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Natural history of Asian lineage Zika virus infection in macaques

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Infection with Asian lineage Zika virus has been associated with Guillain-Barré syndrome and fetal microcephaly[1-4]. Here we show that rhesus macaques are susceptible to infection by an Asian lineage Zika virus isolate that shares more than 99% nucleotide identity with strains currently circulating in the Americas. Following subcutaneous inoculation, Zika virus RNA was detected in plasma one-day post infection (dpi) in all animals (N = 3). Plasma viral loads peaked above 1 x 10^6 viral RNA copies/mL in two of three animals. Viral RNA was also present in saliva, urine, and cerebrospinal fluid (CSF), consistent with case reports from infected humans. Zika virus RNA persisted in both

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plasma and urine at low levels for more than two weeks. Infection was associated with transient increases in proliferating natural killer cells, CD8+ and CD4+ T cells, and plasmablasts, suggesting pathogen sensing by the immune system. These data establish that Asian lineage Zika virus infection in rhesus macaques provides a relevant animal model for human infection. Furthermore, because fetal development is well characterized in rhesus macaques, infections in pregnant macaques will enable important studies of fetal defects associated with Zika virus.

Zika virus (ZIKV) is a mosquito-borne flavivirus first identified in 1947 that is divided into African and Asian lineages based on phylogeny and geographical origin[5]. The primary vector for ZIKV is *Aedes aegypti*, the same species that transmits dengue (DENV) and chikungunya (CHIKV) viruses[6]. Little was known about ZIKV when increased incidence of fetal microcephaly and Guillain-Barré syndrome occurred coincident with epidemic spread of Asian lineage ZIKV in South America. The virologic and immunologic parameters of ZIKV infection remain unknown, including tissue tropism, duration of infection, and the potential for persistence. To date, ZIKV RNA has been detected in plasma, urine, semen, CSF, breast milk and saliva of patients[4,7-10], but the extent to which this represents replication-competent, transmissible virus is unknown. Infectious virus has been isolated from saliva and a small number of case reports suggest that ZIKV may be transmitted by blood transfusion or sexual intercourse, but the likelihood of such non-vectored transmission remains undetermined[11-13]. These and many other questions cannot be resolved quickly by studying case reports of infected individuals alone.

We therefore sought to develop a relevant animal model to understand ZIKV pathogenesis, and to evaluate candidate vaccines and therapeutics. For decades, macaques have been used in both infectious disease and obstetric research. In fact, what is now known as the African lineage of ZIKV was initially discovered in a sentinel rhesus macaque kept in the Zika forest of Uganda for yellow fever surveillance[14]. But does the currently circulating Asian

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lineage virus, which shares ~90% nucleotide identity with the African lineage, also infect macaques[5]?

To determine this, we inoculated three Indian-origin rhesus macaques (*Macaca mulatta*) subcutaneously with 1 x 10^6, 1 x 10^5, or 1 x 10^4 PFU ZIKV derived from a French Polynesian virus isolate (Zika virus/H.sapiens-tc/FRA/2013/FrenchPolynesia-01_v1c1) (Fig. 1a). This range of inocula is based on previous work in related flaviviruses such as West Nile virus (WNV) and DENV, where it was estimated that mosquitoes delivered 1 x 10^4- 1x 10^6 PFU of virus[15]. DENV infection of macaques also provides the precedence for using subcutaneous inoculations[16]. Blood was sampled daily for 11 days post-infection (dpi) and every 3-4 days thereafter. Viral RNA (vRNA) was quantified by qRT-PCR[17]. Plasma viremia was detected in all three animals at 1 dpi and was proportional to the inoculum dose, underscoring the importance of selecting a ZIKV challenge dose in macaques that approximates the dose encountered during either vector-borne or sexual transmission (Fig. 1b)[8,11]. Peak plasma viremia occurred between 2 and 6 dpi, and ranged from 8.2 x 10^4 to 2.5 x 10^6 vRNA copies/mL. The estimated doubling time for the plasma viremia of individual animals ranged from 4.8 to 10.2 hours and appeared to be independent of the infecting dose. By 10 dpi, plasma viral loads were undetectable (<100 copies/mL) in all three animals, although intermittent low-level detection (<550 copies/mL) continued in two of three animals through 17 dpi.

To detect signs of morbidity, animals were evaluated daily for evidence of disease, injury, or psychological abnormalities (e.g., inappetence, dehydration, diarrhea, depression, inactivity, trauma, self-injurious or stereotypical behavior). All animals exhibited mild to moderate inappetence, which led to mild weight loss in two animals. Two animals (912116 and 393422) also developed a very mild rash around the inoculation site one dpi that persisted for 4-5 days. No other abnormal clinical signs were noted (e.g., increased body temperature, joint pain, lymphadenopathy, lethargy).

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To examine potential within-host evolution, we deep sequenced both the challenge stock and plasma virus from each animal at 4 dpi. Like all flaviviruses, the ZIKV genome encodes a single, large polyprotein that is co- and post-translationally processed into three structural (envelope E, membrane precursor prM, capsid C) and seven non-structural (NS1, NS2A, NS2B, NS3, NS4A, NS4B, and NS5) proteins[18]. The ZIKV challenge stock consensus sequence matched the Genbank sequence (KJ776791) of the parental virus, but there were 8 sites where between 5-40% of sequences contained variants (Fig. 2a). Maintenance of some of the challenge stock variants, as well as de novo variants not detected in the challenge stock, were noted in all three animals at 4 dpi (Fig. 2b). Variants were concentrated in the NS1, NS2A and NS2B regions for the both the challenge stock and from the animals.

To measure how selection may be shaping the viral populations during replication in macaques, we calculated measures of synonymous (πS) and nonsynonymous (πN) nucleotide diversity in each coding region of the genome relative to the inoculum for each animal at 4 dpi. The ratio of πS to πN can indicate how selection is acting on a gene. While most coding regions revealed signatures of purifying selection, the regions encoding NS1 and NS2B exhibited elevated levels of nonsynonymous diversity in all three animals (Fig. 2c). NS1 exists in multiple forms, including a membrane--bound mNS1 form and a secreted sNS1 form. These forms are implicated in a number of flavivirus--host interactions, including immune protection, disease pathogenesis, and evasion of the complement system[19,20]. Thus, diversification in NS1 genes could be the result of selective pressures imposed by the macaque host. Further study is required to determine whether NS1 and NS2B are consistently under positive selection in macaques, and to identify candidate selective pressures. However, our data suggest that they may be good targets for future studies of ZIKV immunity and host adaptation.

Animals were monitored using complete blood counts and blood chemistry panels daily for 11 dpi and then every 3 to 4 days thereafter (Extended Data Fig. 1). Similar to other viral infections, a transient lymphopenia was observed beginning 1 dpi, reaching a nadir 4 dpi

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(Extended Data Fig. 1f). The animal that received the highest dose of inoculum developed leucocytosis at 7 dpi that persisted through 28 dpi and was characterized by a mature neutrophilia. Each animal also exhibited evidence of a mild regenerative anemia characterized by varying degrees of polychromasia and anisocytosis, but whether this was secondary to the viral infection or simply a result of frequent blood draws could not be determined. All animals developed an elevated serum creatine kinase (CK), which correlated directly with inoculum dose and peaked 5 dpi (Extended Data Fig. 1). Increased CK is often noted in animals after repeated sedation and venipuncture. Two of three animals also exhibited an approximate two-fold increase in AST and ALT, which peaked at 5 dpi. These findings are not uncommon in acute viral infections in humans. Both cellular dyscrasias and elevated transaminases have been described in human ZIKV case reports; myositis has not been reported[21,22].

We next characterized the immune response to infection in lymphocyte subsets by staining peripheral blood mononuclear cells (PBMC) for multiple lineage and activation markers. Proliferating (Ki-67+) NK cells, CD8+ T cells, and CD4+ T cells expanded significantly above baseline levels by 5 dpi; the most rapid increase occurred in the animal challenged with the highest ZIKV dose (Fig. 3). We also enumerated circulating plasmablasts, defined as CD3-/20-/14-/16-/11c-/123- and CD80+/HLA-DR+ cells, on 3, 7, 11 and 14 dpi (Fig. 3)[23]. The peak plasmablast expansion was inoculum-dose-dependent by day 7 and the decline in plasmablast cells was inversely related to inoculum dose. Together these data provide an estimate that peak activation of the adaptive immune response and antibody production may occur 5-7 dpi and depend on inoculum dose, informing future work to characterize these responses.

In addition to blood, several other body fluids were tested for ZIKV by qRT-PCR including urine, saliva and CSF. Viruria was detected starting at 4 or 5 dpi and continued up to 14 dpi, in urine passively collected from cage pans (Fig. 1b). Despite possible degradation of virus between the time of urination and sample collection and processing, 1 x 10^3-1 x 10^4 vRNA copies/mL urine were detected at multiple timepoints. Virus was also detected in oral swabs

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collected from all three animals, peaking at approximately 1 x 10^4 vRNA copies per sample in 2 of 3 animals (Fig. 1b). Notably, as with urine, the kinetics of virus detection in saliva occurred after peak plasma viremia. Lumbar punctures were performed at 4 and 14 dpi to quantify viral RNA in CSF; vRNA was detectable at 4 dpi in two of three animals (Fig. 1b).

Altogether, our study showed the persistence of ZIKV in the plasma of rhesus macaques for approximately 10 days, similar to other vector-borne flaviviruses that cause acute, typically self-limiting infections in humans. However, the prolonged detection of vRNA in urine and saliva after apparent clearance from the blood, detection of virus in the CSF, and occasional plasma "blips" after initial clearance, suggest that ZIKV could establish a more persistent infection. Given reports of neurologic complications arising from ZIKV infection, the central nervous system may be a reservoir site for persistent virus. Future work in rhesus macaques will seek to determine whether virus from fluids other than blood can infect new hosts and what threshold of viral load at these sites may be required for transmission.

While Asian lineage viruses are associated with fetal microcephaly or Guillain-Barré syndrome, African lineage ZIKV is not, suggesting that the lineages may have important biological differences. To begin to investigate this possibility, we also infected three macaques with the African lineage Zika virus/R.macaque-tc/UGA/1947/MR766-3329, using the same three doses as the Asian lineage infections (Extended Data Fig. 2). Although all three animals became infected, viremia was only detected in two animals 1 dpi. The African lineage virus achieved peak plasma levels at least one order of magnitude lower those observed in animals infected with the Asian-lineage virus (Extended Data Fig. 2). This difference in *in vivo* viral kinetics could be relevant for both virus transmission and pathogenesis. We are currently working to complete the characterization of these infections.

Our study establishes the rhesus macaque as a relevant translational model for ZIKV infection and pathogenesis in humans. Rhesus macaques support replication of both Asian lineage and African lineage ZIKV, providing the possibility to conduct detailed comparative

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pathogenesis studies. The large immunological toolset available for rhesus macaques will enable investigations of immunity and potential vaccines. Pregnancy, the maternal-fetal interface, and fetal development have been described in detail in rhesus macaques, so this model will also enable assessments of the impact of maternal ZIKV infection on the developing fetus.

Methods

Study design

This was a proof-of-concept study was designed to establish the infectivity and viral dynamics of Asian lineage ZIKV. Because nothing is known about ZIKV dosing in macaques, one male rhesus macaque of Indian ancestry was challenged with each of the following ZIKV doses: $1x10^6$, $1x10^5$, and $1x10^4$ PFU ZIKV from each lineage. All macaques utilized in the study were free of Macacine herpesvirus 1, Simian Retrovirus Type D, Simian T-lymphotropic virus Type 1, and Simian Immunodeficiency Virus. In a follow-up study, two additional males and one female macaque were infected with the African lineage ZIKV at the same three challenge doses as the Asian lineage. Primary data from the study is available at goo.gl/rmNCqf.

Care and use of macaques at the Wisconsin National Primate Research Center

All macaque monkeys used in this study were cared for by the staff at the Wisconsin National Primate Research Center (WNPRC) in accordance with the regulations and guidelines outlined in the Animal Welfare Act and the Guide for the Care and Use of Laboratory Animals and the recommendations of the Weatherall report. This study was approved by the University of Wisconsin-Madison Graduate School Institutional Animal Care and Use Committee (Animal Care and Use Protocol Number G005401). For all procedures (i.e., physical examination, virus inoculation, blood and swab collection), animals were anesthetized with an intramuscular dose

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of ketamine (10 mL/kg). Blood samples were obtained using a vacutainer system or needle and syringe from the femoral or saphenous vein.

Inoculations

ZIKV strain H/PF/2013 (GenBank: KJ776791), originally isolated from a 51-year-old female in France returning from French Polynesia with a single round of amplification on Vero cells, was obtained from Xavier de Lamballerie (European Virus Archive, Marseille France). ZIKV strain Zika virus/R.macaque-tc/UGA/1947/MR766-3329 (GenBank:LC002520), originally isolated from a sentinel Rhesus monkey on 20 April 1947 in Zika Forest, Entebbe, Uganda with 149 suckling mouse brain passages and two rounds of amplification on Vero cells, was obtained from Brandy Russell (CDC, Ft. Collins, CO). Virus stocks were prepared by inoculation onto a confluent monolayer of C6/36 mosquito cells. A single harvest of virus with a titer of 1.26×10^6 PFU/mL for the Asian lineage (equivalent to 1.43×10^9 vRNA copies/mL) and 5.9×10^6 PFU/mL for the African lineage was used for all three challenges of each lineage. The stock was thawed, diluted in PBS to the appropriate concentration for each challenge, and loaded into a 1 mL syringe that was kept on ice until challenge. Animals were anesthetized as described above, and 1 mL of inocula was administered subcutaneously over the cranial dorsum. At the conclusion of the procedure, animals were closely monitored by veterinary and animal care staff for adverse reactions and signs of disease.

Viral RNA isolation from plasma

Fresh plasma and PBMC were isolated from EDTA-treated whole blood by Ficoll density centrifugation at 1860 rcf for 30min. The plasma layer was collected and centrifuged for an additional 8 min at 670 rcf to remove residual cells. RNA was extracted from 300 µl of plasma

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using the Viral Total Nucleic Acid Purification Kit (Promega, Madison, WI) on a Maxwell 16 MDx instrument. The RNA was then quantified by quantitative RT-PCR.

Viral RNA isolation from urine

Urine was collected from a pan beneath the animal's cage. Urine was centrifuged for 5 min at 500 rcf to remove cells and other debris. RNA was isolated from 300 µl urine using the Viral Total Nucleic Acid Purification Kit (Promega, Madison, WI) on a Maxwell 16 MDx instrument.

Viral RNA isolation from oral swabs

Oral swab samples were collected from infected animals while anesthetized by gently running a sterile swab under the animal's tongue. Swabs were placed immediately into either RNAlater or viral transport medium (tissue culture medium 199 supplemented with 0.5% FBS and 1% antibiotic/antimycotic) for 60-90 minutes. Samples were vortexed vigorously, then centrifuged for 5 min at 500 rcf before removing the swabs. Samples were stored at either -20°C (RNAlater samples) or -80°C (viral transport medium) until processing. Prior to extraction, virus was pelleted by centrifugation for 1 hour at 4°C at 14000 rpm. Supernatant was removed, leaving the virus in 200 µl media. Viral RNA was extracted from these samples using the Qiamp MinElute Virus Spin kit (Qiagen, Germantown, Maryland) with all optional washes. Viral load data from oral swabs are expressed as vRNA copies/mL eluate.

Quantitative reverse transcription PCR (qRT-PCR)

Viral RNA isolated from plasma, urine, or oral swabs was quantified by qRT-PCR using the primers and probe designed by Lanciotti et al.[17] or using primers with a slight modification to those described by Lanciotti et al. to accommodate the African lineage virus. The sequence of these modified primers are as follows: forward-5'CGYTGCCCAACACAAGG-3', reverse 5'-

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CCACYAAYGTTCTTTTGCABACAT-3', and probe 5'-6fam-AGCCTACCTTGAYAAGCARTCAGACACYCAA-BHQ1-3'. The RT-PCR was performed using the SuperScript III Platinum one-step quantitative RT-PCR system (Invitrogen, Carlsbad, CA) on the LightCycler 480 instrument (Roche Diagnostics, Indianapolis, IN). Primers and probe were used at final concentrations of 600 nm and 100 nm respectively, along with 150 ng random primers (Promega, Madison, WI). Cycling conditions were as follows: 37°C for 15 min, 50°C for 30 min and 95°C for 2 min, followed by 50 cycles of 95°C for 15 sec and 60°C for 1 min. Virus concentration was determined by interpolation onto an internal standard curve composed of seven 10-fold serial dilutions of a synthetic ZIKV RNA fragment based on the Asian lineage. A comparison of the crossing point detected by qRT-PCR found from the standard template, Asian lineage virus and African lineage virus when using either the universal primer set developed by our group, or the primers developed by Lanciotti et al., suggest that the efficiency of each primer set is the same for both lineages of ZIKV (Extended Data Fig. 3).

Viral quantification by plaque assay

All ZIKV screens and titrations for virus quantification were completed by plaque assay on Vero cell cultures. Duplicate wells were infected with 0.1 mL aliquots from serial 10-fold dilutions in growth media and virus was adsorbed for one hour. Following incubation, the inoculum was removed, and monolayers were overlaid with 3 ml containing a 1:1 mixture of 1.2% oxoid agar and 2X DMEM (Gibco, Carlsbad, CA) with 10% (vol/vol) FBS and 2% (vol/vol) penicillin/streptomycin. Cells were incubated at 37°C in 5% CO_2 for four days for plaque development. Cell monolayers then were stained with 3 mL of overlay containing a 1:1 mixture of 1.2% oxoid agar and 2X DMEM with 2% (vol/vol) FBS, 2% (vol/vol) penicillin/streptomycin, and 0.33% neutral red (Gibco). Cells were incubated overnight at 37°C and plaques were counted.

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253

254 *Immunophenotyping*

255 The amount of activated/proliferating NK cells were quantified using a modified version of our
256 protocol detailed step-by step in OMIP-28[24]. Briefly 0.1 mL of EDTA-anticoagulated whole blood
257 samples were incubated for 15 min at room temperature in the presence of a mastermix of
258 antibodies against CD45 (clone D058-1283, Brilliant Violet 786 conjugate), CD3 (clone SP34-2
259 Alexa Fluor 700 conjugate), CD8 (clone SK2, Brilliant Violet 510), NKG2A/C (clone Z199, PE-
260 Cy7 conjugate), CD16 (clone 3G8, Pacific Blue conjugate), CD69 (clone TP1.55.3, ECD
261 conjugate), HLA-DR (clone 1D11, Brilliant Violet 650 conjugate), CD4 (clone SK3, Brilliant Violet
262 711 conjugate), CCR7 (clone 150503, Fluorescein conjugate), CD28 (clone CD28.2, PE
263 conjugate), and CD95 (clone DX2, PE-Cy5 conjugate) antigens. All antibodies were obtained
264 from BD BioSciences, except the NKG2A/C-specific antibody, which was purchased from
265 Beckman Coulter and the CCR7 antibody that was purchased from R&D Systems. Red blood
266 cells were lysed using BD Pharm Lyse, after which they were washed twice in media and fixed
267 with 0.125 ml of 2% paraformaldehyde for 15 min. After an additional wash the cells were
268 permeabilized using Life Technology's Bulk Permeabilization Reagent. The cells were stained
269 for 15 min. against Ki-67 (clone B56, Alexa Fluor 647 conjugate) while the permeabilizer was
270 present. The cells were then washed twice in media and resuspended in 0.125 ml of 2%
271 paraformaldehyde until they were run on a BD LSRII Flow Cytometer. Flow data were analyzed
272 using Flowjo version 9.8.2.

273

274 *Plasmablast detection*

275 Peripheral blood mononuclear cells (PBMCs) isolated from three ZIKV-infected rhesus monkeys
276 at 3, 7, 11, and 14 dpi were stained with the following panel of fluorescently labeled antibodies
277 (Abs) specific for the following surface markers: CD20 FITC (L27), CD80 PE(L307.4), CD123

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PE-Cy7(7G3), CD3 APC-Cy7 (SP34-2), IgG BV605(G18-145) (all from BD Biosciences, San Jose, CA), CD14 AF700 (M5E2), CD11c BV421 (3.9), CD16 BV570 (3G8), CD27 BV650(O323) (all from BioLegend, San Diego, CA), IgD AF647 (polyclonal)(Southern Biotech, Birmingham, AL), and HLA-DR PE-TxRed (TÜ36) (Invitrogen, Carlsbad, CA). LIVE/DEAD Fixable Aqua Dead Cell Stain Kit (Invitrogen, Carlsbad, CA) was used to discriminate live cells. Briefly, cells were resuspended in 1X PBS/1%BSA and stained with the full panel of surface Abs for 30 min in the dark at 4°C, washed once with 1X PBS, stained for 30 min with LIVE/DEAD Fixable Aqua Dead Cell Stain Kit in the dark at 4°C, washed once with 1X PBS, washed again with 1X PBS/1%BSA, and resuspended in 2% PFA Solution. Stained PBMCs were acquired on a LSRII Flow Analyzer (BD Biosciences, San Jose, CA) and the data was analyzed using FlowJo software v9.7.6 (TreeStar, Ashland, OR). Plasmablasts were defined similarly to the method previously described[23] excluding lineage cells (CD14+, CD16+, CD3+, CD20+, CD11c+, CD123+), and selecting CD80+ and HLA-DR+ cells (known to be expressed on rhesus plasmablasts and their human counterpart[25]).

Estimation of plasma viremia doubling time

The doubling time of plasma viremia was estimated in R version 3.2.3 (The R Foundation for Statistical Computing, http://www.R-project.org). For each animal, the slope of the linear portion of the line (between 1 and 2 dpi for the animals treated with $1x10^6$ and $1x10^5$ PFU and between 1, 2, and 3 dpi for the animal treated with $1x10^4$ PFU) was generated by plotting the log of the plasma viral loads. The linear portion represents the exponential growth phase and has been used to estimate doubling time in other systems[26]. The slopes were then used in the equation: log(2)/slope. Each result was then multiplied by 24 hours to produce a simple estimate of doubling time in hours.

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CBC and blood chemistry panels

CBCs were performed on EDTA-anticoagulated whole blood samples on a Sysmex XS-1000i automated hematology analyzer (Sysmex Corporation, Kobe, Japan). Blood smears were prepared and stained with Wright-Giemsa stain (Wescor Aerospray Hematology Slide Stainer; Wescor Inc, Logan, UT). Manual slide evaluations were performed on samples as appropriate when laboratory-defined criteria were met (including the presence of increased total white blood cell counts, increased monocyte, eosinophil, and basophil percentages, decreased hemoglobin, hematocrit, and platelet values, and unreported automated differential values). Individuals performing manual slide evaluations screened both white blood cells (WBC) and red blood cells (RBC) for cellular maturity, toxic change, and morphologic abnormalities.

Whole blood was collected into serum separator tubes (Becton, Dickinson and Company, Franklin Lakes, NJ) for blood chemistry analysis and processed per manufacturer's instructions. Blood chemistry panels were performed on the serum using a Cobas 6000 analyzer (Roche Diagnostics, Risch-Rotkreuz, Switzerland). Results from CBC and blood chemistry panels were reported with species, age, and sex-specific reference ranges.

Zika virus deep sequencing from plasma

A vial of the same ZIKV strain H/PF/2013 virus stock that infected macaques was deep sequenced by preparing libraries of fragmented double-stranded cDNA using methods similar to those previously described[27]. Briefly, the sample was centrifuged at 5000 rcf for 5 min. The supernatant was then filtered through a 0.45-um filter. The Qiagen QiAmp Minelute viral RNA isolation kit (omitting carrier RNA) was used to isolate vRNA. The eluted RNA was then treated with DNAse I. Double stranded DNA was prepared with the Superscript double stranded cDNA synthesis kit (Invitrogen) and priming with random hexamers. Agencourt Ampure XP beads were used to purify double stranded DNA. The purified DNA was fragmented with the Nextera XT kit (Illumina), tagged with Illumina-compatible primers, and then purified with Agencourt

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Ampure XP beads. Purified libraries were then sequenced with 2 x 300 bp kits on an Illumina MiSeq.

Zika virus samples isolated from animals were PCR amplified to create a series of five overlapping amplicons that span the entire viral polyprotein, as described previously for other RNA viruses[28]. Briefly, vRNA was isolated using the Qiagen QiAmp Minelute viral RNA isolation kit, according to manufacturer's protocol (Qiagen). The following primer pairs were used to generate viral cDNA amplicons using the Superscript III One-step RT-PCR kit (Invitrogen) and including MgSO4:

ZFP-1F: GCGACAGTTCGAGTTTGAAGCG; ZFP-1R: ATCCAAAGTCCCAGGCTGTG;

ZFP-2F: AGATCCCGGCTGAAACACTG; ZFP-2R: CCCATGTGATGTCACCTGCT;

ZFP-3F: TACTCACAGCTGTTGGCCTG; ZFP-3R: CACCTCGGTTTGAGCACTCT;

ZFP-4F: TGTTTGGCTGGCCTATCAGG; ZFP-4R: CTGCGGATCCTTTCAATGCG;

ZFP-5F: TATGGGGGAGGACTGGTCAG; ZFP-5R: ACTAGCAGGCCTGACAACAC.

The following cycling conditions were used: 55°C for 30 min; 94°C 2 min; 35 cycles of the following: 94°C 15 sec, 56°C 30 sec, and 68°C 3.5 min; 68°C 10 min. Viral cDNA amplicons were size selected by agarose gel electrophoresis and then purified using the Qiagen MinElute Gel extraction kit. Purified PCR products were pooled and then ~1ng of DNA was fragmented using the Nextera XT kit (Illumina), tagged with Illumina-compatible primers, and then purified with Agencourt Ampure XP beads. Purified libraries were then sequenced with 2 x 300bp kits on an Illumina MiSeq.

Sequences were analyzed using a workflow (https://bitbucket.org/dhoconno/zequencer) implemented in Geneious Pro (Biomatters, Ltd.). Briefly, duplicate reads were initially removed with the BBMap v35.82 dedupe tool (https://sourceforge.net/projects/bbmap/) using a kmer seed length of 31 bp and allowing zero edits or substitutions. Non-duplicate reads were quality trimmed with BBDuk v35.82 to remove Truseq, Nextera, and PhiX adapters from both the left and right ends of sequences. Regions with low quality sequences (average quality < 20) were

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355 removed from both ends of remaining sequence reads, and only reads longer than than 150 bp
356 are retained. BBMap v35.82 was used to map these reads to the ZIKV strain H/PF/2013
357 Genbank reference sequence (KJ776791) in local alignment mode using the normal sensitivity
358 preset. The Geneious Pro 9.1.2 variant caller is used to detect and annotate variants. Variants
359 are called that fit the following conditions: present in >=5% of the mapped reads, have a
360 minimum p-value of 10e-60, and a minimum strand bias of 10e-5. Areas with low coverage
361 (<20x) are annotated.
362
363 *Calculating nucleotide diversity*
364 Synonymous (πS) and Nonsynonymous (πN) nucleotide diversity were calculated for each
365 coding region using Popoolation version 1.2.2[29]reference-based mappings were exported from
366 Geneious as bam files, which were converted to sorted pileup files using SAMtools[30]. πN and
367 πS estimates were calculated for each coding region in the genome using the Syn-
368 nonsyn-at-position.pl script. πS > πN suggests that purifying selection is acting to remove new
369 variation, πN > πS suggests that diversifying selection is favoring new mutations, and πN = πS
370 indicates no strong selection. To reduce the effects of uneven coverage across sites, pileup files
371 were subsampled such that every position was covered by exactly 1000x using the subsample-
372 pileup.pl script in Popoolation. Regions in which coverage was <1000x were excluded from the
373 analysis. πN and πS estimates were then calculated for each coding region in the genome
374 using the Syn-nonsyn-at-position.pl script. Corrections were disabled.
375
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Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

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Author contributions D.H.O., T.C.F., J.E.O., M.T.A., E.M., T.G.G. and D.M.D. designed the experiments. D.H.O., D.M.D., M.T.A., E. M., T.C.F., and L.H.M. drafted the manuscript. M.T.A., and J.E.O. provided and prepared viral stocks and performed plaque assays. A.M.W., G.L-B., and T.C.F. developed and performed viral load assays. K.L.W. and E.G.R. performed immunophenotyping assays. M.S.M., M.E.B., M.N.R., C.M.N., and D.M.D. coordinated and processed macaque samples for distribution. D.D.G., S.L.O., and D.M.D. designed and

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459 performed the sequencing experiments. L.H.M. and T.C.F. performed nucleotide diversity
460 calculations. J.P., N.S-D., H.A.S., S.C., coordinated the macaque infections, sampling, and
461 performed blood chemistries and CBC analysis. J.A.E., M.A.M., and S.R.P. performed the
462 plasmablast experiments.
463
464 **Author information:** Sequences are pending deposition into the Sequence Read Archive at
465 NCBI. The bioSample accession number is SAMN04589872. All data from these studies are
466 available at zika.labkey.com. Reprints and permissions information is available at
467 www.nature.com/reprints. The authors declare no competing interests. Correspondence and
468 requests for materials should be addressed to dhoconno@wisc.edu.
469
470 **Figure Legends**
471 **Figure 1. Study design and ZIKV viral load from rhesus macaque fluids.** a. ZIKV, strain
472 H/FP2013 from the European virus archive (Genbank: KJ776791) was inoculated onto C6/36
473 mosquito cells to generate the stock Zika virus/H.sapiens-tc/FRA/2013/FrenchPolynesia-
474 01_v1c1. Doses administered subcutaneously per animal are shown. b. Viral loads were
475 measured in plasma, urine, saliva and cerebral spinal fluid for each animal through 28 dpi. CSF
476 was collected on days 4 and 14. All animals were negative on day 14.
477
478 **Figure 2. Genetic diversity of the challenge stock and virus isolated from animals.** a. The
479 ZIKV challenge stock was deep sequenced from all three animals. Nucleotide sites where at
480 least 5% of sequences obtained from the challenge stock are different from the Genbank
481 sequence are shown. b. The plasma virus of each animal from 4 dpi was deep sequenced.
482 Non-synonymous variants present in the challenge stock relative to the Genbank consensus
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Non-synonymous variants are shown with dark colored markers, while synonymous variants are shown in a lighter shade. Changes found in both the stock and the animals are connected with a line. c. Deep sequence data from all 3 animals on day 4 post-infection were used to calculate measures of synonymous (πS) and nonsynonymous (πN) nucleotide diversity for each coding region in the genome. The πN/πS ratio for each gene is shown with color corresponding to animal. The capsid, propeptide and membrane coding regions in animal 393422 were excluded from the analysis due to coverage <1000x.

Figure 3. Immune cell expansion following ZIKV infection. Expansion of Ki-67+ (activated) CD4+, CD8+ and NK cells were measured daily for 11 days and then every 3 days for each animal. Total number of activated cells per microliter of blood are plotted for each animal for each cell type over time. The inset of each graph shows the total number of plasmablast cells found in PBMCs collected at 3, 7, 11 and 14 dpi for each animal. Viral load is graphed over the columns as a reference.

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prepared by inoculation onto a confluent monolayer of C6/36 mosquito cells to generate the stock Zika virus/R.macaque-tc/UGA/1947/MR766-3329. The viral loads for the first seven dpi are shown relative to the viral loads of animals receiving the same dose of Asian lineage ZIKV.

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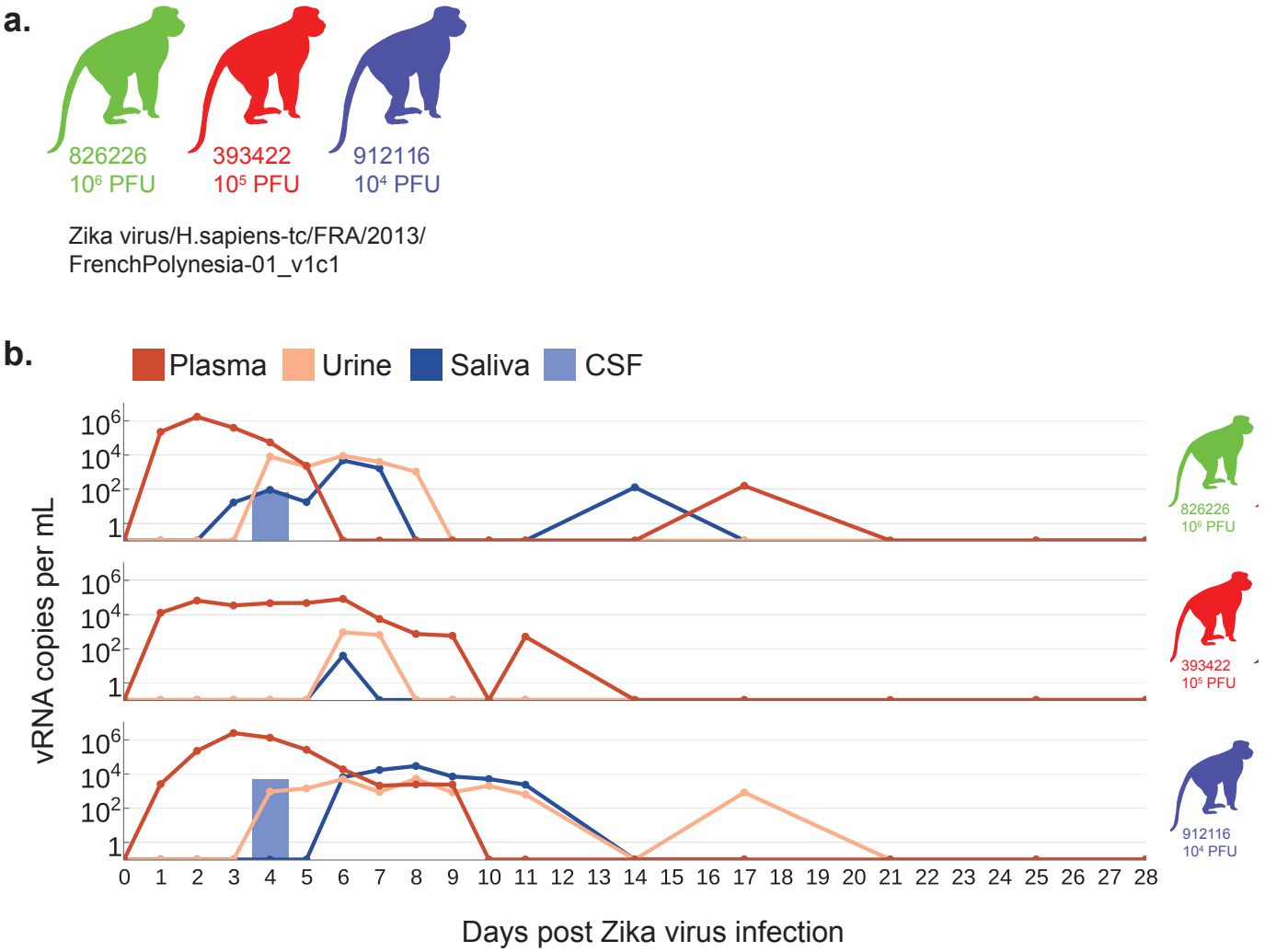

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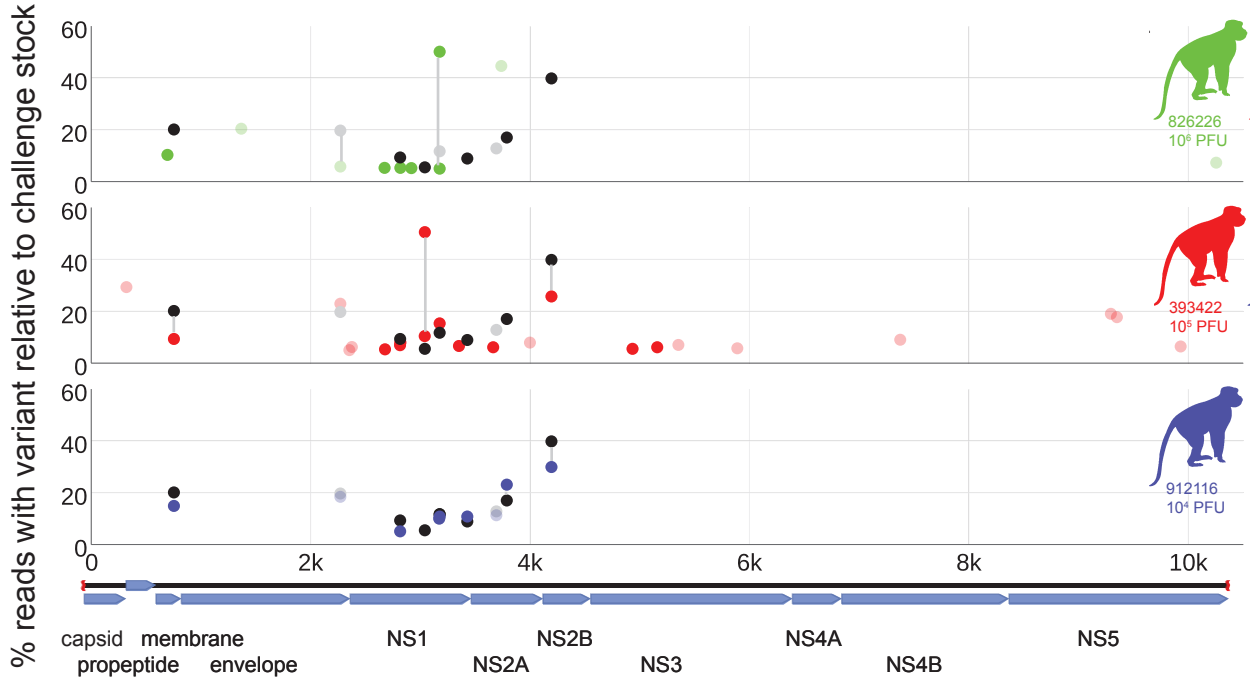
a.

Site relative to KJ776791	Change	Amino Acid Change	Codon Change	Protein Effect	Variant Frequency
801	T -> C	F->L	UUC->CUC	Substitution	17.9%
2318	C -> T		CUC->CUT	None	18.7%
2863	T -> C	L->P	CUC->CCC	Substitution	7.7%
3223	A -> T	K->I	AAA->ATA	Substitution	10.5%
3475	T -> C	M->T	AUG->ACG	Substitution	7.9%
3740	C -> T		GUC->GUT	None	11.1%
3835	C -> T	A->V	GCG->GTG	Substitution	15.8%
4242	G -> C	E->Q	GAG->CAG	Substitution	40.9%

b.



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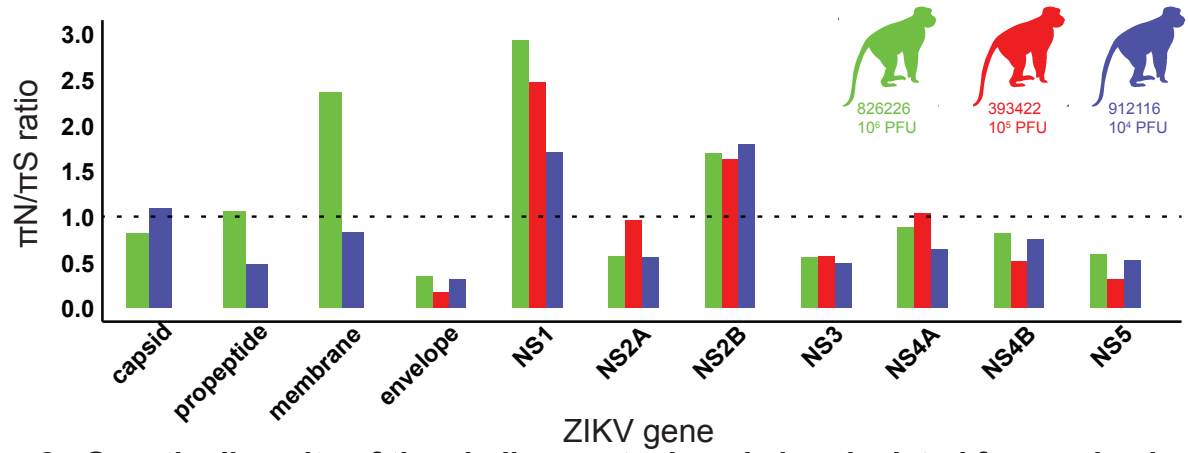

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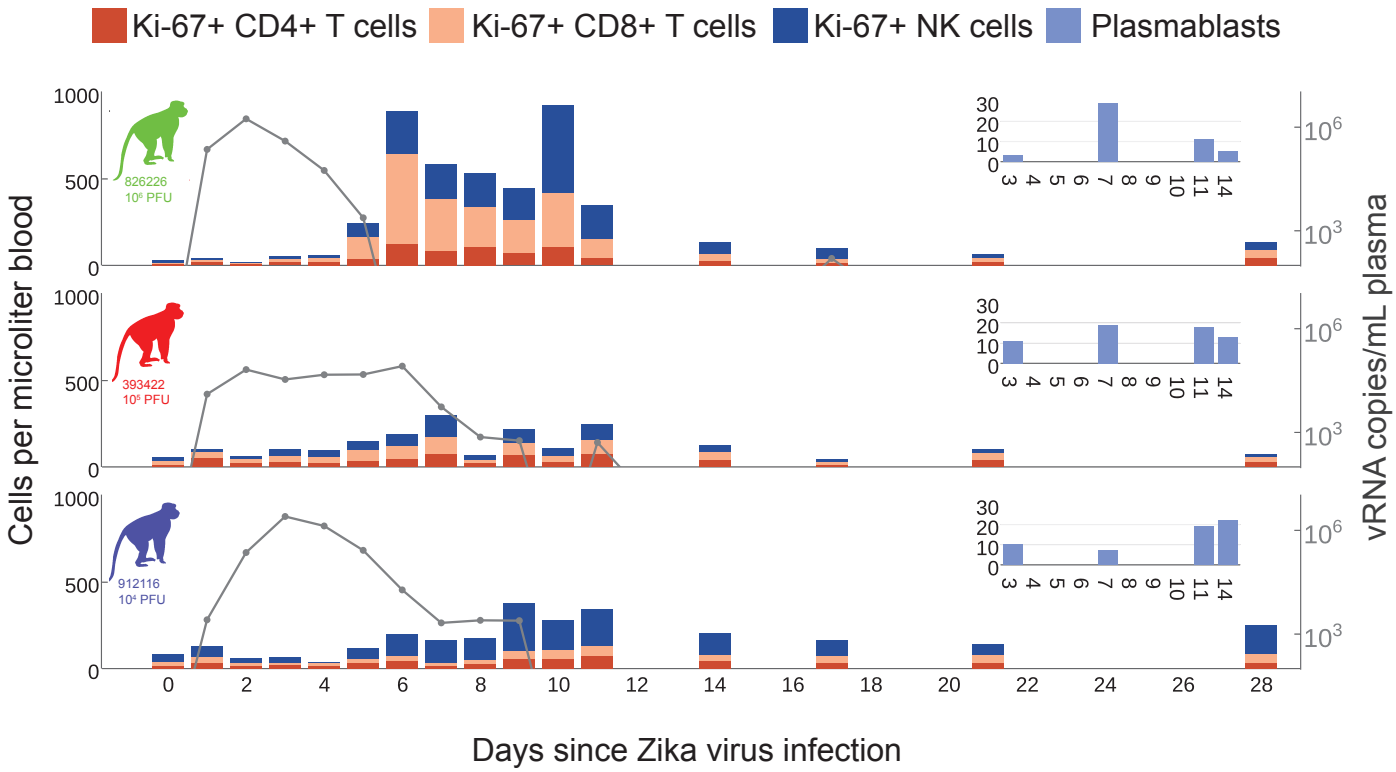

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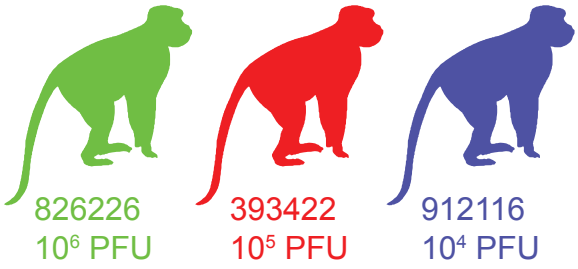

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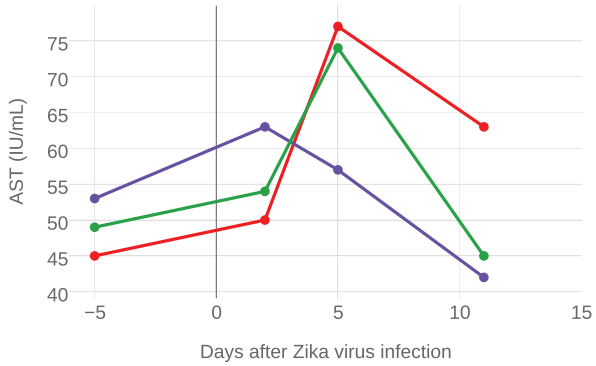

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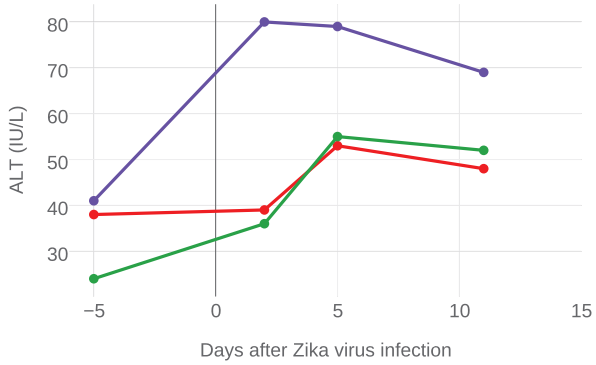

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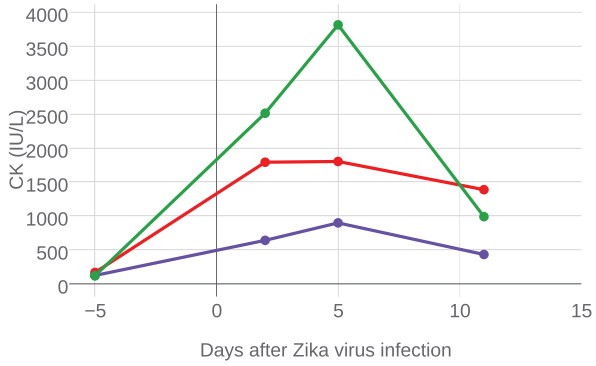

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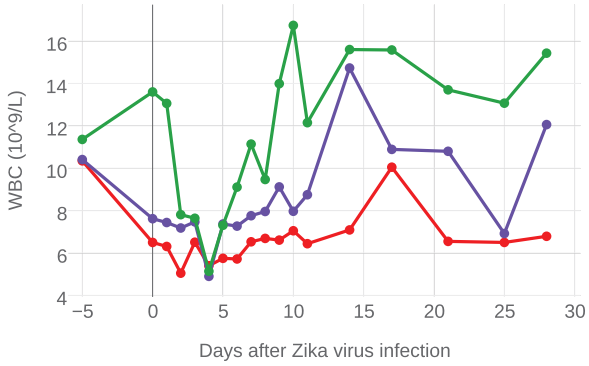

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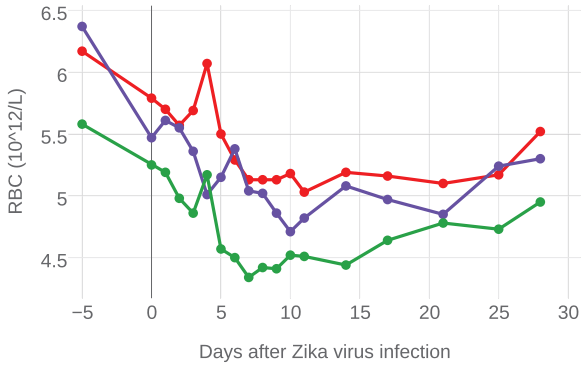

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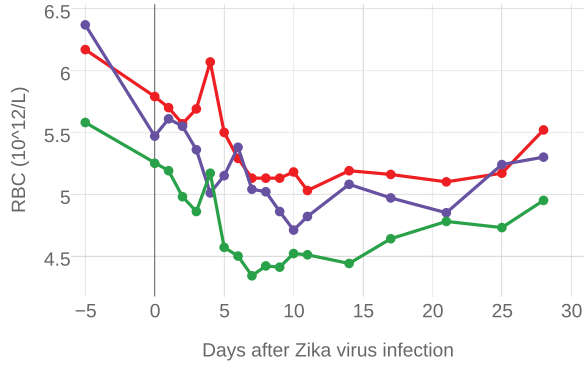

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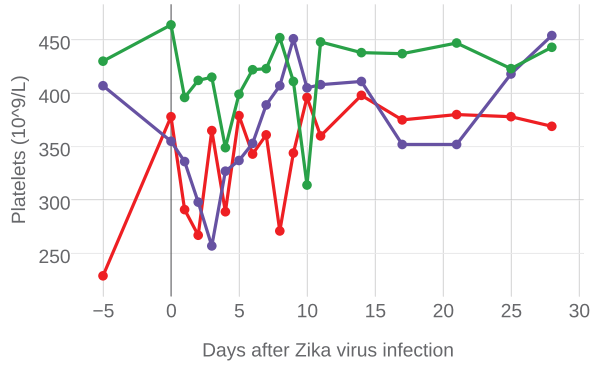

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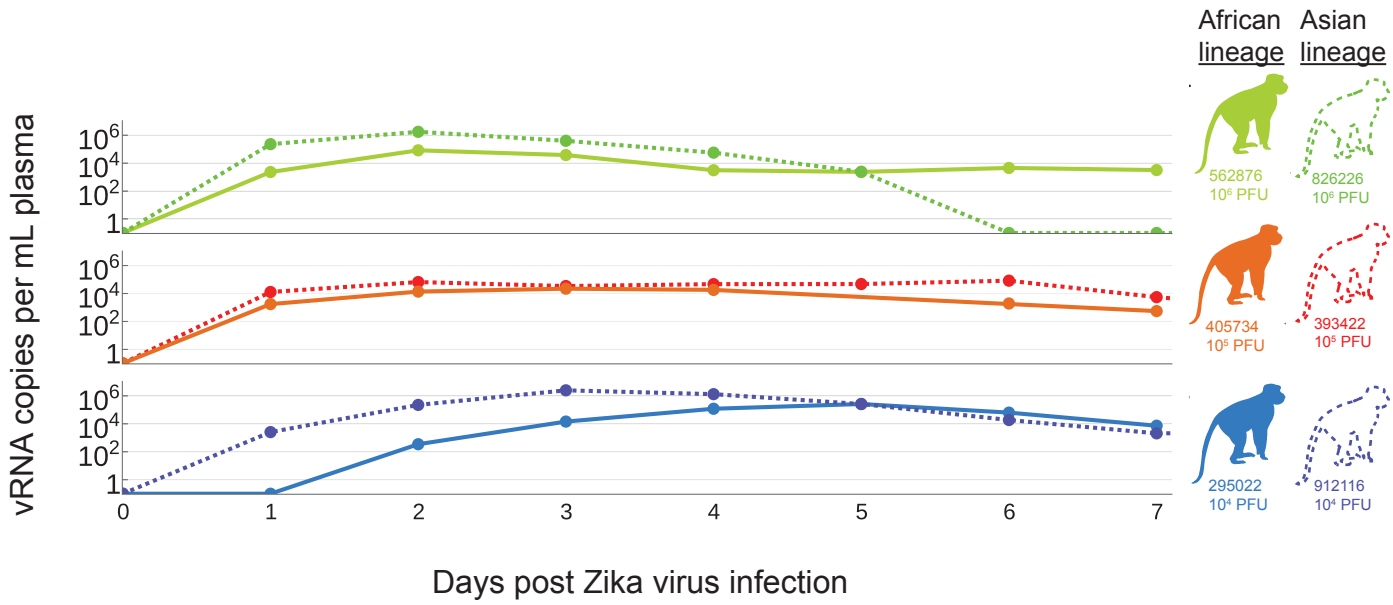

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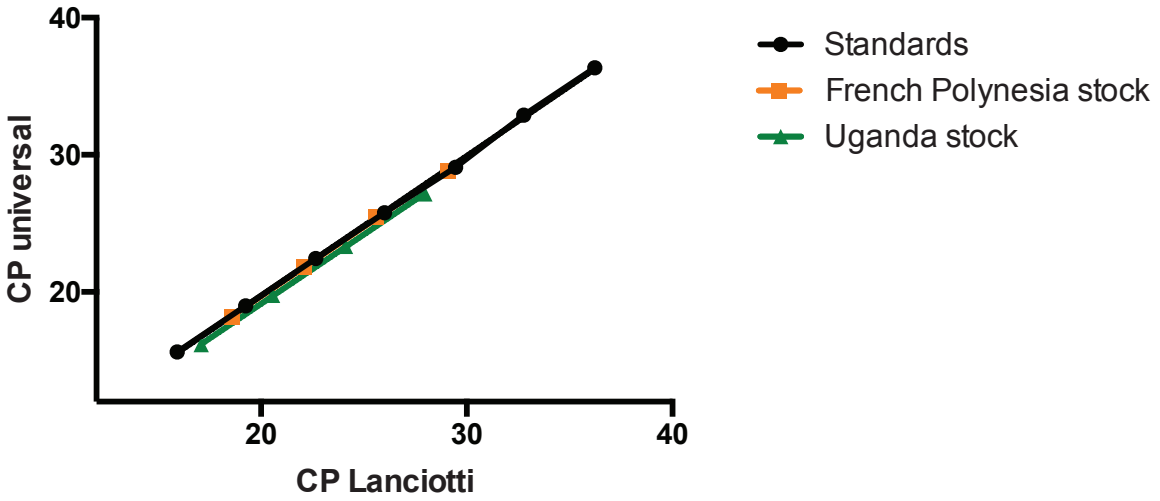

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