Structural dissection of Ebola virus and its assembly determinants using cryo-electron tomography

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Ebola virus is a highly pathogenic filovirus causing severe hemorrhagic fever with high mortality rates. It assembles heterogeneous, filamentous, enveloped virus particles containing a negative-sense, single-stranded RNA genome packaged within a helical nucleocapsid (NC). We have used cryo-electron microscopy and tomography to visualize Ebola virus particles, as well as Ebola virus-like particles, in three dimensions in a near-native state. The NC within the virion forms a left-handed helix with an inner nucleoprotein layer decorated with protruding arms composed of VP24 and VP35. A comparison with the closely related Marburg virus shows that the N-terminal region of nucleoprotein defines the inner diameter of the Ebola virus NC, whereas the RNA genome defines its length. Binding of the nucleoprotein to RNA can assemble a loosely coiled NC-like structure; the loose coil can be condensed by binding of the viral matrix protein VP40 to the C terminus of the nucleoprotein, and rigidified by binding of VP24 and VP35 to alternate copies of the nucleoprotein. Four proteins (NP, VP24, VP35, and VP40) are necessary and sufficient to mediate assembly of an NC with structure, symmetry, variability, and flexibility indistinguishable from that in Ebola virus particles released from infected cells. Together these data provide a structural and architectural description of Ebola virus and define the roles of viral proteins in its structure and assembly.

Mononegavirales | single-stranded RNA virus | virus structure | subtomogram averaging

Ebola virus (EBOV) and Marburg virus (MARV) constitute the family *Filoviridae* within the order *Mononegavirales*. Filoviruses are highly pathogenic, causing severe hemorrhagic fever in monkeys and humans, with high mortality rates (1). Because of the lack of approved vaccines and antiviral drugs, both EBOV and MARV are categorized as biosafety level-4 (BSL-4) pathogens.

The order *Mononegavirales* also contains several other pathogens of clinical importance, such as rabies virus (RABV), mumps virus, measles virus (MeV), and respiratory syncytial virus (RSV) (2). All members of the order possess a nonsegmented, negative-sense RNA genome, which is encapsidated by the viral nucleoprotein (NP). The NP–RNA complex acts as the template for genome replication and assembles into a helical nucleocapsid (NC) along with accessory proteins (3). This characteristic links genome replication mechanisms of mononegaviruses to their NC structure. The NC is recruited to the plasma membrane by the viral matrix protein, where it buds through the membrane to form an enveloped virion. All mononegaviruses share these fundamental characteristics.

EBOV virions contain an RNA genome and seven viral proteins: NP, VP35, VP40, GP (glycoprotein), VP30, VP24, and an RNA-dependent RNA polymerase (L). NP, VP30, VP35, and L are known to associate with the transcription and replication-competent NC (4–6). VP24 is additionally required for NC assembly (7, 8). VP40, the viral matrix protein, binds directly to the viral envelope. Expression of VP40 alone in mammalian cells can

lead to formation and release of enveloped, filamentous virus-like particles (VLPs) (9–12). Expression of NP alone leads to the formation of narrow, tubular structures in the cytoplasm of the cell (13). These narrow structures can be recruited into VLPs by coexpression of VP40 (14). If NP is expressed together with VP24 and VP35, cytoplasmic clusters of NC-like structures are formed that are similar to those seen in infected cells (13). These structures are also recruited into VLPs when VP40 is coexpressed (7, 13–15). Together these studies suggest that a direct interaction between VP40 and NP can recruit NP into released VLPs and that formation of an NC with diameter similar to that in native virions requires co-expression of NP, VP24, and VP35.

Recent cryo-electron microscopy (cryoEM) investigations of MARV described the 3D structure of the MARV NC (16). The MARV NC is a left-handed helix, with the viral NP forming the innermost layer of the structure. Each NP binds to six bases of RNA. Arm-like structures protrude from alternate interfaces between NPs, and immuno-electron microscopy analysis locates VP24 and VP35 to these protrusions. The NC is incorporated into virions by envelopment at the plasma membrane initiated at one end of the NC (16, 17).

In the present study, EBOV virions were imaged using cryoEM and cryo-electron tomography (cryoET) to describe their structure in a near-native state. Image-processing techniques were applied to define the 3D structure of the NC within the virion. The EBOV NP shares ~40% sequence homology with MARV NP (18, 19). Comparison of the morphological parameters and NC structures of EBOV with MARV allowed us to dissect the roles played by the RNA genome and filovirus NPs in determining NC structure.

In addition, Ebola VLPs were produced with different combinations of viral proteins and studied using biochemical, cryoEM, and cryoET techniques. These studies define roles for viral proteins in determining the structure of EBOV virions and their NCs, which range from mediating initial coiling of the NC helix, to helical condensation, to rigid helix formation, to NC envelopment into virions.

Results

CryoEM and CryoET of EBOV. Zaire EBOV virions were harvested from infected Vero cells 1 d after infection in a BSL-4 laboratory. The inactivated virus pellet was released from the BSL-4 laboratory and then imaged using cryoEM. Long, filamentous mem-

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brane-bound particles could be observed along with spherical particles and other irregularly shaped vesicles (Fig. 1*A*). Several virions possessed previously described “moth-eaten” membranes (1). Some filamentous particles lacked NC structures, resulting in a smaller diameter (Fig. 1*A* and *B*, black arrow). Many virions displayed an intact membrane and a clearly visible NC (Fig. 1*B*, white arrow). This variable morphology of EBOV is consistent with previous negative staining EM analysis (1).

CryoEM allows excellent preservation of the specimen in a near-native environment: this allowed us to accurately measure morphological parameters of virions. The distribution of virus lengths (Fig. 1*C*, Fig. S1*A*, and *SI Materials and Methods*) showed a major population of virus particles with a length of $1,028 \pm 69$ nm ($n = 37$), consistent with previously reported values of 970–1,200 nm for the average EBOV virion (20, 21). We also found a second population with a mean length of $1,978 \pm 112$ nm ($n = 8$) (Fig. 1*C*), as well as some longer particles (Fig. S1*A*). The diameter of filamentous EBOV particles that had a continuous membrane and an internalized NC was 90 ± 3 nm ($n = 50$) (Fig. 1*C*, *Right*), which is slightly smaller than that of MARV particles (92 ± 4 nm) (Fig. 1*D*, *Right*).

Fig. 1. CryoEM of EBOV. (*A*) Low-magnification cryoEM images of purified EBOV. Protein density is black. Filamentous particles of varying lengths, spherical particles, and other irregularly shaped particles are observed. (*B*) CryoEM image of a filamentous EBOV virion. White arrow, EBOV virion with an NC. Black arrow, a thin particle without an internalized NC. (*C*) Histograms of virion length (*Left*) and diameter (*Right*) for filamentous EBOV virions containing an NC. (*D*) Corresponding histograms for MARV. More details in Fig. S1 and *SI Materials and Methods*.

To understand the 3D arrangement of the virion, we performed cryoET. A slice through a representative tomogram is shown in Fig. 2*A*. The viral NC appears as a cylinder-like density within the particle center (white arrow in Fig. 2*A*, Movie S1), similar in appearance to the MARV NC (16). Regular repeats at a pitch of ≈ 7 nm could be observed along the length of the NC.

To resolve the structure of the EBOV NC in more detail, we applied subtomogram averaging methods on the tomography data (*SI Materials and Methods*), as described previously (16, 22). All of the reconstructed helices were left-handed with an inner layer decorated with arm-like protrusions in the outer layer. Of NCs whose symmetry could be unambiguously assigned, all were found to contain either 11.8 or 12.8 repeating units per turn. Combining all of the NC helices with the same symmetry into one single subtomogram averaging reconstruction enabled refinement up to 4.1 nm resolution. We also performed iterative real-space helical reconstruction using helical segments extracted from 2D cryoEM images of EBOV (Fig. S2, *Left*) and again obtained a reconstruction with a resolution of 4.1 nm. In contrast, subtomogram averaging and iterative real-space helical reconstruction resolved the MARV NC to better resolutions of 3.4 nm and 2.5 nm, respectively (16). This suggests that the EBOV NC has a higher amount of conformational variability or flexibility than the MARV NC.

We therefore identified the subset of NC helices that aligned successfully (*SI Materials and Methods*) and had a symmetry of 11.8 subunits per turn and combined them into one final reconstruction (Fig. 2*B* and *C*, *Left*) with an improved resolution of 3.6 nm. The final subtomogram averaging reconstruction shows the EBOV NC helix to be left-handed with a pitch of ≈ 7.4 nm (Fig. 2*B* and *C*, *Left*) close to that of MARV (7.5 nm) (16). An inner layer is observed with a diameter of ≈ 28 nm. Boomerang-shaped densities protrude outward from this inner layer, and the diameter of the entire structure is ≈ 40 nm. The protrusions have two lobes. A left-handed helix with a pitch of

Fig. 2. CryoET and 3D reconstruction of the EBOV NC from subtomogram averaging. (*A*) A slice through a reconstructed, filtered tomogram of EBOV. Protein density is black. White arrowhead indicates the rod-like NC within the virion. (*B*) Reconstruction of the EBOV NC from cryoET and subtomogram averaging (*Left*) compared with the MARV NC reconstruction (*Right*) (16). Iso-surfaces have been contoured at 1.5σ away from the mean, and the helical axis is vertical in the plane of the paper. (*C*) The same reconstructions as *B*, viewed along the helical axis.

Order of Protein Assembly and Formation of a Rigid NC Helix. We next prepared a series of VLPs by expression of different combinations of viral proteins along with VP40 in mammalian cells: NP+VP40, NP+VP24+VP40, NP+VP35+VP40, and NP+VP24+VP35+VP40. Previous thin-section EM analyses have indicated the presence of an NC-like structure in these VLPs (13–15). We analyzed recruitment of viral proteins into released VLPs using Western blot analysis with anti-NP, anti-VP24, anti-VP35, and anti-VP40 sera. We found that NP was recruited into VLPs by coexpression with VP40 alone (Fig. 4A). When NP, VP35, and VP40 were coexpressed, all three proteins could be detected in VLPs. These observations are consistent with previous observations that VP40 can recruit VP35 and NP independently into VLPs (13, 26). We found that only a low amount of VP24 was recruited into VLPs when it was coexpressed with NP and VP40. However, VP24 was detected in large amounts when VP35 was additionally expressed (Fig. 4A). These results indicate that NP can be directly recruited into VLPs by VP40, that VP24 and VP35 can be recruited by NP and/or VP40, and that VP35 significantly enhances the recruitment of VP24 into VLPs.

All filamentous VLPs were subjected to cryoEM and cryoET. CryoEM was used to quantify the number of VLPs with and without an internalized NC. CryoET was used to divide VLPs that contained NCs into two structural classes by a visual inspection of the filtered tomograms. The first class had an NC with short stretches of condensed helix broken at multiple points (Fig. 4C). The second class contained a rigid, largely continuous NC structure with outer protrusions (Fig. 4D). We compared the frequencies of the different classes of NCs found in the VLP samples from tomograms (Table S2) with those in EBOV virions. Together the cryoEM and cryoET data showed that in NP+VP40 VLPs, 64% of the VLPs were empty (Fig. 4E), and 36% contained broken,

discontinuous NCs. A rigid, continuous NC could not be observed in any of the NP+VP40 particles. A very similar pattern was found in NP+VP24+VP40 VLPs (73% empty, 27% broken) and in NP+VP35+VP40 VLPs (66% empty, 34% broken) (Fig. 4E).

Although the percentage of empty particles (68%) in NP+VP24+VP35+VP40 VLPs was similar to the other analyzed VLP samples, the NC, when present, was predominantly rigid: 30% of VLPs contained a rigid NC structure, whereas only 2% contained a broken or discontinuous NC. These numbers are comparable to our observations of authentic virions, in which we found that 63% of the particles were empty and 34% contained continuous rigid NCs (Fig. 4E). Unlike authentic virions, the length of the NP+VP24+VP35+VP40 VLPs was not well defined (Fig. S1C). To summarize, in the absence of NP, VP24, or VP35, a rigid NC-like structure was never observed. When NP, VP24, VP35, and VP40 were coexpressed, VLPs were obtained with rigid NCs that were morphologically similar to the full EBOV NC. These data indicate that NP, VP24, and VP35 are all required to form a rigid, continuous NC structure.

Structural Characterization of the VLPs. To detect differences in the NCs between various VLP samples, we next performed 2D classification and averaging of helical segments extracted from cryoEM images of the VLPs (SI Materials and Methods). The NP+VP40 VLPs contained an NC helix with a diameter of ≈ 28 nm, lacking the arm-like protrusions observed in authentic virions (Fig. 5A). Because of discontinuities in the NC helix, the NP layer in the average image appears blurred. Average NC images from NP+VP24+VP40 VLPs and NP+VP35+VP40 VLPs had the same appearance (Fig. S3).

In contrast the NP+VP24+VP35+VP40 VLPs show a rigid NC helix (Fig. 5B) with protrusions emanating from the inner NP layer (Fig. 5B, arrows), appearing similar to NC from authentic virions (Fig. 5C). We performed 3D reconstruction of NCs from the NP+VP24+VP35+VP40 sample using subtomogram averaging and real-space helical reconstruction techniques, exactly as described above for the NC within EBOV virions. The NC helix in the VLPs adopted the same symmetries (11.8 and 12.8 protrusions per turn) and structure as the NC helix in virions (Fig. 5D and E), with an inner layer decorated with boomerang-shaped outer protrusions. The resolution of the real-space helical reconstruction was 4.1 nm (Fig. S2, Right), and a selected subset of the NCs combined with subtomogram averaging reached a resolution of 3.9 nm (Fig. 5D and E). In both cases the reconstructions are the same as the NC reconstruction from virions with the same resolution (compare Figs. 2B and C with 5D and E and Fig. S24, Left vs. Right). Thus,

Fig. 4. Protein recruitment and formation of a rigid NC. (A) Detection of viral proteins in respective VLPs. Purified VLPs were collected, and Western blot analysis using rabbit anti-NP, -40, -35, and -24 antibodies was performed. (B) A tomographic slice through an empty VLP. Protein density is black. (C) Slice through a VLP with a broken NC. Points of breakages in the NC helix have been highlighted with white arrows. (D) A VLP with a rigid NC. (E) Proportion of particles observed with a rigid NC (dark green), with an overall broken NC (orange), and without an NC (gray) in different samples. Data values are in Table S2.

Fig. 5. Location of viral proteins in the EBOV NC. (A) 2D class averages of the NC from NP+VP40 VLPs. (B) 2D class averages of the NC from NP+VP24+VP35+VP40 VLPs. (C) 2D class averages of the NC from EBOV virions. Black arrows indicate protrusions. (D) Subtomogram averaging reconstruction of the NC helix from NP+VP24+VP35+VP40 VLPs. Isosurfaces have been contoured at 1.5σ away from the mean, and the helical axis is vertical in the plane of the paper. (E) The same reconstruction viewed along the helical axis.

the NC helices from NP+VP24+VP35+VP40 VLPs and from EBOV virions are indistinguishable in structure, symmetry, and flexibility.

Discussion

Architecture of EBOV Virions and the EBOV NC. We found that EBOV particles were largely filamentous, but other morphologies, including spherical particles and particles without an internalized NC, were also observed. Such variable morphology is consistent with earlier observations by negative staining EM (1). Within cryoEM images we could see that straight sections of virions contain a cylindrical NC along the center of the virus particle. Most filamentous EBOV virions had a length of $\approx 1,028$ nm, although longer viruses were also observed with lengths that were approximate multiples of this length, suggesting they contain multiple NCs.

A comparison of the EBOV with the recently presented cryoET structure of MARV NC (16) sheds light on factors affecting virus assembly. Many features are shared between the two NCs. The pitch of the EBOV NC helix (7.4 nm) is almost identical to that described by cryoEM for the MARV NC (7.5 nm). The 3D structure of the EBOV NC reveals a left-handed helical structure, just like the MARV NC (16). As in MARV it shows an inner layer made up of the viral NP, which is decorated by boomerang-shaped protrusions. By analogy with MARV, one protrusion emanates from every two NPs in the inner layer. Binding of one copy of the viral phosphoprotein to two copies of the NP has also been observed by x-ray crystallography of a purified rhabdoviral complex (27).

There are also differences between the EBOV and MARV NCs. In all our analyses the EBOV NC was consistently more flexible than the MARV NC. This suggests higher intrinsic conformational flexibility in the repeating asymmetric unit of the EBOV NC. The symmetry of the two filovirus NCs differs: the EBOV NC has fewer NP subunits per turn of the helix but has more turns of the helix per virion, so that EBOV virions are longer than MARV virions. This means that the total number of NPs is approximately the same in EBOV and MARV, and the number of RNA bases per copy of NP is also the same, with each NP binding six RNA bases.

Genome replication in *Mononegavirales* is tightly linked to NC structure (28). A density of six bases per NP in EBOV is consistent with previous observations that only multiples of six bases can be added or removed from the replication promoter region while maintaining function (29). Binding to a multiple of six RNA bases per NP monomer is also observed in paramyxoviruses like Sendai virus and MeV (30, 31), and like EBOV (29) these viruses also have bipartite replication promoters. These facts together suggest that genome replication mechanisms of filoviruses are likely similar to those of Sendai virus and MeV and differ from RSV and rhabdoviruses like VSV and RABV, which package different numbers of RNA bases per NP (32–34).

Structural Roles of EBOV Components in Determining NC Structure.

The expression of VP40 along with NP leads to recruitment of NP into VLPs. This is likely due to binding of VP40 to the C terminus of NP (25). Expression of NP and VP40 together allows recruitment of VP35 into VLPs. For efficient VP24 recruitment into VLPs, NP, VP35, and VP40 must be expressed. This is consistent with previous morphological studies that asserted that NP, VP24, and VP35 are all necessary for NC assembly (7, 13).

The EBOV NP alone, upon binding to RNA, forms a loosely coiled helix. Removal of the C-terminal 288 residues of NP, which are predicted to contain large disordered regions, leads to formation of condensed helices instead of loose coils. The C-terminal region of NP therefore prevents condensation of the N-terminal region of NP into helices. CryoEM observations on MARV and MeV NPs have also shown that purified NP samples could form loose helices, and that C-terminally deleted NPs could assemble condensed helices (16, 23), suggesting that this is a general property shared with other mononegaviruses. The disordered C-terminal domain of EBOV NP contains binding

sites for VP40. NCs purified from VLPs produced by coexpression of NP and VP40 are condensed helices indistinguishable from those formed by C-terminally deleted NP. We therefore propose that binding of VP40 to the C-terminal region of NP during virus assembly relieves its inhibitory effect, allowing the N terminus of NP to assemble a condensed helix. The condensed EBOV NP(1-451) helices have a diameter of ≈ 28 nm, which is the same as the 28-nm diameter of the inner NC helix in the authentic EBOV particle. MARV NP(1-390) assembles into a condensed helix with a diameter of 33 nm, which is the same as the 33-nm diameter of the inner NC helix in the authentic MARV particle (16). This comparison suggests that the N-terminal domain of NP in filoviruses is alone sufficient to define the diameter of the NC helix.

The condensed NC helix retains some flexibility and is punctuated by breaks when packaged into VLPs, or by regions of loose coil when purified. This contrasts with the viral NC, which we found to form rigid helices. Coexpression of both VP24 and VP35 with NP and VP40 was required to release VLPs containing rigid NCs, suggesting that binding of VP24 and VP35 leads to rigidification of the helix. These two proteins form boomerang-shaped protrusions emanating from the inner NP layer. NP, VP24, VP35, and VP40 together are sufficient to assemble an NC that has the same symmetry, structure, variability, and flexibility as the NC within the virion.

The NCs in NP+VP24+VP35+VP40 VLPs do not have a defined length, contrasting with the NC in authentic EBOV and MARV virions that consistently has exactly the length required to package one viral genome at a density of six RNA bases per NP. In some cases more than one NC can be incorporated into a single virion, giving a virus particle with double

Fig. 6. Steps involved in EBOV NC assembly. A schematic illustration of the samples described in this study and their assembly properties. Assembly of a virus particle is indicated by the thick arrow. Initial condensation of the NP-RNA complex can be achieved in vitro by removal of the disordered C-terminal, or in cells by coexpression with VP40 (thin arrows). The condensed helix can be converted into a rigid NC-like helix inside VLPs only if all NP, VP24, VP35, and VP40 are expressed. The resulting NC helix is indistinguishable from that in EBOV virions.

or triple the expected length. These observations imply that NC length is determined by genome length.

By analysis of purified proteins, VLPs, and virions, we can propose distinct structural roles for these components in EBOV assembly. We suggest that formation of a virus particle requires packaging of the RNA genome by N-termini of NP to form a loose coil with a length defined by total genome length. We suggest that it requires condensation of the loose coils into a helix with diameter defined by the N-terminal region of NP and that this can be mediated by binding of VP40 to the C terminus of NP. We suggest that it requires rigidification of the condensed coils into a tight helix with arm-like protrusions by binding of VP24 and VP35 to alternate NPs and that these components are sufficient to define the mature EBOV NC structure (Fig. 6).

Materials and Methods

Purification of Recombinant EBOV NP. NP or its (1–451) truncation mutant were purified from transfected HEK 293 cells using CsCl gradient centrifugation (*SI Materials and Methods*). All samples were prepared in duplicate to control for differences between sample preparations.

Preparation of VLPs and Virus. EBOV proteins were coexpressed with VP40 in HEK 293 cells. Two days after transfection, VLPs were fixed with 1% paraformaldehyde (PFA) and pelleted by ultracentrifugation through a 20% (wt/wt) sucrose cushion. The pellet was resuspended in PBS and stored at 4 °C until further investigation.

All work with infectious EBOV was performed under highest safety precautions in the BSL-4 facility at the Institut für Virologie, Philipps-Universität Marburg. Particles of EBOV that were released from infected Vero cells were collected 1 d after infection, purified by centrifugation through a 20% sucrose cushion, and fixed with 4% PFA to inactivate the virus completely (*SI Materials*

and Methods). All samples were prepared in duplicate and initially analyzed separately to control for differences between preparations.

Western Blot Analysis. Purified VLPs were lysed in SDS sample buffer and separated on a PAGE Tris/glycine gel. Blots were incubated with rabbit anti-NP, anti-VP24, anti-VP35, or anti-VP40 serum as primary antibodies, and with HRP-conjugated anti-rabbit IgG antibody as a secondary antibody. Bands were detected with ECL Plus Western Blotting Detection Reagents (GE Healthcare) and visualized using VersaDoc Imaging System (Bio-Rad).

CryoEM and Image Analysis. For cryoEM studies, vitrified samples were imaged under standard low-dose conditions in a FEI CM120 Biotwin microscope (120 kV). For tomography an FEI TF30 Polara TEM (300 kV) with energy filter was used. Tomographic tilt ranges were typically from +60° to –60°, with a total dose of 6,000–10,000 e[–]/nm². For each VLP sample, 8–20 tomograms were collected, and for virions more than 20 tomograms were collected.

2D data were analyzed using Bsoft (35) and Spider (36). Helical reconstruction was carried out using the real space reconstruction technique (37) implemented in the Spider package (*SI Materials and Methods*). Tomograms were reconstructed using the IMOD software suite (38). Subtomograms were extracted along the length of NCs and iteratively aligned in six dimensions, taking into account the missing wedge as described previously (16, 39). Visualization of image data was carried out in Amira (Visage Imaging) and Chimera (40).

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