIKAV neutralizing antibodies were fully protected from lethal ZIKAV challenge in this experiment, reduced viremia levels and delayed onset of disease in a dose-dependent manner among the groups of mice with detectable levels of circulating ZIKAV neutralizing antibody titers was demonstrated.

[0234] Taken together, preclinical data from both CD-1 and AG129 mouse studies indicate that a PIZV derived from separate and well-characterized viral clones are immunogenic and able to provide protection against challenge with wild-type ZIKAV. Importantly, a low and high vaccine dose elicited a similar neutralizing antibody response after two doses, and provided similar levels of protection against lethal ZIKAV challenge. Interestingly, mice vaccinated with an unadjuvanted PIZV candidate also showed partial protection from ZIKAV challenge. Vaccine antisera significantly diminished viremia in passively immunized AG129 mice, and prolonged survival against lethal ZIKAV challenge. These results also demonstrate that the well-characterized PIZV candidates were highly efficacious against ZIKAV infection in the highly ZIKAV-susceptible AG129 mouse model.

[0235] Additionally, it was found that the sequence of a PRVABC59 (from PRVABC59 P6e) at passage 7 was genetically identical to that of passage 6. This was surprising given that flaviviruses are generally regarded as genetically labile. PRVABC59 P6e was selected as the master virus seed due in part to its genetic stability over 7 passages. Without wishing to be bound by theory, it is believed that this enhanced genetic stability may be due to the single amino acid substitution (W98G) in the wing domain of NS1, as this was the only mutation

observed in the Vero cell-adapted PRVABC59 P6 genome. Additionally, genetic stability and homogeneity is advantageous in that it reduces variability and increases reproducible production of subsequent strains that may be used for vaccine formulation.

CLAIMS

What is claimed is:

1. A vaccine or immunogenic composition comprising one or more antigens from a Zika virus, wherein the Zika virus comprises at least one non-human cell adaptation mutation.

2. The vaccine or immunogenic composition of claim 1, wherein the at least one non-human cell adaptation mutation is in Zika virus Non-structural protein 1 (NS1).

3. The vaccine or immunogenic composition of claim 2, wherein the at least one adaptation mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1.

4. The vaccine or immunogenic composition of claim 3, wherein the at least one adaptation mutation is a Trp98Gly mutation.

5. The vaccine or immunogenic composition of any one of claims 1-4, wherein the at least one adaptation mutation enhances genetic stability as compared to a Zika virus lacking the at least one adaptation mutation.

6. The vaccine or immunogenic composition of any one of claims 1-5, wherein the at least one adaptation mutation enhances viral replication as compared to a Zika virus lacking the at least one adaptation mutation.

7. The vaccine or immunogenic composition of any one of claims 1-6, wherein the Zika virus does not comprise a mutation in Envelope protein E (Env).

8. The vaccine or immunogenic composition of any one of claims 1-7, wherein the non-human cell is a mammalian cell.

9. The vaccine or immunogenic composition of any one of claims 1-8, wherein the non-human cell is a monkey cell.

10. The vaccine or immunogenic composition of claim 9, wherein the monkey cell is from a Vero cell line.

11. The vaccine or immunogenic composition of claim 10, wherein the Vero cell line is a WHO Vero 10-87 cell line.

12. The vaccine or immunogenic composition of any one of claims 1-11, wherein the Zika virus is an African lineage virus or an Asian lineage virus.

13. The vaccine or immunogenic composition of any one of claims 1-11, wherein the Zika virus is an Asian lineage virus.

14. The vaccine or immunogenic composition of claim 13, wherein the Zika virus is from strain PRVABC59.

15. The vaccine or immunogenic composition of any one of claims 1-14, wherein the vaccine or immunogenic composition is a purified antigen vaccine or immunogenic composition, a subunit vaccine or immunogenic composition, an inactivated whole virus vaccine or immunogenic composition, or an attenuated virus vaccine or immunogenic composition.

16. The vaccine or immunogenic composition of any one of claims 1-15, wherein the vaccine or immunogenic composition is an inactivated whole virus vaccine or immunogenic composition.

17. The vaccine or immunogenic composition of any one of claims 1-16, wherein the virus was chemically inactivated.

18. The vaccine or immunogenic composition of claim 17, wherein the virus was chemically inactivated with one or more of a detergent, formalin, beta-propiolactone (BPL), binary ethylamine (BEI), acetyl ethyleneimine, methylene blue, and psoralen.

19. The vaccine or immunogenic composition of claim 17, wherein the virus was chemically inactivated with formalin.

20. The vaccine or immunogenic composition of any one of claims 1-19, further comprising an adjuvant.

21. The vaccine or immunogenic composition of claim 20, wherein the adjuvant is selected from the group consisting of aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions,

virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and Incomplete Freund's Adjuvant (IFA).

22. The vaccine or immunogenic composition of claim 20, wherein the adjuvant is an aluminum salt.

23. The vaccine or immunogenic composition of claim 20, wherein the adjuvant is selected from the group consisting of alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85.

24. The vaccine or immunogenic composition of any one of claims 20-23, wherein at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of the one or more antigens are adsorbed to the adjuvant.

25. The vaccine or immunogenic composition of any one of claims 1-19, comprising from about 0.1 μg Env to about 100 μg Env.

26. The vaccine or immunogenic composition of claim 25, wherein the vaccine or immunogenic composition is unadjuvanted.

27. The vaccine or immunogenic composition of any one of claims 1-26, wherein the Zika virus is a clonal isolate.

28. The vaccine or immunogenic composition of claim 27, wherein the clonal isolate is substantially free of one or more adventitious agents.

29. A vaccine or immunogenic composition comprising: a) an aluminum salt adjuvant; and b) a whole inactivated Zika virus, wherein the Zika virus comprises a non-human cell adaptation mutation, and wherein the non-human cell adaptation mutation is a Trp98Gly mutation at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1.

30. A method of treating or preventing Zika virus infection in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of the vaccine or immunogenic composition of any one of claims 1-29.

31. A method for inducing an immune response in a subject in need thereof, comprising administering to the subject an immunogenic amount of the vaccine or immunogenic composition of any one of claims 1-29.

32. The method of claim 30 or claim 31, wherein the subject is a human.

33. The method of any one of claims 30-32, wherein the subject is pregnant or intends to become pregnant.

34. The method of any one of claims 30-33, wherein administration of the vaccine or immunogenic composition induces a protective immune response in the subject.

35. The method of claim 34, wherein the protective immune response induced in the subject is greater than a protective immune response induced in a corresponding subject administered a vaccine or immunogenic composition comprising one or more antigens from a Zika virus lacking the at least one non-human cell adaptation mutation.

36. The method of any one of claims 30-33, wherein administration of the vaccine or immunogenic composition induces the generation of neutralizing antibodies to Zika virus in the subject.

37. The method of claim 36, wherein the concentration of neutralizing antibodies generated in the subject is higher than a concentration of neutralizing antibodies generated in a corresponding subject administered a vaccine or immunogenic composition comprising one or more antigens from a Zika virus lacking the at least one non-human cell adaptation mutation.

38. The method of any one of claims 30-37, wherein the vaccine or immunogenic composition is administered by a route selected from the group consisting subcutaneous administration, transcutaneous administration, intradermal administration, subdermal administration, intramuscular administration, peroral administration, intranasal administration, buccal administration, intraperitoneal administration, intravaginal administration, anal administration and intracranial administration.

39. The method of any one of claims 30-38, wherein the vaccine or immunogenic composition is administered one or more times.

40. The method of claim 39, wherein the vaccine or immunogenic composition is administered as a first (prime) and a second (boost) administration.

41. The method of claim 40, wherein the second (boost) administration is administered at least 28 days after the first (prime) administration.

42. A method for inactivating a Zika virus preparation comprising:

(a) isolating the Zika virus preparation from one or more non-human cells, wherein the cells are used to produce the virus preparation, and wherein the Zika virus comprises at least one non-human cell adaptation mutation; and

(b) treating the virus preparation with an effective amount of formalin.

43. The method of claim 42, further comprising (c) neutralizing the formalin-treated virus preparation with sodium metabisulfite.

44. The method of claim 43, wherein the virus preparation is neutralized at least five, at least seven, at least nine, at least 11, or at least 14 days after formalin treatment.

45. The method of any one of claims 42-44, further comprising (d) purifying the neutralized virus preparation.

46. The method of claim 45, wherein the neutralized virus preparation is purified by process selected from the group consisting of cross flow filtration (CFF), multimodal chromatography, size exclusion chromatography, cation exchange chromatography, and anion exchange chromatography.

47. The method of any one of claims 42-46, wherein the virus preparation is mixed with an adjuvant.

48. The method of claim 47, wherein the adjuvant is selected from the group consisting of aluminum salts, toll-like receptor (TLR) agonists, monophosphoryl lipid A (MLA), synthetic lipid A, lipid A mimetics or analogs, MLA derivatives, cytokines, saponins, muramyl dipeptide (MDP) derivatives, CpG oligos, lipopolysaccharide (LPS) of gram-negative bacteria, polyphosphazenes, emulsions, virosomes, cochleates, poly(lactide-co-glycolides) (PLG) microparticles, poloxamer particles, microparticles, liposomes, Complete Freund's Adjuvant (CFA), and Incomplete Freund's Adjuvant (IFA).

49. The method of claim 47, wherein the adjuvant is an aluminum salt.

50. The method of claim 47, wherein the adjuvant is selected from the group consisting of alum, aluminum phosphate, aluminum hydroxide, potassium aluminum sulfate, and Alhydrogel 85.

51. The method of any one of claims 47-50, wherein at least 75%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% of one or more antigens in the virus preparation are adsorbed to the adjuvant.

52. The method of any one of claims 42-51, wherein the at least one non-human cell adaptation mutation is in Zika virus NS1.

53. The method of claim 52, wherein the at least one adaptation mutation occurs at position 98 of SEQ ID NO: 1, or at a position corresponding to position 98 of SEQ ID NO: 1.

54. The method of claim 53, wherein the at least one adaptation mutation is a Trp98Gly mutation.

55. The method of any one of claims 42-54, wherein the at least one adaptation mutation enhances genetic stability as compared to a Zika virus lacking the at least one adaptation mutation.

56. The method of any one of claims 42-55, wherein the at least one adaptation mutation enhances viral replication as compared to a Zika virus lacking the at least one adaptation mutation.

57. The method of any one of claims 42-56, wherein the Zika virus does not comprise a mutation in Envelope protein E (Env).

ABSTRACT

The present disclosure relates to Zika virus vaccines and immunogenic compositions having one or more antigens from a non-human cell adapted Zika virus, and methods of manufacture, formulation, testing, and uses thereof.

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