Yellow fever virus capsid protein is a potent suppressor of RNA silencing that binds double-stranded RNA


*Department of Entomology, Texas A&M University, College Station, TX 77843; and †Department of Entomology, Fralin Life Science Institute, Virginia Polytechnic Institute and State University, Blacksburg, VA 24061

Edited by George Dimopoulos, Johns Hopkins School of Public Health, Baltimore, MD, and accepted by Editorial Board Member Carolina Barillas-Mury

Mosquito-borne flaviviruses, including yellow fever virus (YFV), Zika virus (ZIKV), and West Nile virus (WNV), profoundly affect human health. The successful transmission of these viruses to a human host depends on the pathogen’s ability to overcome a potentially sterilizing immune response in the vector mosquito. Similar to other invertebrate animals and plants, the mosquito’s RNA silencing pathway comprises its primary antiviral defense. Although a diverse range of plant and insect viruses has been found to encode suppressors of RNA silencing, the mechanisms by which flaviviruses antagonize antiviral small RNA pathways in disease vectors are unknown. Here we describe a viral suppressor of RNA silencing (VSR) encoded by the prototype flavivirus, YFV. We show that the YFV capsid (YFC) protein inhibits RNA silencing in the mosquito *Aedes aegypti* by interfering with Dicer. This VSR activity appears to be broadly conserved in the C proteins of other medically important flaviviruses, including that of ZIKV. These results suggest that a molecular “arms race” between vector and pathogen underlies the continued existence of flaviviruses in nature.

**Significance**

Until it was demonstrated that the *Aedes aegypti* mosquito transmitted yellow fever, the disease was possibly the most feared pestilence in the western hemisphere. This finding, by Walter Reed’s Yellow Fever Commission, is credited with eradication of the disease in many areas, through sanitation programs designed to eliminate the vector. Since this discovery, the mosquito has been widely believed to be complicit in the transmission of viral diseases, earning the enmity of humans everywhere. However, we show here that yellow fever virus encodes a protein that blocks the mosquito’s immune response, suggesting the pathogen’s continued existence in nature depends on staying one step ahead of the vector’s antiviral defense.

Author contributions: Z.N.A. and K.M.M. designed research; G.H.S., M.W., and K.M.M. performed research; G.H.S., M.W., and A.B. contributed new reagents/analytic tools; G.H.S., Z.N.A., and K.M.M. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission. G.D. is a Guest Editor invited by the Editorial Board.

Freely available online through the PNAS open access option.

Data deposition: The data reported in this paper have been deposited in the Gene Expression Omnibus (GEO) database, www.ncbi.nlm.nih.gov/geo (accession no. GSE80691).

Present address: Center for Genome Sciences, United States Army Medical Research Institute of Infectious Diseases, Fort Detrick, MD 21702.

*To whom correspondence should be addressed. Email: mylesk@tamu.edu.

This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1600544113/-/DCSupplemental.

**Yellow fever virus | flavivirus | capsid | Zika virus | RNA interference**

Arthropod-borne viruses (arboviruses) are maintained in nature through transmission cycles that involve alternating replication in hematophageous insect vectors and vertebrate hosts. The *Flavivirus* genus of the family *Flaviviridae* contains a number of important mosquito-borne viruses responsible for large epidemics of human disease. These include Zika virus (ZIKV), West Nile virus (WNV), dengue virus (DENV; serotypes 1–4), Japanese encephalitis virus (JEV), and the type virus for the family, yellow fever virus (YFV). The YFV genome is a single-stranded RNA of positive polarity that is ~11 kb in length. The 5′-capped RNA encodes the structural proteins (C-prM-E) and nonstructural proteins (NS1-NS2A-NS2B-NS3-NS4A-2K-NS4B-NS5) (1). Nonstructural components are assembled, with host factors, into membrane-bound replication complexes, where the various enzymatic functions of these proteins (e.g., RNA-dependent RNA polymerase, helicase, capping machinery) replicate the genome. The relatively smooth surface of the flavivirus virion is decorated with the envelope (E) and membrane (M) proteins, anchored into the host-derived lipid bilayer via C-terminal transmembrane domains. Within the virion is a nucleocapsid core comprising a single copy of the RNA genome in complex with multiple capsid (C) proteins (2–4). In an urban transmission cycle, the yellow fever mosquito, *Aedes aegypti*, acquires the virus after feeding on an infected human host. After a brief incubation period, the mosquito may transmit the virus to other susceptible humans. Although infection of the vertebrate host is often associated with pathology and disease, a persistent nonlethal infection is established in the insect vector.

When vector mosquitoes become infected with arboviral pathogens, antiviral immunity is essential to the survival of the insect (5). Although a sufficiently robust immune response may be sterilizing, the antiviral defenses of the mosquito are not always an effective barrier to the transmission of viral pathogens. In plant and invertebrate organisms, double-stranded RNA (dsRNA) replicative intermediates (RIs) produced during viral infection activate an antiviral defense based on RNA silencing (6). In flies, the ribonuclease Dicer-2 (Dcr-2) recognizes dsRNA (7). Cleavage of dsRNA RIs by Dcr-2 generates viral small interfering RNAs (siRNAs) ~21 nt in length. These siRNA duplexes are incorporated into the RNA-induced silencing complex (RISC). RISC maturation involves loading a duplex siRNA, choosing and retaining a guide strand, and ejecting the antiparallel passenger strand (8–11). The guide strand directs Argonaute 2 ( Ago-2), an essential RISC component possessing endonuclease activity, to complementary RNAs in the cell, leading to their sequence-specific degradation. When infected with RNA viruses, *Drosophila melanogaster* dcr-2 or ago-2 loss-of-function mutants exhibit an “enhanced disease phenotype,” characterized by elevated levels of virus replication and increased mortality (12–14).

In an evolutionary “arms race,” the genomes of numerous plant and insect viruses have evolved to encode one or more viral suppressors of RNA silencing (VSRs) (15, 16). For example, the flock house virus (FHV) B2 is a well-characterized dsRNA-binding protein that shields the RIs produced during viral infection from processing by Dcr-2 and interferes with the incorporation of vsiRNA duplexes into the RISC (17–20). Well-documented examples of plant and insect virus proteins that interfere with small RNA pathways suggest that arboviruses may encode similar viral suppressors of RNA silencing.
proteins; however, the mechanisms by which medically important arboviruses modulate small RNA pathways to productively infect disease vectors remain unknown. Elucidating these mechanisms would fundamentally alter our understanding of pathogen transmission by arthropod vectors.

We have previously shown that an antiviral response directed by vsiRNAs is essential to limiting the pathogenesis of arbovirus infections in mosquitoes (5). Suppressing the accumulation of vsiRNAs in *Ae. aegypti* by infection with a recombinant Sindbis virus (SINV) expressing the FHV B2 resulted in elevated levels of virus replication and increased mortality, i.e., a disease phenotype. In the present study, we show that this phenotype can be used to reliably identify VSRs that interfere with RNA silencing by diverse mechanisms. Thus, to investigate the precise presence of VSRs encoded in the genomes of flaviviruses and other mosquito-borne pathogens, we infected *Ae. aegypti* with recombinant SINVs (*Alphavirus* genus) expressing heterologous virus sequences. Here we report suppression of RNA silencing in *Ae. aegypti* mosquitoes infected with a recombinant SINV expressing the yellow fever virus capsid (YF). Furthermore, through a series of biochemical assays, we demonstrate that the observed antagonism of RNA silencing is mediated by the binding of YFC to long dsRNAs, interfering with the production of vsiRNAs by Dicer.

**Results**

Studies identifying VSRs have often been guided by previous experimental evidence implicating a particular viral protein or sequence as a virulence factor. Infection of an invertebrate host with arboviruses tends to be avirulent, however, making the identification of VSR activity in these viruses more challenging. We previously showed that expression of FHV B2 protein from recombinant alphaviruses dramatically increases the virulence of infection in the mosquito host, resulting in a disease phenotype (5). We hypothesized that this phenotype would be predictive of VSR functions in other medically important flaviviruses, including that of ZIKV.

**Fig. 1.** A disease phenotype in *Ae. aegypti* infected with recombinant SINVs expressing heterologous VSR proteins. (A) Schematic of a recombinant SINV expressing a VSR from an engineered subgenomic promoter (SG2). (B) Survival of mosquitoes after infection with recombinant SINVs expressing well-known VSR proteins (red triangles) or with viruses containing non-translatable versions of the VSR sequences (black circles). Survival curves represent cohorts of ≥40 adult female mosquitoes injected with 500 pfu of virus. Significance was determined using the Mantel-Cox log-rank test.

As an initial test of this hypothesis, we created recombinant SINV's expressing well-characterized VSR proteins from a duplicated subgenomic promoter (Fig. 1A). *Ae. aegypti* mosquitoes were infected with SINV expressing the Cricket paralysis virus 1A protein (SINV/CrPV-1A) (12, 21), *D. melanogaster* C virus 1A protein (SINV/DCV-1A) (14), Carnation Italian ringspot tombusvirus P19 protein (SINV/CIRV-P19) (22–24), or the influenza A virus NS1 protein (SINV/H1N1-NS1) (25–27). Control viruses were engineered to contain nontranslatable versions of these proteins expressed from the second subgenomic promoter. We previously showed that the survival of *Ae. aegypti* infected with recombinant SINV is distinguishable from that of uninfected mosquitoes (5). In the present experiment, however, mortality was significantly higher in *Ae. aegypti* infected with SINV/CrPV-1A, SINV/DCV-1A, SINV/CIRV-P19, or SINV/H1N1-NS1 compared with mosquitoes infected with SINV or control viruses expressing nontranslatable transcripts (Fig. 1B). These results suggest a strong correlation between the expression of heterologous VSR proteins and a disease phenotype in mosquitoes infected with SINV, even under circumstances in which the heterologous VSRs are derived from a diverse range of viruses employing different mechanisms of RNA interference (RNAi) suppression.

We next tested whether the expression of candidate VSRs, previously identified in the genomes of WNV, DENV, and Bunyamwera virus (BUNV; *Orthobunyavirus* genus), correlated with a disease phenotype in mosquitoes infected with SINV. *Ae. aegypti* were infected with viruses expressing the WNV sfRNA (SINV/WNV-sfRNA) (28), DENV NS4B protein (SINV/DENV-NS4B) (29), or BUNV NSs protein (SINV/BUNV-NSs) (30). Control viruses expressed inactive mutant versions of the WNV sfRNA or BUNV NSs protein, containing the previously described deletions sfRNA IRA (31) or delNSs (30), respectively. In contrast to the results described above, there was no significant difference between the survival of mosquitoes infected with control viruses and those infected with SINV/WNV-sfRNA, SINV/DENV-NS4B, or SINV/BUNV-NSs (Fig. 2A). Real-time PCR (RT-PCR) analysis of SINV RNA in the infected mosquitoes corroborated the survival results, indicating no significant increase in levels of virus replication in mosquitoes infected with SINV/ WNV-sfRNA, SINV/DENV-NS4B, or SINV/BUNV-NSs compared with mosquitoes infected with control viruses (Fig. 2B). These results suggest that the WNV sfRNA, DENV NS4B, and BUNV NSs provide little to no protection from an RNA silencing response induced by virus replication in *Ae. aegypti*.

Whereas a disease phenotype is evident in mosquitoes infected with alphaviruses expressing heterologous VSRs, similar experiments had not been performed with flaviviruses. Therefore, we assessed the replication of YFV in *Ae. aegypti* in which Dicer-2 has been genetically ablated (32). Our results show that the replication of YFV is significantly higher in dcr-2 null mutant mosquitoes compared with wild type (WT) mosquitoes at the same time point, suggesting that RNAi is an important antiviral mechanism targeting flaviviruses in the vector host (Fig. S1). Although not a feature of all VSRs, dsRNA binding is essential to the antiviral activity of many well-characterized proteins that interfere with small RNA pathways (14, 17–20, 32, 33). Even though a dsRNA-binding function has not previously been assigned to the flavivirus capsid, the protein has physical properties common to other VSRs that indiscriminately bind dsRNAs, shielding them from Dicer and RISC assembly; for example, evidence indicates that the flavivirus capsid protein binds RNA through nonspecific electrostatic interactions (35, 36), similar to FHV B2 (17–20) and Influenza A virus NS1 (25–27). Therefore, we assessed the potential of the YFC protein to act as an antagonist of antiviral immunity in the disease vector host. To do so, we infected *Ae. aegypti* with a virus expressing the YFC protein (SINV/YFC) or a control virus containing a nontranslatable version of the protein-coding sequence (SINV/YFC-NT). Similar to the results obtained with well-known VSR proteins, mortality was
significantly higher in mosquitoes infected with SINV/YFC compared with those infected with the virus containing the nontranslatable version of the protein-coding sequence (Fig. 3A). Also consistent with a disease phenotype, RT-PCR analysis indicated significantly higher levels of virus replication in mosquitoes infected with SINV/YFC compared with those infected with SINV/YFC-NTR (Fig. 3B).

To confirm that the observed disease phenotype in mosquitoes infected with SINV/YFC is due to a specific VSR activity present in the expressed YFV capsid, and not to some other function of the protein, we infected dcr-2 null mutant Ae. aegypti defective for the production of siRNAs with the recombinant SINVs (32). Not surprisingly, dcr-2 null mutant mosquitoes infected with the control virus also exhibited a disease phenotype, as would be expected in the absence of a functional siRNA pathway (12–14). However, in contrast to infections of the WT mosquitoes, survival curves for the dcr-2 mosquitoes infected with SINV/YFC or SINV/YFC-NTR were remarkably similar, indicating that in the absence of the primary antiviral response, the expression of YFC did not dramatically enhance virus pathogenicity (Fig. 3C). Consistent with this, the replication of SINV/YFC-NTR did not differ significantly from that of SINV/YFC in the mutant mosquitoes (Fig. 3D), indicating that the dcr-2 null genotype genetically rescued replication of the control virus. Overall, these experiments suggest VSR activity in YFC capable of suppressing RNA silencing in the natural disease vector host, Ae. aegypti.

Because small RNAs ultimately serve as the effector molecules that direct the sequence-specific degradation of cognate target RNAs in the cell, we next analyzed the ratio of viral small RNAs (through Illumina-based sequencing) to viral RNAs in mosquitoes infected with recombinant SINVs (Fig. 3B and Fig. S2). The number of vsiRNAs targeting SINV/YFC decreased approximately fourfold compared with those targeting SINV/YFC-NTR in a WT genetic background (Fig. 4A), suggesting that the VSR activity of YFC is mediated by specific inhibition of Dicer. Thus, to further characterize the VSR activity of the YFV capsid, we expressed and purified a polyhistidine-tagged (His-tag) YFC protein (His-YFC) from Escherichia coli.

To assess the effect of His-YFC on the processing of a dsRNA substrate into siRNAs, we performed in vitro dicing assays. In the absence of His-YFC, dsRNA was efficiently processed into siRNAs by the recombinant Dicer enzyme (Fig. 4B, lanes 1 and 2); however, addition of His-YFC to the in vitro dicing assay inhibited the production of siRNAs in a dose-dependent manner (Fig. 4B, lanes 3–5). Similarly, His-tagged influenza A virus NS1, a well-characterized dsRNA-binding protein with previously demonstrated VSR activity (25–27), also inhibited dicing of the dsRNA substrate in this assay (Fig. 4B, lanes 6–8). In contrast, an unconjugated His-tag peptide or a green fluorescent protein (GFP) had no effect on the activity of Dicer (Fig. 4B, lanes 9 and 10). These results confirm that YFC inhibits cleavage of dsRNA substrates by Dicer.

To determine whether the mechanism of Dicer inhibition by the YFC is similar to that of NS1 (i.e., nonspecific binding to dsRNAs) (25–27), we performed electrophoretic mobility shift assays (EMSAs) with both a long dsRNA and a 21-bp duplex siRNA. Although YFC was able to bind the longer dsRNA with high affinity (Fig. 4C), there was little evidence of interaction of the recombinant protein with the shorter duplex siRNA (Fig. 4D), suggesting that YFC does not interfere with loading of the RISC. Whereas there was an observable shift in the mobility of both the long dsRNA and the duplex siRNA with the recombinant NS1 protein, there was no interaction of either RNA with the unconjugated His-tag, even at the equivalent highest concentration. Finally, we also observed binding of YFC with long ssRNAs derived from YFV or luciferase sequences (Fig. S3), indicating that the recombinant protein is able
pared with the capsid of ZIKV (SINV/ZIKC), WNV (SINV/WNC), DENV-2 infected with SINV, we generated recombinant SINVs expressing interferes with the cleavage of long dsRNAs by Dicer.

The YFV capsid binds dsRNA and interferes with Dicer processing. Fig. 4.

A disease phenotype in Ae. aegypti infected with recombinant SIN viruses expressing flavivirus capsid proteins (red triangles). Significance was determined by comparing mosquito survival with recombinant SIN containing a nontranslatable YFC (black circles; replicated for each panel containing a nontranslatable YFC (black circles; replicated for each panel to the survival of mosquitoes infected with recombinant SINV or negative-stranded DENV RNAs of differing lengths is consistent with the severe morbidity and mortality they cause in humans. Transmission of these viruses to human or animal hosts depends on the pathogen’s ability to overcome the potentially sterilizing RNAi-based immune response of the mosquito vector; however, the specific mechanisms by which this may occur remain unknown.

There are numerous examples of VSRs encoded in the genomes of plant and insect viruses, which presumably evolved as a result of an evolutionary arms race with antiviral silencing pathways (15, 16). However, for various reasons, interrogating mosquito-borne viruses for the presence of VSR proteins has proven challenging. In this paper, we describe a simple but effective assay that is highly informative with respect to identifying VSR functions in proteins and other sequences. With this assay, we have identified a VSR encoded in the genome of YFV, the prototype flavivirus. Results obtained both in vivo and in vitro indicate that the YFC protein antagonizes RNA silencing by binding long dsRNAs with high affinity, interfering with efficient processing by Dicer. Our studies also suggest that this VSR function is broadly conserved among flavivirus C proteins.

Precedent for a VSR function in a viral structural protein can be found in the coat protein (CP) of Turnip crinkle virus, which has been shown to suppress RNA silencing through a mechanism that likely also involves the binding of dsRNAs (38). In comparison with other positive-strand viruses, the flavivirus nucleocapsid has a poorly defined and unique architecture (2, 3, 39). Reported production of noninfectious virus-like particles by overexpressing only precursor membrane (prM) and E protein indicates that the C protein is not required for formation of the virion’s spherical shell (40, 41). Indeed, there appears to be little interaction between the C protein and the E and M proteins, raising questions about how nucleocapsids are efficiently incorporated into budding membranous structures containing the flavivirus surface proteins (42). A packaging signal has not been identified in genomes of flaviviruses, and in vitro formation of nucleocapsid-like particles with positive- or negative-stranded DENV RNAs of differing lengths is consistent with this.

to bind nonspecifically to both dsRNAs and ssRNAs. Overall, these results demonstrate that YFC has a dsRNA-binding function that interferes with the cleavage of long dsRNAs by Dicer.

To investigate a possible correlation between the expression of other flavivirus C proteins and a disease phenotype in mosquitoes infected with SINV, we generated recombinant SINVs expressing the capsid of ZIKV (SINV/ZIKC), WNV (SINV/WNC), DENV-2 (SINV/DENC), or Rio Bravo virus (RBV; SINV/RBC). Compared with Ae. aegypti infected with SINV/YFC-NTR, mosquitoes infected with the recombinant SIN viruses expressing the heterologous C proteins exhibited evidence of viral pathogenicity (Fig. 5). Given that these experiments included the C proteins of the genetically distant ZIKV, DENV-2 (vectored by Ae. aegypti), WNV (vectored by culex species mosquitoes), and a representative of the group of flaviviruses without any known vector, RBV (isolated from bats) (37), these results strongly suggest that the VSR activity identified in the YFC is broadly conserved in the C proteins of other flaviviruses as well.

Discussion

Although many questions remain regarding the role of mammalian small RNA pathways in antiviral defense, it is clear that the presence of conserved antiviral silencing pathways in invertebrate organisms plays an important role in the transmission of agents of human disease. Mosquito-borne viruses are responsible for a broad spectrum of human diseases, including arthritis, hemorrhagic fever, encephalitis, and recently microcephaly and Guillian-Barré syndrome. Members of the Flavivirus genus, such as ZIKV, WNV, DENV 1–4, JEV, and YFV, are among the most important mosquito-borne pathogens because of their widespread prevalence and the severe morbidity and mortality they cause in humans. Transmission of these viruses to human or animal hosts depends on the pathogen’s ability to overcome the potentially sterilizing RNAi-based immune response of the mosquito vector; however, the specific mechanisms by which this may occur remain unknown.

There are numerous examples of VSRs encoded in the genomes of plant and insect viruses, which presumably evolved as a result of an evolutionary arms race with antiviral silencing pathways (15, 16). However, for various reasons, interrogating mosquito-borne viruses for the presence of VSR proteins has proven challenging. In this paper, we describe a simple but effective assay that is highly informative with respect to identifying VSR functions in proteins and other sequences. With this assay, we have identified a VSR encoded in the genome of YFV, the prototype flavivirus. Results obtained both in vivo and in vitro indicate that the YFC protein antagonizes RNA silencing by binding long dsRNAs with high affinity, interfering with efficient processing by Dicer. Our studies also suggest that this VSR function is broadly conserved among flavivirus C proteins.

Precedent for a VSR function in a viral structural protein can be found in the coat protein (CP) of Turnip crinkle virus, which has been shown to suppress RNA silencing through a mechanism that likely also involves the binding of dsRNAs (38). In comparison with other positive-strand viruses, the flavivirus nucleocapsid has a poorly defined and unique architecture (2, 3, 39). Reported production of noninfectious virus-like particles by overexpressing only precursor membrane (prM) and E protein indicates that the C protein is not required for formation of the virion’s spherical shell (40, 41). Indeed, there appears to be little interaction between the C protein and the E and M proteins, raising questions about how nucleocapsids are efficiently incorporated into budding membranous structures containing the flavivirus surface proteins (42). A packaging signal has not been identified in genomes of flaviviruses, and in vitro formation of nucleocapsid-like particles with positive- or negative-stranded DENV RNAs of differing lengths is consistent with this.

to bind nonspecifically to both dsRNAs and ssRNAs. Overall, these results demonstrate that YFC has a dsRNA-binding function that interferes with the cleavage of long dsRNAs by Dicer.

To investigate a possible correlation between the expression of other flavivirus C proteins and a disease phenotype in mosquitoes infected with SINV, we generated recombinant SINVs expressing the capsid of ZIKV (SINV/ZIKC), WNV (SINV/WNC), DENV-2 (SINV/DENC), or Rio Bravo virus (RBV; SINV/RBC). Compared with Ae. aegypti infected with SINV/YFC-NTR, mosquitoes infected with the recombinant SIN viruses expressing the heterologous C proteins exhibited evidence of viral pathogenicity (Fig. 5). Given that these experiments included the C proteins of the genetically distant ZIKV, DENV-2 (vectored by Ae. aegypti), WNV (vectored by culex species mosquitoes), and a representative of the group of flaviviruses without any known vector, RBV (isolated from bats) (37), these results strongly suggest that the VSR activity identified in the YFC is broadly conserved in the C proteins of other flaviviruses as well.

Discussion

Although many questions remain regarding the role of mammalian small RNA pathways in antiviral defense, it is clear that the presence of conserved antiviral silencing pathways in invertebrate organisms plays an important role in the transmission of agents of human disease. Mosquito-borne viruses are responsible for a broad spectrum of human diseases, including arthritis, hemorrhagic fever, encephalitis, and recently microcephaly and Guillian-Barré syndrome. Members of the Flavivirus genus, such as ZIKV, WNV, DENV 1–4, JEV, and YFV, are among the most important mosquito-borne pathogens because of their widespread prevalence and the severe morbidity and mortality they cause in humans. Transmission of these viruses to human or animal hosts depends on the pathogen’s ability to overcome the potentially sterilizing RNAi-based immune response of the mosquito vector; however, the specific mechanisms by which this may occur remain unknown.

There are numerous examples of VSRs encoded in the genomes of plant and insect viruses, which presumably evolved as a result of an evolutionary arms race with antiviral silencing pathways (15, 16). However, for various reasons, interrogating mosquito-borne viruses for the presence of VSR proteins has proven challenging. In this paper, we describe a simple but effective assay that is highly informative with respect to identifying VSR functions in proteins and other sequences. With this assay, we have identified a VSR encoded in the genome of YFV, the prototype flavivirus. Results obtained both in vivo and in vitro indicate that the YFC protein antagonizes RNA silencing by binding long dsRNAs with high affinity, interfering with efficient processing by Dicer. Our studies also suggest that this VSR function is broadly conserved among flavivirus C proteins.

Precedent for a VSR function in a viral structural protein can be found in the coat protein (CP) of Turnip crinkle virus, which has been shown to suppress RNA silencing through a mechanism that likely also involves the binding of dsRNAs (38). In comparison with other positive-strand viruses, the flavivirus nucleocapsid has a poorly defined and unique architecture (2, 3, 39). Reported production of noninfectious virus-like particles by overexpressing only precursor membrane (prM) and E protein indicates that the C protein is not required for formation of the virion’s spherical shell (40, 41). Indeed, there appears to be little interaction between the C protein and the E and M proteins, raising questions about how nucleocapsids are efficiently incorporated into budding membranous structures containing the flavivirus surface proteins (42). A packaging signal has not been identified in genomes of flaviviruses, and in vitro formation of nucleocapsid-like particles with positive- or negative-stranded DENV RNAs of differing lengths is consistent with this.
with the idea that these sequences do not exist (36). Rather, interactions between RNA and the C protein appear to be mediated by nonspecific electrostatic interactions (35, 36), and our results are consistent with this idea. Evidence indicates that packaging requires active synthesis of viral RNA, suggesting that replication and encapsidation are tightly coupled (43). This may explain the apparent specificity with which genomic RNA is packaged into the virion, despite a lack of organization in the way in which C proteins and viral RNA assemble (2, 3, 39). Our results, presented here, suggest that the C protein interacts with viral dsRNA RiIs in the infected cell.

Similar to other positive-strand viruses, the replication complexes of flaviviruses are believed to assemble on internal cellular membranes (42). In DENV-infected cells, this process involves extensive rearrangements of the endoplasmic reticulum to form a continuous network of vesicles consisting of series of invaginations of the endoplasmic reticulum membrane (44). Similar structures have been observed in mosquito cells infected with DENV (45). Detection of dsRNA within membrane vesicles suggests that these are sites of viral replication within the infected cell (44, 45). Although evidence suggests that membrane vesicles are connected to the cytosol by pores (44, 45), it has been postulated that these structures may protect the viral RNA from the cellular components of innate immunity (46). However, the existence of VSR proteins encoded in the genomes of flaviviruses suggests that replication complexes are accessible to RNA silencing pathways, and possibly other antiviral responses as well.

In mammalian cells, viral infection induces the synthesis and secretion of type I IFNs (α/β). Activation of signaling pathways by the secreted IFN results in the transcription of hundreds of genes, contributing to the establishment of a general antiviral state in the cell. A number of viruses have been found to express proteins that are antagonists of the IFN α/β response. Similar to VSR proteins, viruses have evolved diverse mechanisms for inhibiting the IFN α/β response (47). Also similar to VSR proteins, dsRNA binding is a molecular mechanism by which some viral proteins inhibit the IFN α/β system. For example, the dsRNA-binding function of the influenza NS1 protein counteracts the establishment of an antiviral state in the cell by blocking virus-mediated activation of cellular transcription factors that induce synthesis of IFN α/β (26, 27, 47). Thus, a particularly intriguing possibility is that the dsRNA-binding domain of the flavivirus C protein is capable of blocking RNAi of the mosquito, suggesting coevolution of these viruses in response to the strong selective pressure exerted by the RNA silencing pathways of the host, a classic “red queen hypothesis.” Thus, a molecular arm race occurring between vector and pathogen underlies the continued existence of these viruses in nature, illustrating the importance of RNA silencing pathways in the transmission of many important human diseases.

Materials and Methods

Mosquito Infections. Ae. aegypti (Liverpool) WT and dcr-2 null strains were reared at 28 °C and 80% relative humidity, with a 14/10-h day/night light cycle. dcr-2 null mosquitoes were obtained as described previously (32). WT null mosquitoes were 8 d old due to the screening process. dcr-2 null mosquitoes were 8 d old due to the screening process.

Recombinant Virus Production. Recombinant SIN viruses were generated by inserting heterologous sequences encoding known or suspected VSRs into a multiple cloning site, located downstream of a duplicated subgenomic promoter. Recombinant viruses were rescued as described previously (5).

RNA Isolation and Detection. Total RNA was extracted from pools of mosquitoes with Tri Reagent RT (Molecular Research Center) according to the manufacturer's instructions. SINV mRNA levels were determined using a strand-specific quantitative RT-PCR Taqman assay (Life Technologies) as described previously (54), with the following primers and Taqman probe: forward, 5′-ATACAATA-TGCGAACAGAAAGAAG-3′; reverse, 5′-CTGTCGTTGCGAAATGTTG-3′; probe, 5′-CTAAAACAGCGCCGAACCT-3′. The following primers and Taqman probe were used for YFV: forward, 5′-GGTTCAATGAGGCGTGGCTAT-3′; reverse, 5′-GGCGAGCAGGCTAAGATTG-3′; probe, 5′-CAAGCTGAAGATGGGGT-3′. A minimum of three independent biological replicates were tested, and results were analyzed using the t test to determine significance.

Small RNA Library Preparation and Analysis. Libraries were prepared from total RNA isolated with Tri Reagent (Molecular Research Center) from adult female mosquitoes at 96 h after infection with recombinant SINVs. In brief, small
RNAs (18–35 nt) were recovered by PAGE separation, and libraries were prepared with the TruSeq Small RNA Sample Preparation Kit (Illumina) according to the manufacturer’s instructions. To reduce sources of nonbiological variation, libraries were multiplexed and biological replicates were sequenced in a single lane of a HiSeq (Illumina) flow cell. Following removal of the 3′ adaptors, 10 million small RNA reads were selected at random from each dataset and mapped to the SINV genome. Differential expression of sRNAs was determined as described previously (55). The small RNA libraries used in this study are available for download from the Gene Expression Omnibus (accession no. GSE80691).

**Dicing Assays.** Processing of long dsRNA into sRNAs was analyzed with an ultra-active form of recombinant human Dicer, capable of cleaving more than 95% of dsRNA template into sRNAs within 2 h (Genentech, South San Francisco, CA). The small RNA libraries used in this study are available for download from the Gene Expression Omnibus (accession no. GSE80691).

**Processing of long dsRNA into siRNAs was analyzed with an ultra-active form of recombinant human Dicer, capable of cleaving more than 95% of dsRNA template into sRNAs within 2 h (Genentech, South San Francisco, CA). The small RNA libraries used in this study are available for download from the Gene Expression Omnibus (accession no. GSE80691).**

**ACKNOWLEDGMENTS.** This work was supported by National Institute for Allergy and Infectious Diseases Grants AI077726, AI103265, and AI119081. The funding agency had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.