Overlapping Local and Long-Range RNA-RNA Interactions Modulate Dengue Virus Genome Cyclization and Replication

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The dengue virus genome is a dynamic molecule that adopts different conformations in the infected cell. Here, using RNA folding predictions, chemical probing analysis, RNA binding assays, and functional studies, we identified new *cis*-acting elements present in the capsid coding sequence that facilitate cyclization of the viral RNA by hybridization with a sequence involved in a local dumbbell structure at the viral 3' untranslated region (UTR). The identified interaction differentially enhances viral replication in mosquito and mammalian cells.

Dengue virus (DENV) is a member of the *Flaviviridae* family that includes other important pathogens such as yellow fever virus (YFV), West Nile virus (WNV), Saint Louis encephalitis virus (SLEV), and Japanese encephalitis virus (JEV). The DENV genome is a plus-stranded RNA molecule that contains a single open reading frame flanked by highly structured 5' and 3' untranslated regions (UTRs) (1–3). RNA elements located within these regions are responsible for translation initiation and genome replication (4–7). The 5' UTR is about 100 nucleotides (nt) long and includes three different elements: (i) stem-loop A (SLA), which is the promoter for viral polymerase binding and activation (8–10); (ii) stem-loop B (SLB), which contains a sequence known as 5' upstream of the AUG region (5' UAR) that is complementary to a sequence present at the 3' UTR (3' UAR) and mediates long-range RNA-RNA interactions between the ends of the genome (11); and (iii) a spacer sequence between SLA and SLB rich in U's, which functions as an enhancer of viral replication (10). The viral 3' UTR is about 450 nucleotides long and comprises four defined domains: domain A1, which features a variable region (VR) (12); domains A2 and A3, which present two almost-identical dumbbell-like secondary structures (DB1 and DB2), which appear to work as enhancers for viral RNA replication (13–15); and domain A4, which contains a small hairpin (sHP) and the 3' stem-loop (3' SL), which are essential elements for viral replication (3, 16). In addition to RNA structures defined in the UTRs that play different roles during infection, important RNA elements have been described in the protein coding region. In this regard, essential sequences that mediate long-range RNA-RNA interactions known as 5' cyclization sequence (5' CS) and 5' downstream of AUG region (5' DAR) are located within the capsid coding sequence (11, 13, 17–21). Also, a hairpin known as cHP, located between 5' CS and 5' DAR, has been shown to be necessary for efficient RNA replication (22). The current model for viral RNA synthesis includes the interaction of the viral polymerase NS5 with the 5'-end SLA promoter and its transfer to the 3'-end initiation site by cyclization of the viral genome (9). Despite great advances in knowledge of *cis*-acting RNA elements in the flavivirus genomes, the molecular details and mechanisms by which many of them function during viral replication are still not well understood.

Intrigued by dual roles of RNA sequences in viral protein coding regions, we examined the complete capsid coding sequence for the presence of new *cis*-acting RNA elements. First, we used representative sequences of the six different genotypes of DENV type

2 (DENV2) to examine the conservation of the first 450 nucleotides of the viral genome, including the 5' UTR and the complete capsid coding sequence. The highest nucleotide conservation was observed in the first 300 nucleotides of the viral genome (Fig. 1A). To evaluate the ability of the nucleotide sequences to form RNA secondary structures, we performed an evolutionary conservation analysis using RNAz and RNAalifold software (23–25). Interestingly, the base pairing probability plot drastically dropped around nucleotide 310 (region indicated as C3 in Fig. 1B), showing high double-strand probability for 5' SLA, SLB, and cHP structures (black boxes) as expected but also highlighting two additional structured regions indicated as C1 and C2 (gray boxes). To confirm the RNA structure predictions, a fragment of the first 450 nt of the viral genome was probed with *N*-methylisotoic anhydride (NMIA) employing SHAPE (selective 2'-hydroxyl acylation analyzed by primer extension) technology (26, 27). This chemical preferentially reacts with unpaired nucleotides and poorly with those that are constrained in base pairs. The samples were resolved by capillary electrophoresis, and data were analyzed using ShapeFinder software v1.0 (28). NMIA reactivity of each nucleotide position is color coded and shown in a summary plot in Fig. 1C. The secondary structure prediction was obtained using RNAstructure software v5.4 (29–31) (Fig. 1D). A notable correlation of the pairing probability and the SHAPE reactivity was observed (compare Fig. 1B and C), showing high SHAPE reactivity downstream of nucleotide 310. Importantly, the analysis supports the presence of RNA structures within nucleotides 150 and 310 of the genome (C1 and C2 regions). For the C1 region, two small hairpins were

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FIG 1 Conserved RNA structures in the DENV capsid coding sequence. (A) Nucleotide conservation of the first 450 nucleotides of the viral genome using representative sequences of the six different DENV2 genotypes. Window size = 10. (B) Double-stranded probability of the first 450 nucleotides of the viral genome. Three regions were defined: high double-strand probability for 5' SLA, SLB, and cHP (black boxes); C1 and C2 structures (gray boxes); and unstructured C3 region (white boxes). (C) Plot showing SHAPE reactivity at each nucleotide position of DENV2 RNA. (D) Viral RNA structures based on conservation, predictions, and SHAPE reactivity. Red notations correspond to unpaired nucleotides, and black indicates double-stranded regions, whereas residues of intermediate reactivity are noted in orange. Abbreviations: UTR, untranslated region; SLA, stem-loop A; SLB, stem-loop B; cHP, capsid hairpin; 5' CS, 5' cyclization sequence; PK, pseudoknot; C1, C2, and C3, RNA structures identified in the capsid coding sequence. Data shown are averages of three experiments.

mapped. Nucleotides located in the top of the first hairpin (GAGAA) and a downstream complementary region (UUCUC) were unreactive for NMIA, predicting the formation of a pseudoknot structure (PK) (Fig. 1D), in agreement with data recently reported using DENV4 (32). In the C2 region, two large stem-loops were predicted, while in the C3 region, low-stability structures were observed.

In order to study the functional significance of these RNA ele-

ments, a DENV2 system previously described (33), carrying a luciferase gene, was modified to generate a new construct that allowed manipulation of the complete coding sequence of capsid but retained the ability to produce infectious particles. This construct allowed uncoupling *cis*-acting signals from coding sequences of capsid. The new reporter virus (named FullCapDV-Luc) contained the viral 5' UTR, the complete capsid coding region followed by the *Renilla* luciferase gene, which was flanked

FIG 2 Functional significance of RNA structures in the capsid coding sequence. (A) Schematic representation of the DENV reporter FullCapDVLuc. FullCap and MCAE stand for full capsid coding sequence and minimal *cis*-acting elements, respectively. (B) Replication of new DENV reporter construct. Plots show *Renilla* luciferase activity as a function of time post-RNA transfection in BHK and C6/36 cells. The luciferase values are means $\pm$ standard deviations ($n = 4$). (C and D) Translation and replication of DENV reporter RNAs containing the indicated deletions in BHK cells (C) and C6/36 cells (D). Luciferase activity was measured as a function of time and is represented as arbitrary units (a.u.). The luciferase values are means $\pm$ standard deviations ($n = 4$).

by two foot-and-mouth disease virus 2A (FMDV2A) protease coding sequences (QLNFDLLKLAGDVESNPGP), and the rest of the viral genome, including a second copy of the capsid protein coding sequence followed by the rest of the open reading frame and the 3' UTR (Fig. 2A). Two FMDV2As were introduced to ensure the release of the luciferase and avoid changes in its enzymatic activity due to fusion of additional amino acids of the capsid coding sequence. In order to evaluate the ability of this virus to replicate, the *in vitro*-transcribed RNA was transfected into mosquito (C6/36) and mammalian (BHK) cells, and luciferase activity was measured as a function of time (Fig. 2B). A replication-impaired control (mutations in the polymerase NS5) was also constructed in the described backbone (MutNS5). The results indi-

cate that the reporter FullCapDVLuc is fully active for translation and replication in both cell types.

To examine the requirement of the predicted RNA structures in the capsid coding sequence, we constructed a set of mutants with a deletion of each structure (Mut Δ C1, Mut Δ C2.1, Mut Δ C2.2, Mut Δ C3, and Mut Δ C123) in the context of the FullCapDVLuc. The mutants Mut Δ C2.1 and Mut Δ C2.2 correspond to deletions of each of the two hairpins of structure C2 (Fig. 2A). RNAs corresponding to the mutants were transfected into C6/36 and BHK cells and monitored by luciferase activity as a function of time. The mutant with simultaneous deletion of all three regions, Mut Δ C123, showed delayed and reduced replication in both cell lines. A reduction of about 100-fold with respect to wild-type

(WT) control was observed at 48 h in BHK cells, while the reduction was more than 1,000-fold in C6/36 cells at 72 h (Fig. 2C and D, black arrows). Mutants Δ C2.1, Δ C2.2, and Δ C3 showed luciferase levels that were similar to those of the WT FullCapDVLuc in both cell lines, indicating that these structures do not play an important role in translation or RNA synthesis. On the other hand, the reporter Mut Δ C1 presented a delay and a 10-fold reduction in viral replication in mosquito cells, and a 6-fold reduction in mammalian cells, at 72 and 48 h, respectively (Fig. 2C and D). Although deletion of C1 showed a clear effect on viral replication, deleting each structure individually was not as drastic as the deletion of the 3 elements at the same time, suggesting an additive effect when these RNA structures are deleted collectively. For all the transfected viral RNAs, similar levels of luciferase activity were observed at 4 h, indicating that the deletions tested did not alter translation of the input RNA. The results suggest that the RNA structures present in the capsid coding sequence are important for efficient DENV amplification, with a more pronounced requirement when the virus replicates in mosquito cells than when it replicates in mammalian cells (compare Fig. 2C and D). It is important to mention that the deleted RNA structures are duplicated downstream (in the coding region of the capsid protein); thus, we cannot rule out a possible underestimation of the effect observed by RNA structure deletions due to complementation via the duplicated structures.

It has been suggested in a recent report that RNA elements present in the capsid coding sequence of DENV4, DCS-PK, homologous to C1, could modulate the conformation of RNA elements at the 5' end (5' UAR, 5' DAR, cHP, and/or 5' CS) during RNA cyclization (32); however, no evidence of how this RNA structure would participate during this viral process was provided. Based on the finding that the C1 region enhances viral RNA synthesis, we examined a possible direct contribution of its sequence in long-range RNA-RNA interactions. First, we evaluated conserved patterns of possible RNA-RNA interaction based on sequence alignments of the capsid coding regions and 3' UTRs from different DENV2 genotypes. To this end, the RNAaliduplex software was used for the analysis (25), and the R-CHIE package was used to plot the data (34). The program takes two RNA sequence alignments, predicts optimal and suboptimal interactions and hybridization energies, and provides information about conserved interactions. The alignments and details of specific sequences used for this study are described in Fig. S1 in the supplemental material. Interestingly, this analysis resulted in a model that predicted a novel conserved interaction between a C1 sequence and a sequence contained in the conserved DB1 structure of the 3' UTR (Fig. 3A and B). The predicted model is complex because it involves the interaction between sequences contained in C1 and DB1 structures, which also form local PK interactions. The PK formed between the conserved nucleotides at the loop of DB1 (GCUGU) and downstream sequence (CGACG) has been well characterized in previous studies (14, 35, 36). The proposed long-range RNA-RNA hybridization competes and disrupts not only the stem-loops in C1 and DB1 but also the two PKs. Thus, the C1-DB1 long-range hybridization and the C1-DB1 local structures represent mutually exclusive forms of the DENV genome, which may exist in equilibrium in the infected cell. We have previously described that alternative conformations of the DENV genome, linear and circular, are necessary for viral RNA replication and that a balance between these forms of the genome is crucial for

DENV infectivity (37). To confirm the formation of the new predicted structures, SHAPE studies using 5' UTR-capsid and 3' UTR RNAs were performed. Although most of the nucleotides involved in the long-range RNA-RNA interaction are also involved in local base pairings, we detected nucleotides (151-GAG AGAAA-158) at the loop of the first hairpin of C1 (TL1) that significantly change the reactivity to NMIA in the presence or absence of the 3' UTR (Fig. 3C, red dashed box). This observation shows that upon C1-DB1 interaction, the TL1 nucleotides (which are involved in the C1-PK) become more reactive (Fig. 3C), while the nucleotides of C1-PK do not change the reactivity, since they mostly interact with nucleotides of DB1 (Fig. 3C, circular form).

To further examine whether the predicted RNA-RNA complementarity between C1 and DB1 contributes in stabilizing a complex between the ends of the viral genome, electrophoretic mobility shift assays were performed with different RNA molecules (Fig. 3D). The affinity between RNA molecules containing the sequence of the 3' UTR and the 5' end of the genome including the C1 sequence or not was evaluated. A uniformly 32 P-labeled RNA probe corresponding to the entire DENV2 3' UTR was *in vitro* transcribed and purified in a polyacrylamide gel. The radiolabeled 3' UTR RNA was incubated with increasing concentrations of unlabeled RNAs corresponding to the 5' UTR-FullCap (full length capsid), 5' UTR-FullCap Δ C1 (with a deletion of the C1 structure), or 5' UTR-MCAE (minimal *cis*-acting elements). The complexes were analyzed in native 4% polyacrylamide gels as previously described (11). Total radioactivity for each lane was determined by quantifying the bands corresponding to the RNA-RNA complex and the free probe. For each case, the apparent dissociation constant (K_d) was estimated by nonlinear regression analysis. RNA titrations indicate a significantly higher affinity between the 5' UTR-FullCap and the 3' UTR ($K_d = 16 \pm 5$ nM) than the affinities of the 5' UTR-FullCap Δ C1 ($K_d = 87 \pm 6$ nM) and the 5' UTR-MCAE ($K_d = 200 \pm 14$ nM) with the 3' UTR (Fig. 3D). These results confirm that new sequences in the capsid coding region contribute in stabilizing the RNA-RNA complex formed between the ends of the DENV genome and highlight the relevance of the C1 region.

Next, we evaluated the functional significance of a possible C1-DB1 hybridization during DENV replication. To this end, we constructed recombinant viruses with nucleotide changes disrupting or restoring the predicted interactions. The design of these viruses was very complex because five different structures had to be taken into account simultaneously: C1 structure and local PK, DB1 structure and local PK, and the long-range RNA-RNA interaction (see linear and circular forms of the RNA in Fig. 4A). Mutations were incorporated along with compensations to maintain local structures. Mutant O (MutO) includes substitutions only in the 5' end of the genome that are predicted to disrupt C1-DB1 interaction but maintain the C1 secondary structure. Mutant R (MutR) includes substitutions only in the 3' end that are predicted to debilitate C1-DB1 interaction but maintain the local DB1 and TL1-PK2 pseudoknot structures by compensatory mutations. Finally, a reconstitution mutant (MutO+R) predicted to restore the long-range C1-DB1 interaction contained the substitutions of both MutO and MutR. For clarity, the location of the mutations is indicated in the two alternative structures (linear and circular forms, Fig. 4A). RNAs corresponding to the three mutants (MutO, MutR, and MutO+R) and the RNA of the parental virus (WT) were transfected into C6/36 and BHK cells, and replication

FIG 3 Sequence in the capsid protein coding region facilitates long-range RNA-RNA interactions between the ends of the viral genome. (A) Schematic representation of the conserved patterns of long-range interactions based on alignments of different DENV2 genotypes. Details of nucleotide sequences, alignments, and predictions are described in Fig. S1 in the supplemental material. (B) Representation of the circular conformation of the DENV genome indicating the known and new complementary elements. The predicted RNA-RNA interaction between C1 (red) and DB1 (green) is indicated. (C) Analysis of 5'-end RNA secondary structure in the absence and presence of the 3' UTR RNA (ratio of 1:5). NMIA probing profiles of an RNA corresponding to the 5' end of the genome (5' RNA) and the probing of the same RNA in the presence of a second RNA containing the viral 3' UTR sequence (5' + 3' RNA) are shown on the left. The y axis contains the sequence and nucleotide numbers; the x axis depicts chemical reactivity. On the right, secondary structures and NMIA reactivity are indicated for the predicted 5' end alone and the 5'-3' hybridized form. Red dashed frames highlight a region with different probing profiles. (D) Mobility shift assays indicate that C1-DB1 hybridization stabilizes the RNA-RNA complex. The 3' UTR RNA probe was incubated with increasing concentrations of unlabeled RNAs corresponding to molecules with viral 5'-end sequences as indicated. Mobilities of the 3' UTR RNA probe and RNA-RNA complex as well as the apparent dissociation constants ($K_d \pm$ standard error) are indicated. On the right, the percentage of RNA probe bound was plotted as a function of unlabeled RNA concentration for the three molecules used.

was monitored by luciferase activity. The levels of luciferase at 4 h posttransfection were comparable for all the viruses in the two cell lines, indicating efficient translation of the input genomes. The reporter MutO showed delayed RNA replication with an 8-fold

reduction of luciferase activity at 48 h in BHK cells, while in mosquito cells, replication was reduced about 50-fold (Fig. 4B). The MutR replicated slightly less than the WT virus in both cell lines. However, when the C1-DB1 interaction was reconstituted

FIG 4 Relevance of C1-DB1 interaction in DENV RNA replication. (A) Design of mutation disrupting and reconstituting long-range C1-DB1 RNA-RNA interaction in the context of the viral genome. *cis*-acting elements in the 5' end and the 3' end of the genome are indicated. Nucleotide sequences of C1 and DB1 and the substitutions incorporated are shown in the two alternative conformations, linear and circular. Location, nucleotide changes, and names of mutants are indicated. MutO (green) shows substitutions in C1, and MutR (orange) shows substitutions in DB1. MutO+R contains both substitutions, restoring the interaction and maintaining C1 and DB1 structures. In addition, this mutant maintains the pseudoknot TL1/PK2 by compensatory mutations. (B) Replication of the three mutant RNAs in BHK and C6/36 cells together with the WT control. Luciferase activity was measured as a function of time and represented as arbitrary units (a.u.). The luciferase values are means $\pm$ standard deviations ($n = 3$).

(MutO+R), the impaired replication of MutO was partially restored in both cell lines (Fig. 4B). The restitution of viral RNA replication of MutO by compensatory mutations supports the relevance of the identified long-range interaction. In addition, the data highlight a different requirement of the identified RNA elements for efficient viral replication in mosquito and mammalian cells.

Here, we identified sequences in the DENV genome that enhance viral RNA replication by stabilizing long-range RNA-RNA interactions, in agreement with the proposed function of genome cyclization during minus-strand RNA synthesis (9). The identified complementary sequences are located within the capsid coding region at the 5' end of the RNA and in a well-characterized dumbbell structure at the 3' UTR, and both form PK structures,

stabilizing local RNA elements at each end of the genome. Long-distance hybridization of these RNA sequences disassembles the hairpin and dumbbell structures. This observation, together with previous studies, supports the idea that conservation of alternative (mutually exclusive) functional RNA structures is a common feature in the flavivirus genomes. For instance, in the case of the conserved 5'-3' CS interaction, the 3' CS lies in a region that forms a local PK with the loop sequence of the DB2 structure (located downstream of DB1) (14, 15). In the case of the 5'-3' UAR complementary sequence, the 3' UAR adopts local structures that are essential for viral replication, including the sHP and the base of the terminal 3' SL (16, 21, 37–39). It is possible that conservation of competing structures in the viral genome provides a mechanism for fine tuning RNA conformations required for different viral processes. In this regard, the presence of regulatory RNA elements in protein coding sequences may be relevant for controlling functional RNA structures by translating ribosomes. In the infected cell, it is likely that RNA binding proteins modulate different viral RNA conformations. Many proteins have been previously reported as binders of the DENV RNA (40). Among them, proteins with RNA helicase and chaperone activities have been described to interact with the UTRs of flavivirus genomes (41–43). In this regard, the host helicase DDX6 was reported to bind specifically to the DENV DB1 and DB2 structures; however, the function of these interactions is still unknown (44). Another example is the host protein AUF1, which was shown to bind the WNV RNA and enhance RNA replication likely by rearranging RNA conformations (45). Moreover, the different requirements of viral RNA structures for viral RNA replication in mosquito and mammalian cells may also reflect distinct ways of controlling viral RNA conformations in different host cell environments.

A previous study has investigated the role of RNA sequences within the capsid coding region in DENV infectious particle production. These studies suggested a function of RNA elements during viral assembly in mosquito cells (46). The role of C1 in enhancing viral RNA synthesis observed in our studies limited the possibility of evaluating its function in viral assembly or encapsidation, steps that take place downstream of genome amplification. Nevertheless, we cannot rule out the possibility that C1 sequences play dual functions in viral replication and encapsidation. This possibility should be further investigated.

We propose that DENV conserves RNA sequences that participate in alternative local and long-range structures as a mechanism to modulate the viral RNA architecture for efficient replication. In addition, based on the impact of deletions and mutations of the identified viral RNA structures for replication in mosquito and mammalian cells, we propose that distinct RNA conformations may play different roles in the two host cell environments.

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