A glycerophospholipid-specific pocket in the RVFV class II fusion protein drives target membrane insertion

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The Rift Valley fever virus (RVFV) is transmitted by infected mosquitoes, causing severe disease in humans and livestock across Africa. We determined the x-ray structure of the RVFV class II fusion protein Gc in its postfusion form and in complex with a glycerophospholipid (GPL) bound in a conserved cavity next to the fusion loop. Site-directed mutagenesis and molecular dynamics simulations further revealed a built-in motif allowing en bloc insertion of the fusion loop into membranes, making few nonpolar side-chain interactions with the aliphatic moiety and multiple polar interactions with lipid head groups upon membrane restructuring. The GPL head-group recognition pocket is conserved in the fusion proteins of other arthropod-borne viruses, such as Zika and chikungunya viruses, which have recently caused major epidemics worldwide.

The Rift Valley fever virus (RVFV), a member of the genus Phlebovirus within the order Bunyavirales (https://talk.ictvonline.org/taxonomy/), is a mosquito-borne virus first isolated in Kenya in 1930 (1). RVFV is responsible for the occurrence of epizootones throughout Africa, with devastating economic consequences for the livestock industry (2), and can cause serious disease in humans (3). In 2000, a large-scale RVFV outbreak in the Arabian Peninsula showed that the virus can potentially invade other continents (4). The fact that RVFV can be transmitted to humans by several different mosquito species, a number of which are spread across the planet, is a source of considerable concern (5).

RVFV virions display two envelope proteins, Gn and Gc, at their surface; these proteins are associated as heterodimers arranged with icosahedral symmetry (6–8). The virus enters cells via receptor-mediated endocytosis (9), with the glycoprotein Gc catalyzing an acid-induced membrane fusion reaction with the endosome for entry. The x-ray structure of RVFV Gc in its prefusion form (10) shows the characteristic fold of class II membrane fusion proteins, initially identified in other arboviruses such as those in the genus Flavivirus (11) (which includes Zika, dengue, and yellow fever viruses) and in the genus Alphavirus (12) (which includes the pathogenic chikungunya virus). Class II membrane fusion proteins encompass homologous proteins with a characteristic three-domain β-sheet–rich fold, lacking the typical central α-helical coiled coil of class I and III fusion proteins (13). Despite their unrelated molecular architectures, all three classes appear to use the same overall fusogenic mechanism, with an initial conformational change allowing exposure of a nonpolar segment—termed “fusion loop” or “fusion peptide”—that inserts into the target membrane while the protein adopts an extended intermediate conformation bridging the two membranes. This intermediate then collapses into a “hairpin” that brings the membranes into close apposition (14). The initial insertion into the target membrane is a crucial step and is best understood for the class I fusion proteins, where it involves a structural reorganization of the fusion peptide to expose a highly hydrophobic platform projecting multiple bulky nonpolar side chains (9 to 20, depending on the virus) inserting into the outer lipid leaflet (15, 16). Efficient insertion also requires specific lipids (17). In contrast, the class II and III fusion loops are integral parts of the folded protein and undergo no major reorganization during the fusogenic conformational change (18–22). The energetics of insertion are not understood, as the fusion loops expose only two bulky nonpolar side chains for insertion into the membrane, which is not sufficient to provide strong enough membrane anchoring to withstand downstream forces throughout the fusion process. A requirement for specific lipids has also been documented for class II proteins (23–25), and a direct interaction with cholesterol was demonstrated by photochemical cross-linking in the case of the alphaviruses (26). Furthermore, changes in lipid requirements for insertion correlate with potential alphavirus mosquito vector usage (27), as a variant with a single A226–V226 (A226V) mutation in the fusion protein affects interactions with lipids, which in turn allowed transmission by a different mosquito species to cause a major outbreak of chikungunya disease in Indian Ocean territories in 2005–2006 (28).

We report here structural, biochemical, and molecular dynamics (MD) studies showing that RVFV Gc has a binding pocket for binding glycerophospholipid (GPL) head groups tucked underneath the fusion loop, resulting in a multitude of polar interactions for binding with high affinity, despite insertion of only two bulky nonpolar side chains per subunit. We also show that additional small–head-group lipids (in this case, cholesterol) are required to create space between bulky head groups to allow fusion loop insertion. Finally, we show that a similar pocket is present in the flavivirus and the alphavirus class II fusion proteins and that the residue controlling the selection of the bound GPL corresponds to the site of the A226V mutation in chikungunya virus mentioned above.

We crystallized the recombinant ectodomain of RVFV Gc (29) and determined the structure by molecular replacement (table S1), showing a trimer with the typical organization of its three domains in the postfusion form (Fig. 1), as demonstrated recently for a related tick-borne phlebovirus, the severe fever with thrombocytopenia syndrome virus (SFTSV) (20). The new feature provided by the RVFV Gc structure is the C-terminal region downstream domain III (the “stem”) (Fig. 1, A and B, and fig. S1), which was not resolved in the structures of postfusion class II fusion proteins of arboviruses reported to date. This segment adopted an extended conformation, making two additional β strands running on the side of the elongated tip of domain II, to reach the fusion loop and complete the Gc postfusion hairpin (Fig. 1B). The observed path of the stem is very similar to that of its counterpart in the somatic cell class II fusion protein EFF-1 from Caenorhabditis elegans (30). Comparison with the prefusion form (10) showed that domains I and II remain as a single rigid body with no hinge in between (fig. S2), a notable difference with the alphavirus (19) and flavivirus (18, 21) fusion proteins. All structural changes are confined to the end of domain I opposite to domain II.
where both domain I β sheets are substantially reorganized due to the insertion of an additional β strand, J0 (fig. S2D and movies S1 to S3), entailing changes in the hydrophobic core of the β sandwich. RVFV Gc crystals grew in a solution containing MES [2-(N-morpholino)-ethanesulfonic acid] buffer, and an MES molecule was found bound in a pocket lined by strictly conserved residues adjacent to the fusion loop (fig. S3). Because MES mimics a zwitterionic lipid head group, we tested cocrytalization in the presence of zwitterionic lipids with short aliphatic tails for increased solubility. These experiments led to crystals of a different space group.

Fig. 1. Structure of the postfusion trimer of RVFV Gc and the lipid–head-group binding pocket. (A) Organization of the RVFV middle (M) genomic segment. Oblique stripes mark the noncoding 3′ and 5′ ends, with the single open-reading frame in between. The signal sequences for the NSm, Gn, and Gc proteins are shown in light blue, and predicted transmembrane regions are in dark gray. The first residue of each mature protein is indicated above. The second row outlines the domain organization of Gc, colored according to domains, with domain I red, the two segments that make up domain II distinguished in yellow and beige, domain III blue, and the stem magenta. Regions not present in the structure are in gray [membrane-proximal region (MPR) and cytosolic tail] or black [transmembrane (TM) segment]. This color scheme is maintained throughout the paper. vRNA, viral RNA. (B) Gc postfusion trimer in surface representation, except for the protomer in the foreground, which is represented as ribbons colored by domains. Disulfide bonds are shown as green sticks and are numbered. The bound C3PC lipid (framed) and the N-linked glycan chains are drawn as sticks colored according to atom type (carbon, white; nitrogen, blue; oxygen, red; phosphorus, orange). Lowercase letters denote β strands in domain II, following the standard nomenclature for class II fusion proteins. (C) Comparison of the chemical structures of MES and C3PC. (D) C3PC binding groove. The Gc trimer is shown in surface representation and colored according to electrostatic potential, with C3PC shown as sticks. Relevant residues are labeled. (E) Electron micrograph of liposomes after incubation with Gc and negative staining with uranyl acetate. Scale bar, 100 nm. (F) Binding of Gc to DOPC membranes requires cholesterol. Sensorgrams measured by bio-layer interferometry (BLI), showing the interaction of Gc with liposomes containing different amounts of DOPC and cholesterol (CHL) as indicated.

Fig. 2. Lipid binding specificity. Comparative binding of WT Gc and mutants at position 961 (i/loop) to liposomes made of DOPC:DOPS:cholesterol at molar ratios 4:0:4 (left), 3:1:4 (middle), and 2:2:4 (right) (i.e., 50% cholesterol and increasing DOPS concentration). (A and B) Surface plasmon resonance (SPR) (A) and BLI (B) sensorgrams measured with the WT Gc ectodomain (magenta) and with mutants D961N (green), D961K (light blue), and W821H (dark blue, used as control). All experiments were performed as duplicates (shown by double lines). (C) BLI sensorgrams showing the interaction of Gc WT, D961N, and D961K with liposomes made of 30% cholesterol and different compositions of SM, DOPC, and DOPE as indicated [note that this color key corresponds only to (C)]. In the SPR experiments (A), the liposomes were immobilized on the chip surface and the protein flown on top, whereas in the BLI experiments [(B) and (C)], the protein was immobilized in the tip and the liposomes were in solution.
diffracting to higher resolution (table S1) in the presence of the GPL dipropionyl phosphatidylcholine (C$_3$PC). The crystal structure revealed a C$_3$PC molecule (Fig. 1D) bound in the same way as MES (fig. S3), with the two nonesterized oxygen atoms of the phosphate group engaging in bidentate hydrogen bonds with the R776 side chain at the very end of the $\beta$ strand $\beta$ at the tip of domain II (see Fig. 1B and fig. S1 for class II secondary structure nomenclature). The glycerol moiety fit between the disulfide bond 5, which is conserved across all class II fusion proteins, and the side chain of V780 at the end of $\beta$ strand $\beta$. The two propionyl aliphatic tails projected out, at the very top, packing against the side chain of F826 (Fig. 1D and fig. S3C), the only nonpolar side chain of the fusion loop (cd loop) that is strictly conserved across the Phlebovirus genus. In line with this observation, the F826N mutation in RVFV Gc abolishes fusion in a cell-to-cell membrane fusion assay (31), and the F699S mutation of its counterpart in SFTSV (fig. S4A) results in nonviable virus (20). The presence of the bound lipid did not alter the conformation of the fusion loop main chain compared to its prefusion form (fig. S4A). Notably, the fusion loop of the flavivirus fusion protein was also in the same conformation in both its pre- and postfusion forms (18, 21, 32), unlike the alphavirus fusion protein, which has a longer fusion loop found in alternative conformations (18, 19, 32).

Liposome coflotation experiments in density gradients, together with electron microscopy, showed that the wild-type Gc ectodomain binds to phosphatidylcholine (PC)-containing liposomes, with the Gc ectodomains bound to membranes via the tapered end (i.e., via the domain II tip) and making clusters at the liposome surface (Fig. 1E and fig. S4B), as shown earlier for other class II fusion proteins (18, 33). To further understand the lipid requirements for binding, we monitored the interaction with liposomes containing PC as the main component and varying amounts of other lipids. The Gc ectodomain bound to dioleoyl phosphatidylcholine (DOPC) liposomes at neutral and acidic pH (fig. S4C) only if the liposomes had at least 20% cholesterol. Binding increased to be highest at the maximum cholesterol/PC ratio tested (50%) (Fig. 1F). In the structure, the positively charged interactions, with the residues of the cd loop represented as sticks. Carbon, nitrogen, and oxygen in white, blue, and red, respectively. Hydrogen bonds are shown as dashed lines. (E) MD simulation statistics as function of DOPC/cholesterol ratio. (Top panel) Membrane penetration depth ($D_d$; see materials and methods) relative to 0% cholesterol. (Middle panels) Total protein-lipid potential energy $V$ and number of protein-lipid hydrogen bonds, respectively. (Bottom panel) Cholesterol enrichment at the membrane-protein contact area, computed as described in (29). Circles represent average values from 2 $\mu$s of simulation collected from several independent runs, with error bars (SEM) calculated by block averaging. The average values from individual simulation runs are shown as triangles. For a full description, see (29).
In the MD simulations, DOPC head groups consistently found their way into the GPL pocket (which was left empty by removing the bound C3PC molecule at the beginning of the calculations), demonstrating that the lipid-binding mode revealed in the crystal structure is also used in a membrane environment. The simulations revealed long-lived strong hydrogen bonds between the DOPC-phosphate group and the R776 guanidinium trimethylamine group of the zwitterionic PC head group faces the strictly conserved D961. Thus, we explored the effect of neutralizing and reversing the electrostatic charge at this location on liposome-binding properties by making mutants D961N and D961K. The WT Gc ectodomain bound consistently better in the absence of dioleoyl phosphatidyl serine (DOPS), a GPL with a negatively charged head group (Fig. 2), whereas the D961N mutant bound to a similar extent with or without DOPS. The D961K mutant had the reverse effect compared with the wild type: The mutant showed almost no binding in the absence of DOPS, and although it remained poor, binding increased with increasing DOPS, as predicted from the positive charge at position 961 in the ij loop. Thus, the residue at the tip of the ij loop affects the membrane-binding properties and is important to specify the Gc being recognized.

We further analyzed lipid specificities by replacing PC with sphingomyelin (SM), which also has a phosphocholine head group but is connected to a ceramide instead of a diacetylcerol backbone (fig. S5A). The Gc ectodomain did not bind to 100% SM liposomes, and the addition of cholesterol at various ratios did not rescue binding (fig. S5B). Because liposomes containing SM and cholesterol did not support Gc insertion, we also tried replacing PC with phosphatidylethanolamine (PE), as PE does not make lipid bilayers by itself and cannot be tested on its own. Adding PE efficiently restored binding (Fig. 2C and fig. S5C), which led to increased binding with respect to the liposomes containing PC and cholesterol. The effect of the mutants at position 961 was the same as for the PC-containing liposomes: The neutral D961N mutant bound less well, and the D961K mutant did not bind appreciably (fig. S5C).

To obtain a molecular model for Gc-membrane interactions, we ran MD simulations of the Gc trimer in contact with membranes composed of DOPC and cholesterol at different ratios (fig. S6) (29). In agreement with the liposome-binding experiments, the simulations showed that, with increasing cholesterol content, the protein reached deeper into the membrane and formed more favorable contacts (Fig. 3), as quantified by a decrease in protein-lipid potential energies and an increase in the number of protein-lipid hydrogen bonds (Fig. 3E and movie S5). The enhanced protein-lipid interactions correlated with an enrichment of cholesterol at the protein-membrane interface (Fig. 3, A and E (bottom panel), and fig. S7A). Cholesterol promoted binding mainly by creating space between the bulky DOPC head groups (Fig. 3, B and C), allowing the Gc fusion loop to penetrate the head-group region (Fig. 3D). In addition, multiple hydrogen bonds with cholesterol were formed by the main chain of the fusion loop and with side chains of inserted residues such as W821, N827, and N829 (Fig. 3D), in line with the reported lack of fusion activity of the mutant N827A (31).
group, whereas the interaction of the DOPC-trimethylamine group with the D961 side chains was more dynamic. Indeed, the methyl groups partially shield the positive charge of the central amine, resulting in frequent on-off interactions (fig. S7B), in line with the observed stronger binding to PE-containing membranes (Fig. 2C, left panel). Finally, to rationalize the experimentally determined lack of binding of the W921H mutant and the W921A/F826A double mutant (Fig. 2 and fig. S4D), we computed the change in membrane binding affinity ΔΔG caused by the mutations (Fig. S6B). We obtained ΔΔG values of 45.9 and 60.2 kJ mol⁻¹ for the single- and double-mutant trimers, respectively, confirming the crucial role of these fusion loop aromatic side chains for binding. For the class I fusion protein hemagglutinin (HA) of influenza virus, the membrane binding energy of the fusion peptide was measured to be roughly 32 kJ mol⁻¹ (34), which implies 96 kJ mol⁻¹ per HA trimmer. The HA fusion peptide inserts nine nonpolar side chains into the membrane (35), a considerably more hydrophobic surface, with a total of 27 hydrophobic residues per trimer. Although the obtained ΔΔG figures are relative (wild type versus mutant) and not absolute binding affinities—and thus cannot be directly compared with the HA case—they still reveal that the effect of inserting only two aromatic side chains per RVFV Gc protomer involves a considerable difference in binding energy, which appears to be within a factor of 2, at most, from the absolute value measured for the HA trimer.

Inspection of the structures of the flavivirus E and alphavirus E1 fusion proteins revealed the presence of a groove at the same location of the GPL pocket of RVFV Gc (Fig. 4A, middle and right panels, respectively). The disulfide bond that interacts with the glycerol moiety is conserved, and a positively charged arginine or lysine side chain at the end of β-strand b is also in position to interact with the phosphate group. Moreover, the ij loop is also poised to provide the relevant contact that would specify the particular GPL being recognized. In line with the reported dengue virus requirement for anionic GPLs such as phosphatidyl serine in the target membrane for fusion (36), the ij loop in protein E projects a lysine side chain into the pocket (Fig. 4A, middle panel). To test the GPL requirement for binding, we ran liposome coexclusion experiments (fig. S8) using the ectodomain of flavivirus E protein (from Zika virus; fig. S8A) and the alphavirus E1 (from chikungunya virus; fig. S8B), confirming that in the absence of GPLs, these proteins display no detectable interaction with membranes, even in the presence of cholesterol to provide space between the bulky lipid head groups, similar to the SM and cholesterol results in the case of RVFV Gc (fig. S5B). In contrast, the control herpesvirus class III fusion protein gB interacted with liposomes made of SM as long as cholesterol was present, indicating that it does not rely on

REFERENCES AND NOTES

29. Materials and methods are available as supplemental materials. See https://www.sciencemag.org/content/358/6363/663/suppl/DC1.

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SUPPLEMENTARY MATERIALS

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Materials and Methods

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Unmasking class II membrane fusion
Rift Valley fever virus (RVFV) is transmitted by mosquitos and enters cells through receptormediated endocytosis. The infection process requires class II membrane fusion proteins, which insert a hydrophobic fusion loop into cell membranes and then refold. Guardado-Calvo et al. report the high-resolution crystal structure of RVFV class II fusion protein Gc in its postfusion form complexed with phosphatidylcholine. They find that Gc does not restructure its fusion loop after insertion. Rather, it uses an integrated system that accommodates glycerophospholipid head groups and then initiates membrane reorganization by concentrating cholesterol at the insertion site. Comparison with class I fusion proteins from other virus families suggests a common mechanism, which may provide a target for future antiviral therapies.

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