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Haiyan Zhao
Lily Xu
Robin Bombardi
Rachel Nargi
Zengqin Deng

See next page for additional authors

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Authors
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ARTICLE

Mechanism of differential Zika and dengue virus neutralization by a public antibody lineage targeting the DIII lateral ridge

Haiyan Zhao1, Lily Xu1, Robin Bombardi2, Rachel Nargi2, Zengjin Deng3, John M. Errico3, Christopher A. Nelson4, Kimberly A. Dowd4, Theodore C. Pierson4, James E. Crowe Jr.2,8,9, Michael S. Diamond1,5,6, and Daved H. Fremont1,5,7

We previously generated a panel of human monoclonal antibodies (mAbs) against Zika virus (ZIKV) and identified one, ZIKV-116, that shares germline usage with mAbs identified in multiple donors. Here we show that ZIKV-116 interferes with ZIKV infection at a post-cellular attachment step by blocking viral fusion with host membranes. ZIKV-116 recognizes the lateral ridge of envelope protein domain III, with one critical residue varying between the Asian and African strains responsible for differential binding affinity and neutralization potency (E393D). ZIKV-116 also binds to and cross-neutralizes some dengue virus serotype 1 (DENV1) strains, with genotype-dependent inhibition explained by variation in a domain II residue (R204K) that potentially modulates exposure of the distally located, partially cryptic epitope. The V-J reverted germline configuration of ZIKV-116 preferentially binds to and neutralizes an Asian ZIKV strain, suggesting that this epitope may optimally induce related B cell clonotypes. Overall, these studies provide a structural and molecular mechanism for a cross-reactive mAb that uniquely neutralizes ZIKV and DENV1.

Introduction

Zika virus (ZIKV) typically causes a self-limiting febrile illness, with most infected individuals exhibiting minimal or no symptoms (Duffy et al., 2009). However, ZIKV infection can result in severe neurological disease (Mlakar et al., 2016), including neurodevelopmental defects in infants after congenital infection (Moore et al., 2017; de Paula Freitas et al., 2016). Dengue virus (DENV) is genetically related to ZIKV, infects nearly 400 million people annually, and causes variable clinical disease ranging from a mild to severe febrile illness and life-threatening dengue shock syndrome (Bhatt et al., 2013). Since its introduction and spread in the Western hemisphere in 2015–2016, ZIKV has emerged as a significant global health concern.

Both ZIKV and DENV are principally transmitted by mosquitoes (Cao-Lormeau et al., 2016) and belong to the Flavivirus genus of the Flaviviridae family of single-stranded positive-sense RNA viruses, which also include West Nile (WNV), Japanese encephalitis (JEV), yellow fever, and the tick-borne encephalitis viruses (Lazar and Diamond, 2016). Flavivirus genomes encode a single polyprotein that is cleaved by viral and cellular proteases into three structural proteins (capsid protein, precursor membrane protein, and envelope [E] protein) and seven nonstructural proteins. Cryo-electron microscopy (cryo-EM) models of mature flaviviruses show 90 anti-parallel E protein dimers lying flat against the virion surface with T = 3 quasi-icosahedral symmetry (Zhang et al., 2013; Kostyuchenko et al., 2016; Qiu et al., 2018). E protein is the primary target of neutralizing antibodies and is composed of three ectodomains: domain I (DI), which links DII and DIII together; DII, which contains a fusion loop that mediates viral fusion with host endosomes; and DIII, which adopts an Ig-like fold that undergoes a substantial repositioning during viral fusion (Rey et al., 1995; Dai et al., 2016; Modis et al., 2004). Antibodies against flaviviruses map to epitopes in all three domains, and those against DIII are among the most potent at neutralizing infection (Nybakkken et al., 2005; Robbiani et al., 2017; Zhao et al., 2016; Shrestha et al., 2010; Sukupolvi-Petty et al., 2010).

Correspondence to Daved H. Fremont: fremont@wustl.edu.

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While the affinity of antibody binding governs the proportion of epitopes occupied under steady state conditions (Robinson et al., 2015), it does not always correlate with flavivirus neutralization. Another factor that influences antibody neutralization is the valency of virion engagement, where potent neutralization can be obtained for a bivalent binding antibody even in the setting of relatively weak monovalent affinity (Edeling et al., 2014). A third important factor is epitope accessibility, which is influenced by virion maturation as well as the capacity for dynamic motion and affects the stochiometry of antibody binding and efficiency of neutralization (Pierson et al., 2007; Pierson and Diamond, 2012).

Germline selection and affinity maturation of broadly neutralizing mAbs have been studied extensively for HIV and influenza virus and have allowed for the development of novel vaccine strategies (Pappas et al., 2014; Liao et al., 2013; Duan et al., 2018). Germline precursors generally show weak or undetectable affinity for target immunogens; thus, vaccine antigens may need to be engineered to induce neutralizing antibodies. For flaviviruses, most cross-reactive mAbs against the E protein target the highly engineered to induce neutralizing antibodies. For flaviviruses, most cross-reactive mAbs against the E protein target the highly conserved fusion loop in DIII. The accessibility of the fusion loop is dependent on the maturation state of the virus, with limited exposure on mature virions, and most fusion loop–directed mAbs exhibit weak neutralization potency (Zhao et al., 2016; Cherrier et al., 2009; Rey et al., 2018). Another group of cross-reactive mAbs has also been identified from DENV-infected donors that bind a quaternary E-dimer epitope and can neutralize both DENV and ZIKV infection efficiently (Dejmirattisai et al., 2015; Fernandez et al., 2017). These E-dimer epitope mAbs have shown significant potency against ZIKV both prophylactically and therapeutically in murine models of infection (Fernandez et al., 2017).

We and others have reported mAbs from multiple human donors that all use the VH3–23 heavy chain and Vk1–5 light chain and have the unique capacity to neutralize both ZIKV and DENV with varying potencies (Sapparapu et al., 2016; Robbiani et al., 2017; Magnani et al., 2017; Niu et al., 2019). Several of these public lineage mAbs have demonstrated protection in animal models of ZIKV challenge. Herein, we combine virological and biophysical assays to define the mechanistic basis for the neutralizing activity of ZIKV-116, a mAb derived from a donor infected by ZIKV in 2015 in Brazil. Functional studies in cell culture show that ZIKV-116 primarily inhibits infection at a post-cellular attachment step by blocking viral fusion with host membranes. X-ray crystallographic studies reveal that ZIKV-116 recognizes a DIII lateral ridge (LR) epitope, with one residue distinguishing binding and consequent neutralization of Asian versus African ZIKV strains. ZIKV-116 also potently neutralizes infection of DENV1, with genotype-specific neutralization best explained by differential epitope exposure. Lastly, we constructed the inferred V-J germline sequence of ZIKV-116 and demonstrated that it binds and neutralizes diverse ZIKV and DENV1 strains, albeit with lower potency than the affinity-matured clone.

**Results**

**ZIKV-116 primarily inhibits infection at a post-attachment step**

To investigate how ZIKV-116 inhibited virus infection, we performed pre- and post-attachment neutralization assays with a French Polynesian ZIKV strain (H/PF/2013). As anticipated, prebinding with ZIKV-116 efficiently limited ZIKV infection (Fig. 1A). ZIKV-116 similarly inhibited ZIKV infection after virus was bound to the target cell surface (half maximal inhibition concentration [IC50] of 52 and 105 ng/ml for pre- and post-attachment neutralization, respectively). To corroborate these studies, we evaluated whether ZIKV-116 could inhibit cellular attachment of ZIKV. Different concentrations of ZIKV-116 or an isotype control mAb were preincubated with ZIKV and then added to Vero cells. After extensive washing, cell-associated viral RNA was measured by quantitative real-time PCR (qRT-PCR). ZIKV-116 did not block virus attachment relative to isotype control near the IC50 of 0.1 µg/ml (Fig. 1B). However, a modest inhibition of attachment (~60%) was observed at concentrations ~10- to 100-fold higher than the IC50 value, suggesting that attachment blockade is not the dominant inhibitory mechanism of ZIKV-116.

**ZIKV-116 can block viral fusion with host membranes**

We next examined whether ZIKV-116 could block viral fusion. Normally, flaviviruses enter cells via receptor-mediated internalization, and pH-triggered fusion occurs with endosomal membranes (Modis et al., 2004). However, under low-pH conditions, cell-associated virus can be induced to fuse at the plasma membrane, a process termed “fusion from without” (Edwards and Brown, 1986; Liao and Kielland, 2005; Thompson et al., 2009; Fernandez et al., 2018). ZIKV was preadsorbed to Vero cells on ice and subsequently treated with ZIKV-116 or control ZV-2, a ZIKV-specific nonneutralizing mAb (Zhao et al., 2016). Virus-plasma membrane fusion was triggered by a brief exposure to low pH, and infection was monitored at 24 h after initial treatment. In all steps, 25 mM NH4Cl was added to prevent ZIKV entry and infection via endosomal fusion (Helenius et al., 1982). As expected, under neutral pH conditions, ZIKV infection was rarely detected. In comparison, exposure of cell-adsorbed ZIKV to pH 5.8 resulted in a 10-fold increase in infection. ZIKV-116 completely blocked this pH-triggered infection, whereas mAb ZV-2 did not (Fig. 1, C and D).

To block fusion in cells, ZIKV-116 must retain binding at the acidic pH of the endosome. To evaluate this question, we performed biolayer interferometry (BLI) with ZIKV-116 and its ligand ZIKV DIII at pH values corresponding to the extracellular and endosomal milieu. ZIKV-116 was immobilized and dipped into solutions of recombinant ZIKV DIII (H/PF/2013) ranging from 3.125 nM to 100 nM concentration. At pH 7.4, DIII bound to ZIKV-116 with a kinetically derived binding affinity (Kd) of 1.15 nM and t1/2 of 25.70 min. The complex was only slightly less stable at pH 5.5, with a Kd of 2.24 nM and t1/2 of 16.06 min (Fig. 1, E and F). These results suggest that ZIKV-116 engagement of virions should endure within the acidic environment of endosomes such that viral fusion can be blocked.

**X-ray crystal structure of ZIKV-116 in complex with ZIKV DIII**

To better understand ZIKV-116-mediated neutralization, we determined the x-ray crystal structure of ZIKV-116 antigen-binding fragments (Fab) in complex with ZIKV DIII (H/PF/2013) at 2.3 Å resolution (Fig. 2 and Table S1). ZIKV-116 binds to
the LR of DIII by engaging the N-terminal region as well as the BC-, CC-, FG-loops. The total buried surface area at the antibody-antigen interface is 1,551 Å², with a shape complementarity factor of 0.71. The heavy chain accounts for 67% of the interface, with a combined buried surface of 1,042 Å² (542 Å² on DIII and 500 Å² on the heavy chain), and the light chain contributes the remaining buried surface area (264 Å² on DIII and 245 Å² on the light chain). The interaction is mediated by contacts between 12 DIII residues and residues from 5 ZIKV-116 complementarity-determining regions (CDRs: CDR-H1, CDR-H2, CDR-H3, CDR-L1, and CDR-L3) plus 2VN framework residues. Two of the DIII residues (T£309 and K£394) are contacted by both heavy and light chains. Binding of ZIKV-116 to DIII is mediated by 11 direct hydrogen bonds and 9 water-mediated networks in addition to van der Waals contacts. The N-terminal and FG-loop regions contain the majority of the interactions (79 of 99 contacts) with ZIKV-116. The FG-loop has the most extensive contact network, forming five direct hydrogen bonds and four water-mediated hydrogen bonds with ZIKV-116. The N-terminal region of DIII contributes 35 contacts and forms 5 additional direct hydrogen bonds with residues from the CDR-L3 and CDR-H2 of ZIKV-116 (Tables S2–S6). We note that there are two complexes in the crystallographic asymmetric unit that adopt overall similar conformations, with a root mean square
deviation of 1.19 Å over 332 Cα atoms of DIII and variable domains of ZIKV-116 (Fig. S1, A and B). However, presumably due to crystal packing, we observed a different conformation of the DIII CC9-loop in one complex where it is not engaged by ZIKV-116 CDR loops but rather by an adjacent Fab (Fig. S1, C and D).

Variation at residue 393 modulates the binding kinetics of ZIKV-116

We performed infection assays and found that ZIKV-116 neutralized four ZIKV Asian strains (H/PF/2013, Brazil Fortaleza, Cambodia_FSS13025, and Puerto Rico_PRVABC58) more effectively than two African strains (MR-766 and Dakar 41519; Table S7), consistent with our previous observations (Sapparapu et al., 2016). Sequence analysis of the ZIKV-116 structural epitope among the seven representative ZIKV strains showed that 11 out of 12 contact residues are invariant. However, African strains (MR-766 and Dakar 41519) have an Asp at residue 393, while most Asian strains have a Glu. Interestingly, one Asian strain (Malaysia_P6740), which has an Asp393 like African strains, was less effectively neutralized by ZIKV-116 compared with other Asian strains (Table S7).

In our structure, Glu393 is directly contacted by ZIKV-116 CDR-H3 and participates in an extensive hydrogen-bonding network, forming three direct hydrogen bonds and two water-mediated indirect hydrogen bonds with ZIKV-116 (Fig. 3 A and Tables S4–S6). To explore whether the residue 393 impacts ZIKV-116 activity, we used BLI to compare the binding of ZIKV-116 to different purified DIII proteins, including WT H/PF/2013 DIII, E393D H/PF/2013 DIII, and WT MR-766 DIII. ZIKV-116 bound MR-766 DIII weakly (100-fold higher KD [121 nM] and t1/2 of 0.25 min) compared with H/PF/2013 DIII (Fig. 3, B and C; and Fig. 1 E). Correspondingly, the E393D substitution of H/PF/2013 DIII (to the residue in MR-766) significantly decreased the binding affinity and half-life, with a KD of 62 nM and t1/2 of 0.52 min. Thus, it appears that a single methyl group of residue 393 in ZIKV strains can determine ZIKV-116 binding strength, most likely because the shorter side chain is incapable of optimally forming a hydrogen-bonding network with residues in CDR-H3, specifically ArgH96 and GluH100C, which engage the DIII Glu393 carboxylate and main chain, respectively (Fig. 3 A).

The amount of ZIKV-116 required to neutralize Malaysia_P6740 strain is higher than that required to inhibit Asian strain H/PF/2013, and lower than the amount required to neutralize African strains (Table S7). Consistently, the DIIDP(E393D) mutant, which is identical to the naturally occurring DIII of Malaysia_P6740 strain, bound ZIKV-116 with an ~50-fold lower affinity than DIIIWT and ~2-fold higher affinity than WT DIIIMR. While the structurally defined epitope is strictly conserved among analyzed ZIKV strains with the exception of residue 393, the African MR-766 strain varies at three additional positions in DIII regions that could distally affect the epitope and thereby mAb binding (Fig. S2).

Variation at residue 393 modulates the neutralizing activity of ZIKV-116 against different ZIKV strains

To assess the role of residue 393 in differential ZIKV-116 neutralization, we generated ZIKV reporter virus particles (RVPs) with H/PF/2013 and MR-766 structural proteins and incorporated...
the reciprocal changes at residue 393. Exchanging the amino acid at residue 393 dramatically reversed the ZIKV-116 neutralization phenotype between the H/PF/2013 and MR-766 strains. The IC50 values of WT H/PF/2013 and E393D variant increased from 69 to 4,855 ng/ml, making the neutralization of E393D H/PF/2013 RVPs similar to the neutralization of WT MR-766. Reciprocally, D393E MR-766 RVPs were neutralized by ZIKV-116 similarly to WT H/PF/2013 RVPs (approximately twofold difference in IC50; Fig. 3 D). In comparison, the presence of D393 or E393 had little effect on mAb ZV-67, which binds an overlapping DIII-LR epitope and neutralizes H/PF/2013 and MR-766 strains equivalently (Fig. 3 E; Zhao et al., 2016). Taken together with our binding studies, these observations strongly suggest that differential ZIKV neutralization by ZIKV-116 is primarily controlled by residue 393 variation, and that mAb-binding affinity correlates well with neutralization potency against distinct ZIKV strains.

Differential neutralization of DENV1 genotypes

Sequence analysis of the ZIKV-116 mAb showed that it used the VH3-23 heavy chain and VK1-5 light chain genes, and VH3-23/VK1-5 mAbs against ZIKV have been reported to cross-react with DENV1 (Sapparapu et al., 2016; Robbiani et al., 2017; Magnani et al., 2017; Niu et al., 2019). To investigate cross-reactivity, we tested the specificity of ZIKV-116 against a panel of recombinant proteins from different flaviviruses by ELISA (Fig. 4, A and B). In this qualitative assay, ZIKV-116 recognized H/PF/2013 DIII, E, and E-FL (a fusion loop mutant) equally well as MR-766 DIII. We also found that ZIKV-116 can recognize DIII from DENV1, but not DIII from other DENV serotypes (DENV2, DENV3, or DENV4) or E or DIII proteins from other flaviviruses (WNV, JEV, Spondweni, or Powassan viruses). Based on these binding properties, we assessed whether ZIKV-116 could inhibit infection of different DENV1 strains. We found that ZIKV-116 neutralized a DENV1 genotype 2 strain (16007) more effectively than a genotype 4 strain (West Pac-74), with IC50 values of 11 and 73 ng/ml, respectively (Fig. 4 C). No neutralization activity was observed to DENV2. Sequence comparison revealed only two amino acid differences in DIII between 16007 and West Pac-74, both located in the CC9-loop (T339S and A345V). Residue 345 but not 339 is located within the predicted ZIKV-116 footprint on DENV1 DIII (Fig. 5 A and Fig. S2).

A residue distal to the ZIKV-116 epitope modulates DENV1 strain neutralization sensitivity

Since ZIKV-116 less efficiently neutralized DENV1 West Pac-74 than 16007, we evaluated the binding affinity of ZIKV-116 to DIII from these two viruses. BLI experiments revealed that ZIKV-116
Reciprocally, exchange of K204R on the West Pac-74 background led to neutralization curves to that seen with WT 16007 RVPs (IC50 of the affinity-matured ZIKV-116 is encoded by VH3 binding and neutralizing ZIKV and DENV1 strains. A reverted, unmutated ancestor form of ZIKV-116 is capable of variantaD E N V 1 R V P s (Fig. 5 F).

To investigate whether this same residue 204 variation also accounts for the different neutralization activity of ZIKV-WT 116, we performed neutralization assays using DENV1 16007 and West Pac-74 RVPs bearing reciprocal changes at E residue 204. Exchange of K204R on the West Pac-74 background shifted the neutralization curves to that seen with WT 16007 RVPs (IC50 of 29 ng/ml for K204R West Pac-74 versus 16 ng/ml for WT 16007). Reciprocally, exchange of R204K on the 16007 background led to an 11-fold reduced ZIKV-116 neutralization potency (Fig. 5 D). In contrast, a control mAb, WNV-E60, which binds the E-FL epitope in DII, showed similar neutralization potency with all WT and variant DENV1 RVPs (Fig. 5 F).

A reverted, unmutated ancestor form of ZIKV-116 is capable of binding and neutralizing ZIKV and DENV1 strains

The affinity-matured ZIKV-116 is encoded by VH3-23 and VK1-5 germline genes, with 18 aa substitutions in the V1 domain and 10 aa substitutions in the V2 domain. Only 2 of the 28 residues mutated from V and J gene segments are part of the ZIKV-116 paratope, S131N in CDR-H1 and T149K adjacent to CDR-H2, and both residues make only modest contacts with the DIII-LR epitope (Tables S2–S6). The remaining 26 residue substitutions are located mainly in the V1 framework regions, CDR-H2 and CDR-L1/2 (Fig. 6, A and B). To investigate the properties of the ZIKV-116 clonal lineage, we reverted ZIKV-116 to the inferred germline sequence (reverted, unmutated ancestor mAb, ZIKV-116-V.J.Rev) and produced it using the same IgG1 expression vectors as the affinity-matured mAb. Both the V and J gene segments were reverted to the respective templated germline sequences. The reverted antibody retained the original CDR-H3 of the mature WT ZIKV-116 antibody, which possesses one somatically mutated residue when compared with the inferred germline D gene segment IGHD3-10’01.

We next evaluated the binding of ZIKV and DENV1 DIII proteins to immobilized ZIKV-116-V.J.Rev using BLI (Fig. 6 C). Surprisingly, ZIKV-116-V.J.Rev showed only a moderately decreased binding to ZIKV H/PF/2013 DIII (Kd of 48.9 nM and t1/2 of 1.18 min). On the other hand, binding to DIII from ZIKV MR-766 and the two DENV1 strains was markedly weaker (Kd of ~10 μM for DIII MR-766 and Kd of ~7 μM for DIII 16007 and DIII West Pac-74). The decreased binding to DIII of all four viruses was due primarily to increased dissociation rates (Fig. 6 C and Table S8).

Neutralization assays revealed that ZIKV-116-V.J.Rev has better than West Pac-74 (A) ZIKV-116 or control mAbs (WNV-E111 [flavivirus cross-reactive] and DENV4-E88 [DENV4 type-specific]) were tested for binding to the indicated flavivirus proteins (ZIKV E, ZIKV E-FL [mutant strain of H/PF/2013], ZIKV DIII [strains of H/PF/2013 and MR-766], WNV E and DII, SPOV E [Spondweni virus], and DENV1–4 DII) as indicated in ELISA. (B) ZIKV-116 or control mAbs (JEV 306 [JEV DIII-specific] and POWV-61 [Powassan DIII-specific]) were tested for binding to the indicated DII of ZIKV H/PF/2013, JEV, and POWV. The results are representative of two independent experiments performed in triplicate. (C) Neutralization curves of DENV2 (D2S20) and DENV1 (16007 and West Pac-74 [WP]) by ZIKV-116 and control mAb DENV1-E111. The indicated virus was incubated with serial-diluted mAbs for 1 h at 37°C followed by addition of the mixture to Vero cells. Then, FRNT assays were performed, and the percentage of infection was calculated by comparing to no mAb–treated wells. Data were pooled from two or more independent experiments performed in triplicate or quadruplicate.
with 30- to 50-fold lower activity compared with the somatically mutated mAb. However, ZIKV-116-V.J.Rev neutralization of ZIKV MR-766 was only modest, with an apparent 100-fold reduced activity that failed to block infection of 50% of the African virus even at the highest mAb concentration tested. We note that the binding and neutralization activity to all ZIKV and DENV strains increased during somatic evolution of the ZIKV-116 lineage (Fig. 6, E and F). Further, the increased neutralization potency of the affinity-matured mAb relative to the germline configuration is highly correlated for each virus with the monotypic binding affinity we measured for DIII (Fig. 6 G).

Figure 5. Variation at residue 204 in DII is responsible for the neutralization difference between two DENV1 strains. (A) Cartoon of DENV1 E protein (DI, red; DII, yellow; DIII, blue). ZIKV-116 epitope is highlighted in magenta. Amino acid differences (339 and 345) between 16007 and West Pac-74 (WP) DIII are shown in cyan. Residue 204, which is responsible for the difference in neutralization between two DENV1 strains, is marked as a black triangle in DII. (B and C) Comparison of $K_S$ and $t_{1/2}$ of ZIKV-116 to DIII from H/PF/2013, 16007, and West Pac-74. The results from at least two independent experiments are shown. Error bars represent SD. (D–F) Effects of residue 204 on ZIKV-116 neutralization of DENV1 strains 16007 and West Pac-74. Neutralization profiles of mAbs against WT DENV1 strains 16007 and West Pac-74 RVPs compared with mutant RVPs containing reciprocal amino acid at residue 204 (upper). IC50 values are from three independent experiments (bottom). DENV1-E111, which was reported to display differential neutralization against DENV1 genotypes, is used as a positive control mAb, and WNV-E60 serves as a negative control mAb. Statistical significance was determined by one-way ANOVA with a Tukey’s test (B and C) or two-tailed t tests (D–F). ns, not significant; *, P < 0.05; **, P < 0.01; ***, P < 0.001. FP, H/PF/2013.

The ZIKV-116 epitope is not fully exposed in the cryo-EM model of mature ZIKV

Three cryo-EM structures of mature ZIKV H/PF/2013 have been reported to date (Sirohi et al., 2016; Kostyuchenko et al., 2016; Sevvana et al., 2018), and we used the highest-resolution model to evaluate how ZIKV-116 might interact with the virion (Protein Data Bank [PDB] 6C08). Analysis of the virion structure indicates that the DIII-LR epitope is predominantly solvent exposed, with the exception of some parts of the CC9-loop (Fig. 7 A). However, docking of our ZIKV-116-DIII complex to all three unique T = 3 environments on the virus revealed that steric

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hindrance imposed by adjacent E subunits would cause clashes with the ZIKV-116 VH domain (Fig. 7, B–D). Thus, the ZIKV-116 epitope can be considered to be partially cryptic in the context of the mature, icosahedrally averaged structure. In fact, the predicted virion binding orientation of ZIKV-116 places the Fab lying nearly parallel to the virus membrane, similar to a typical CC loop engaging mAb (i.e., DENV1-E111; Austin et al., 2012) and in contrast to some other DIII-LR mAbs, which project away.
from the virion surface (i.e., DENV1-E106; Edeling et al., 2014; Fig. 7, E–G). We speculate that as claimed for other viruses (e.g., WNV, DENV, norovirus, HIV, and hepatitis C virus; Dowd et al., 2011; Sukupolvi-Petty et al., 2013; Lindesmith et al., 2014; Keck et al., 2016; Mengistu et al., 2015), dynamic motions or “breathing” of virions might enable cryptic epitopes to become transiently exposed, allowing for antibodies like ZIKV-116 to bind and neutralize infection. Virion breathing and its effects on epitope accessibility have been well studied with other flaviviruses, and time-dependent increases in mAb neutralization have been observed for mAbs that recognize cryptic epitopes (Austin et al., 2012; Dowd et al., 2011; Li et al., 2018; Lok et al., 2008). To experimentally address this issue, we evaluated the time and temperature dependence of ZIKV-116 neutralization (Fig. S3). We observed mildly enhanced ZIKV neutralization by ZIKV-116 following prolonged incubation times, although these changes do not appear particularly temperature dependent.

Discussion

Previous studies have shown that antibody-mediated immune responses against flaviviruses are protective (Nybakken et al., 2005; Sapparapu et al., 2016; Shrestha et al., 2010; Fernandez et al., 2018). The identification of the mechanisms of neutralization and key epitopes of potently neutralizing antibodies has implications for antibody-based therapies and vaccine development. Here, using structural and functional approaches, we characterized a DIII-LR mAb that is commonly elicited in humans against ZIKV and found to cross-react with DENV1, and determined the molecular basis of its inhibitory activities.
Mechanism-of-action experiments showed that ZIKV-116 could efficiently block infection even after the virus has attached to target cells. Indeed, ZIKV-116 could block viral fusion with cell membranes using an in vitro assay. We and others have previously described DIII-LR mAbs against WNV (Thompson et al., 2009), JEV (Fernandez et al., 2018), and ZIKV (Wang et al., 2019) that are capable of blocking viral membrane fusion, and similar mechanisms of action have been reported for flavivirus mAbs against other epitopes (Gollins and Porterfield, 1986; Füzik et al., 2018; Zhang et al., 2016; Vogt et al., 2009; Zhang et al., 2015; Cockburn et al., 2012). Consistent with its ability to block fusion, ZIKV-116 retained high affinity binding to DIII at an acidified endosomal pH. Fusion inhibition may be a particularly important neutralization mechanism for flavivirus mAbs since viruses can apparently enter target cells through multiple receptors and entry pathways (Pokidysheva et al., 2006; Hamel et al., 2015; Sirohi and Kuhn, 2017; Perera-Lecoin et al., 2013; Cruz-Oliveira et al., 2015).

We found that the neutralization potency of ZIKV-116 to different ZIKV strains is highly correlated with the identity of residue 393 in the FG-loop of DIII. Using engineered reporter viruses, we found that an E393D change in the Asian ZIKV H/PF/2013 strain resulted in significantly diminished neutralization, while the reciprocal D393E substitution in the African ZIKV MR-766 strain dramatically improved neutralization. These results support the idea that E393 is critical for optimal ZIKV-116 engagement, and indeed, neutralization is highly correlated with ZIKV DIII binding affinity and half-life. We previously described E393 as a contact residue for a murine mAb, ZV-67 (Zhao et al., 2016), which also engages the DIII-LR epitope, albeit from a different orientation compared with ZIKV-116. However, ZV-67 neutralized all examined ZIKV strains equivalently, and activity is unperturbed by E393D substitutions. The likely explanation for this disparity is that E393 coordinates an elaborate hydrogen-bonding network at the ZIKV-116 paratope–epitope interface while it is a minor player in the murine mAb interface.

Our studies have revealed that ZIKV-116 can neutralize DENV1, with higher potency measured for genotype 2 strain 16007 relative to genotype 4 strain West-Pac-74. Quantitative binding experiments showed that DIII from both viruses binds ZIKV-116 equivalently. Similar differences in genotype-dependent neutralization of DENV1 were observed for the murine mAb DENV1-E111, which primarily recognizes a cryptic DIII CC’-loop epitope (Austin et al., 2012; Dowd et al., 2015). For DENV1-E111 the genotype-dependent differences in neutralization were found to be due in part to DII residue 204, which may allosterically affect the display of DIII epitopes (Fig. S5 E; Dowd et al., 2015). Here, we have found that residue 204 plays a similar role in modulating ZIKV-116 neutralization sensitivity of DENV1 strains, with R204-containing viruses better inhibited than K204 viruses. Similar to DENV1-E111, our analysis suggests that the DIII-LR epitope engaged by ZIKV-116 is cryptic. Indeed, while the epitope is predominantly solvent exposed, docking of our crystal structure onto cryo-EM models of mature viruses indicated steric clashes of the Fab that could only be resolved by envelope rearrangements. While it is tempting to speculate that DENV1 residue 204 could play a role in such virion conformational dynamics, as it is located ~60 Å away from the DIII-LR epitope of the same E protein, it is formally possible that it is instead a direct contact residue, as it is located only ~30 Å distant by icosahedral symmetry (Fig. S4). The precise role of residue 204 may be best addressed in future studies of virion-antibody complexes using cryo-EM.

Human mAbs that target DIII and cross-react between ZIKV and DENV1 have been described from at least eight donors examined by four independent groups (Robbiani et al., 2017; Sapparapu et al., 2016; Magnani et al., 2017; Niu et al., 2019). Remarkably, all of these mAbs use the same germline precursors as ZIKV-116, specifically the \( V_{\beta}3-23 \) heavy chain and \( V_{\gamma}1-5 \) light chain, arguing strongly that this pairing represents a broadly shared, public antibody lineage. Similar to ZIKV-116, many of these \( V_{\beta}3-23/V_{\gamma}1-5 \)-derived mAbs exhibit better neutralization of Asian ZIKV strains relative to African strains (Niu et al., 2019; Magnani et al., 2017). Further, protection studies in macaques with one of these mAbs, Z004, led to a viral escape mutant with an E393D substitution (Keeffe et al., 2018), a result supporting our conclusion that natural variation at residue 393 determines the differential neutralization of ZIKV strains by ZIKV-116. In recent years, germline-targeted vaccine development has been explored for HIV, with immunogens engineered for optimal engagement of broadly neutralizing mAb precursors (Jardine et al., 2016; Steichen et al., 2016; Duan et al., 2018; Burton, 2019). We found that the V-J reverted germline form of ZIKV-116 can broadly bind and neutralize ZIKV and DENV1 strains, with the highest potency against Asian ZIKV strain H/PF/2013. Thus, to induce \( V_{\beta}3-23/V_{\gamma}1-5 \)-lineage mAbs like ZIKV-116, vaccine strategies should probably include ZIKV E proteins that encode E393, although suboptimal protection against African ZIKV strains is possible.

In summary, we have defined the structural and molecular mechanism of a representative DIII-LR–specific mAb from a common germline lineage in humans. We found that the potency of ZIKV-116 against ZIKV strains is tightly correlated with a single epitope residue that modulates binding affinity, while differential DENV1 strain neutralization depends on a residue outside the epitope, which presumably modulates epitope accessibility on the virion. These studies highlight how the choice of virus strain can impact antibody responses and inform future epitope-based and germline-directed vaccine strategies against flavivirus infections.

Materials and methods
Key resources can be found in Table S9.

Viruses and cells
Vero, C6/36, and HEK 293T cells were maintained in DMEM supplemented with 7% FBS, 100 U/ml penicillin/streptomycin, 10 mM Hepes, 1 mM sodium pyruvate, and 1× nonessential amino acids. Raji cells (Raji-DCSIGNR) which express the attachment factor DC-SIGNR (dendritic cell–specific ICAM-3-grabbing nonintegrin related) were maintained in RPMI 1640 medium with the same media supplements.
**Virus and RVPs**

ZIKV strains H/PF/2013 (French Polynesia, 2013) and MR-766 were grown in Vero cells at 37°C. DENV1 16007 (genotype 2) and West Pac-74 (genotype 4) were amplified in C6/36 Aedes albopictus cells at 28°C. ZIKV and DENV1 RVPs were generated as described previously (Dowd et al., 2016, 2015). HEK 293T cells were cotransfected with a plasmid encoding the structural C-prM-E proteins and another plasmid encoding a WNV replicon with a GFP reporter gene, which can be used to monitor infection by flow cytometry. RVP-containing supernatants were harvested between 72 and 120 h after transfection and frozen at -80°C.

**Focus-forming and reduction assays**

A focus-forming assay (FFA) was used to determine virus titer on Vero cells. Vero cell monolayers were inoculated with virus or virus-mAb mixture and incubated at 37°C for 1 h. Cells then were overlaid with 1% (wt/vol) methylcellulose in MEM supplemented with 4% FBS and incubated at 37°C. 40 h (ZIKV) or 68 h (DENV) later, the cells were fixed with 4% paraformaldehyde and incubated with 500 ng/ml WNV-E60 mAb (Olliphan et al., 2006) in PBS buffer supplemented with 0.1% saponin and 0.02% Tween 20 for 2 h. Washing, cells were stained with HRP-conjugated goat anti-mouse IgG, and virus-infected foci were developed using TrueBlue peroxidase substrate (KPL) for 30 min and counted by an ImmunoSpot 5.0.37 macroanalyzer (Cellular Technologies).

Focus reduction neutralization tests (FRNT) were conducted with indicated ZIKV strains and mAbs in Vero cells in 96-well plates as described previously (Zhao et al., 2016). The infection frequency of the wells inoculated with virus in the presence of mAb was compared with that of wells inoculated with virus and medium alone. The IC50 values were determined using nonlinear regression analysis (GraphPad Prism).

**RVP titration and neutralization assay**

ZIKV and DENV1 RVP neutralization assays were performed as previously described (Dowd et al., 2016, 2015). To determine virus titer, twofold dilutions of RVPs were used to infect Raji-DCSIGNR in duplicate technical replicates at 37°C. GFP-positive infected cells were detected by flow cytometry 40 h later. In subsequent neutralization assays, RVPs were sufficiently diluted to within the linear range of the virus-infectivity dose-response curve to ensure antibody excess at informative points. To measure neutralization, ZIKV or DENV1 RVPs were mixed with serial dilutions of mAb for 1 h at 37°C, followed by infection of Raji-DCSIGNR cells in duplicate technical replicates. Infections were performed at 37°C, and GFP-positive infected cells were quantified by flow cytometry 2 d later. Results were analyzed by nonlinear regression analysis to estimate the dilution of mAb required to inhibit 50% of infection (IC50).

**ELISA**

All the recombinant E and DIII (5 µg/ml) from distinct flaviviruses were coated onto a MAXISORP 96-well plate (Nunc) and blocked with PBS supplemented with 0.02% Tween 20, 5% BSA, and 5% goat serum. 1 h later, plates were incubated with ZIKV-

**Protein production, purification, and crystallization**

ZIKV DIII (H/PF/2013 and MR-766, residues 299–407), DENV1-DIII (residues 293–399; Austin et al., 2012), WNV DIII and E (New York, 1999; residues 296–401 [DIII] and residues 1–406 [E]; Luca et al., 2012; Nybakken et al., 2005), JEV DIII (SA14-14-2, residues 299–399; Fernandez et al., 2018), POWV DIII (Spuoner, residues 298–399), ZIKV E (H/PF/2013, residues 1–407), and SPOV E (SM6 V-1, residues 1–411) were cloned into the pET21a vector (Novagen) and expressed as inclusion bodies in Escherichia coli BL21(DE3). Isolated inclusion bodies were solubilized and refolded as previously described (Dai et al., 2016; Nybakken et al., 2005). ZIKV-116 heavy and light chain variable regions were codon-optimized for mammalian expression and synthesized on a BioXP 3200 DNA synthesis instrument (SGI-DNA). The resulting gene fragments were cloned directly into CMV-driven mammalian expression vectors that had already been fused with C2l-2-3 for IgGl, C2l4 for Fab, and C2l5 for IgK. The mAb or Fab heavy and light chain vectors were cotransfected into Expi293F or ExpiCHO cells (Thermo Fisher) for transient expression. After 7 d of culture, the culture supernatants were separated by centrifugation, filtered, and purified by fast protein liquid chromatography on an AKTA instrument using a HiTrap MabSelect Sure column (GE Healthcare) or a CaptureSelect IgG-CH1 column (Thermo Fisher), respectively. The purified mAb or Fab then was buffer-exchanged into PBS.

Purified ZIKV-116 Fab was incubated with excess DIII for 2 h followed by complex isolation through size exclusion chromatography (Superdex 75 16/600). The complexes were collected and crystallized in hanging drops at 20°C. Diffraction quality crystals were obtained in the condition of 30% 2-Methyl-2,4-pentanediol, 0.1 M imidazole, and 10% polyethylene glycol 4K, at 80°C.

**Structure determination and refinement**

Diffraction data were collected at the Advanced Photon Source beamline 24-ID-C and processed with the HKL2000 program. Structural phase was determined by molecular replacement in Phenix (Adams et al., 2010) using the Phaser GUI (McCoy et al., 2007), and previously solved ZIKV DIII from the complex of ZV-DIII (PDB 5KVG) and human Fab structure from Z20-e (PDB 5GZO) were used as the search models, respectively. Cycles of model building and refinement were performed in Coot (Emsley et al., 2010) and Phenix. The structural figures were generated in PyMOL software. The interaction of Fab and DIII was analyzed in the ligplot program suite using default settings (McDonald and Thornton, 1994). The buried surfaces and shape
complementarity were calculated using PISA (http://www.ebi.ac.uk/msd-srv/prot_int/).

BLI
The binding affinity of recombinant DIII with ZIKV-116 was monitored by BLI using an Octet-Red96 device (Pall ForteBio). ZIKV-116 and ZIKV-116-V.J.Rev mAbs were biotinylated using EZ-Link-NHS-PEG4-Biotin reagent and then loaded onto streptavidin biosensors (ForteBio). Binding at neutral pH was determined in HBS-EP buffer (10 mM Hepes, pH 7.4, 150 mM NaCl, 3 mM EDTA, 0.005% P20, and 3% BSA). Low pH binding was performed in a similar assay buffer with the pH adjusted to 5.5 using 2-ethanesulfonic acid buffer. Buffer alone with loaded mAb and sensor alone with the highest concentration of DIII were added in parallel to serve as negative controls. The experimental curves were analyzed using Biaevaluation 4.1 (GE Healthcare) and fitted by a 1:1 Langmuir binding model to determine the association constant (k_a), dissociation constant (k_d), kinetic binding affinity (K_D, kinetic), and steady-state affinity (K_D, equilibrium).

Attachment inhibition assay
Attachment blockade was measured by qRT-PCR on an ABI 7500 Real Time-PCR system (Applied Biosystems). 2 × 10^5 Vero cells were seeded in 24-well plates and incubated for 24 h. ZIKV (2 × 10^6 focus-forming units [FFU]) were preincubated with ZIKV-116 or a human isotype control (anti-influenza IgG1) mAb (0.1 µg/ml, 1 µg/ml, 10 µg/ml) for 1 h, followed by addition of the mAb-virus complex to chilled Vero cells for 1 h at 4°C. Cells were extensively washed with cold DMEM on ice, and total RNA was extracted using an RNeasy Mini Kit (Qiagen). ZIKV RNA levels were determined using a Taqman RNA-to-Ct 1-Step Kit (Thermo Fisher) normalized to the internal control GAPDH. Primers and probes used are as follows: ZIKV-E-Fwd, 5'-CCACCAATGTTC TCTTGCAGACATATTG-3'; ZIKV-E-Rev, 5'-TTGGGACAGCCG TTGTCCAACACAG-3'; Probe ZIKV-E, 56-FAM/AGCCTACCT/ZEN/TGACAGGAATATGC/3IABkFQ; GAPDH-Fwd, 5'-TTGAGTTGA GGATTTGGTC/3IABkFQ. The atomic coordinates of ZIKV-116 Fab in complex with ZIKV DIII have been submitted to the RCSB PDB with accession no. 6PLK.

Online supplemental material
Fig. S1 shows that there are two ZIKV-116–DIII complexes in a crystallographic asymmetric unit, which is related to Fig. 2. Fig. S2 shows amino acid differences in DIII between two representative ZIKV strains (H/PF/2013 and MR-766) and two DENV1 strains (16007 and West Pac-74), which is related to Fig. 5. Fig. S3 shows time- and temperature-dependent neutralization curves for ZIKV-116 against ZIKV strain H/PF/2013. Fig. S4 shows the distance of residue 393 within structural ZIKV-116 epitope to residue 204DENV1/209ZIKV from the same E protein or adjacent E protein on the virion surface. Table S1 presents crystallographic data collection and refinement statistics. Tables S2 and S3 provide van der Waals contacts for ZIKV-116–DIII complex. Tables S4–S6 present hydrogen bond contacts between ZIKV-116 and DIII. Table S7 shows the neutralizing values of ZIKV-116 and ZV-67 against different ZIKV strains. Table S8 provides binding parameters for ZIKV-116 to recombinant DIII protein of distinct ZIKV and DENV1 strains measured by BLI. Table S9 shows key resource information used in this study.

Acknowledgments
We acknowledge staff at the Advanced Photon Source Beamlines 24-ID-C/E, especially K. Rajashankar, K. Perry, and N. Sukumar, for aid in data collection.

This work was supported by National Institute of Allergy and Infectious Diseases grant R01AI073755 (to M.S. Diamond and D.H. Fremont), R01AI127828 (to J.E. Crowe Jr. and M.S. Zhao et al. Antbody neutralization of Zika and dengue virus
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References


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Antibody neutralization of Zika and dengue virus

journal of experimental medicine

https://doi.org/10.1088/jem.20191792

Submitted: 23 September 2019
Revised: 9 October 2019
Accepted: 18 October 2019

PhD candidate, University of California, Berkeley. The authors declare no competing financial interests.

Author contributions: H. Zhao, L. Xu, K.A. Dowd, T.C. Pierson, J.E. Crowe Jr., M.S. Diamond, and D.H. Fremont designed the experiments. H. Zhao, L. Xu, and K.A. Dowd performed the experiments. R. Bombardil and R. Nargi generated mAbs and Fab. C.A. Nelson provided SPOV E and JEVDIII proteins. J.M. Errico provided POWV DIII protein. H. Zhao, Z. Deng, and D.H. Fremont performed the cryocrystallography experiments. T.C. Pierson, J.E. Crowe Jr., M.S. Diamond, and D.H. Fremont obtained funding. H. Zhao and D.H. Fremont analyzed the data and wrote the manuscript, with all authors providing editorial comments.

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Journal of Experimental Medicine

https://doi.org/10.1088/jem.20191792

Submitted: 23 September 2019
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Rey, F.A., F.X. Heinz, C. Mandl, C. Kunz, and S.C. Harrison. 1995. The en- velope glycoprotein from tick-borne encephalitis virus at 2 A resolu-

Sirohi, D., and R.J. Kuhn. 2017. Zika virus structure, maturation, and re-