Formation of Higher-Order Foot-and-Mouth Disease Virus 3Dpol Complexes Is Dependent on Elongation Activity

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The replication of many viruses involves the formation of higher-order structures or replication “factories.” We show that the key replication enzyme of foot-and-mouth disease virus (FMDV), the RNA-dependent RNA polymerase, forms fibrils in vitro. Although there are similarities with previously characterized poliovirus polymerase fibrils, FMDV fibrils are narrower, are composed of both protein and RNA, and, importantly, are seen only when all components of an elongation assay are present. Furthermore, an inhibitory RNA aptamer prevents fibril formation.

Foot-and-mouth disease virus (FMDV) is one of the most important animal pathogens of agricultural significance, yet key aspects of its replication are still unclear. The virus occurs as seven distinct serotypes and is a member of the Picornaviridae, which also includes important human pathogens, such as poliovirus (PV) and rhinovirus. Replication of the positive-stranded RNA genome occurs via a negative-stranded intermediate and involves several viral proteins as well as specific sequences within the untranslated regions (UTRs) at the 5’ and 3’ ends. The 5’ UTR is exceptionally long (~1,200 nucleotides), constituting one-seventh of the viral genome. It includes a 400-nucleotide predicted to be double stranded in structure (3), a poly(C) tract of ~200 nucleotides (20), a series of pseudoknots (3, 6), a cis-acting replication element (cre) necessary for the uridylation of the peptide primer of replication (VPg) (13, 15), and an internal ribosome entry site (IRES) necessary for the initiation of protein synthesis (18). The 3’ UTR is shorter but includes two stem-loop structures and a poly(A) tract. Several of these RNA structural elements are known or thought to be important in genome replication. The engine of genome replication is an RNA-dependent RNA polymerase enzyme (RdRp), termed 3Dpol, with the involvement of its precursor, 3CD. RNA synthesis is primed by a uridylation of VPg, which is the product of the 3B gene. Uniquely, the FMDV genome encodes three tandem copies of 3B (VPg), and it is tempting to speculate that this feature is causally linked to the exceptionally fast replication rate characteristic of the virus. Some strains of FMDV complete the entire life cycle in only 3 h in vitro (20). Uridylation of VPg to form VPgpUpU is carried out by 3Dpol [3Dpol(C)] was purified as described previously (7). The 3Dpol (serotype O) gene was subcloned from pUb-3Dpol (15) into pET22b-3Dpol. The protein was expressed in Escherichia coli as a C-terminal hexahistidine fusion and was affinity purified by adsorption to a nickel-Sepharose column (in 50 mM phosphate [pH 7.8], 300 mM NaCl, and elution with a gradient of up to 500 mM imidazole). The 3Dpol samples were more than 95% pure, as shown in Fig. 1A. The wild-type (WT) proteins were shown to be active in elongation assays in which [32P]UTP incorporation was measured using a poly(A) template and an oligo(U)15 primer (5). However, under these conditions the relationship between enzyme concentration and nucleotide incorporation was not linear, indicative of a degree of cooperativity (Fig. 1B). This is particularly evident with the less-active serotype O enzyme. The addition of ZnCl2 (60 μM) to enhance the activity according to previously reported PV protocols (2, 8, 9, 12, 21) had little effect. Using a similar assay, we showed that the D388A mutant was unable to incorporate [32P]UTP, as expected (Fig. 1C). Interestingly, cooperativity had been demonstrated previously for other viral polymerases, including PV 3Dpol (11, 17).

Previously, we selected RNA aptamers to 3Dpol of the serotype C virus. RNA aptamers are derived from the process of in vitro selection and bind to their target molecules with high affinity and specificity, making them useful as molecular tools and potentially as inhibitors (5). We characterized aptamers that inhibited the...
activity of 3Dpol in an elongation assay. One of the aptamers was truncated to 32 mer (F47tr, 5’-GGGUUACAGAAAACCUCAG UUGCUGGUGUGU-3’), maintaining both its inhibitory capacity and its affinity for 3Dpol. Here, we also show that preincubation of 3Dpol with 47tr resulted in no incorporation of \[^{32}\text{P}]\text{UTP} (Fig. 1C).

To investigate the products of elongation assays (as described in Fig. 1C), samples were negatively stained prior to examination by transmission electron microscopy (TEM) according to a previously described protocol (22). PV 3Dpol was included as a control and shown to form fibrils similar to those described previously (11, 23) (Fig. 2A). In contrast, FMDV 3Dpol alone was disordered and aggregated, showing no evidence of defined higher-order structures (Fig. 2C). This result is in agreement with crystallographic data in which differences in crystal packing between the PV and FMDV enzymes suggest that FMDV 3Dpol alone is unlikely to form higher-order structures (7). Products of the FMDV elongation assay, however, appeared as ordered fibrils by TEM (Fig. 2B). The fibrils were 250 ± 27 Å in width and were narrower and more regular than the fibrils observed with isolated PV 3Dpol (460 ± 32 Å; n = 30). Fibrils formed by both PV 3Dpol and FMDV 3Dpol were variable in length. Both C and O 3Dpol were capable of fibril formation (Fig. 2B and D, respectively); however, the fibrils formed with the more active serotype C enzyme were more regular in morphology. There was no evidence for the formation of defined higher-order structures other than fibrils, unlike PV 3Dpol, which also formed sheets and tubes. Also in contrast with PV 3Dpol, all components of an elongation assay were necessary for the formation of the FMDV 3Dpol structures (i.e., primer, template, magnesium ions, and NTPs). The presence of dithiothreitol (DTT) or \(\text{Zn}^{2+}\) had no effect; however, EDTA inhibited both [\(\text{[\text{a-}^{32}\text{P]}\text{UTP}}\)] incorporation in the elongation assay and the formation of fibrils as seen by TEM (Table 1), indicating that fibril formation was not a spontaneous event, but rather requires active elongation to occur.

The TEM data correlated with light-scattering measurements of the assay undertaken over the same time scale (Fig. 3). A rapid increase in absorbance at 350 nm was observed with samples of isolated PV 3Dpol. A similar, but slower, increase was seen with FMDV 3Dpol (saturating after 25 to 30 min) when all components of an elongation assay were present. No increase in light scattering was detected when FMDV 3Dpol alone was incubated over time. In addition, substitution of WT 3Dpol for the D388A mutant also resulted in no increase in light scattering over time and no fibril formation visible by TEM (Fig. 3 and Table 1). Furthermore, fibrils were not observed in samples treated with RNase A (0.5 \(\mu\text{g}\) per 20-\(\mu\text{l}\) reaction mixture) or proteinase K (10 \(\mu\text{g}\) per 20-\(\mu\text{l}\) reaction mixture) at the endpoint of the assay (i.e., after 30 min) (Table 1). This indicates that the fibrils are composed of both 3Dpol and single-stranded RNA, rather than an RNA duplex. The walls of the fibrils were measured to be 60 to 70 Å in thickness, consistent with the presence of a single polymerase molecule (7) and suggestive of the structures being tubular in nature. However, further structural studies will be required to confirm this hypothesis. Interestingly, fibrils were also observed when part of the 5’ UTR (negative sense) was used as a template in an elongation assay, although these were less uniform, probably due to the heterogeneity of the RNA used (16). However, this does indicate that the fibrils are not an artifact resulting from the use of a poly(A) template and that 3Dpol is able to unfold structured RNA in order to produce a product.

In order to determine the effects of aptamer 47tr on fibril formation, this molecule was included in the assays. Preincubation of 3Dpol(C) with 47tr (at an equimolar concentration for 30 min prior to the addition of other components of the elongation assay) inhibited the formation of fibrils observable by TEM (Fig. 2E). This result correlated with no increase in light scattering detect-
able over time in the presence of the aptamer (Fig. 3). However, addition of the aptamer at the endpoint of the assay had no effect. The inhibitory effect of the aptamer on enzyme activity therefore correlates with the absence of fibrils, again demonstrating that these formed only when the polymerase was active. We had demonstrated previously that the aptamer had no inhibitory effect on the ability of PV 3Dpol to incorporate $^{32}$PUTP in an elongation assay (5). In agreement with this, the aptamer had no effect on the formation of PV 3Dpol fibrils by TEM (Fig. 2F).

The formation of a fibrillar replication lattice could have several benefits in viral replication. It could serve to increase the local

![FIG 2](image1) Formation of fibrils during the polymerase elongation assay. (A) PV 3Dpol alone (3 μM). (B) FMDV 3Dpol(C) (1 μM) in the context of a full elongation assay after 30 min. The FMDV fibrils have a regular width averaging 250 Å but exhibit considerable variations in length. (C) FMDV 3Dpol(C) alone. (D) Same as panel B but with the substitution of 3Dpol(O) for 3Dpol(C). (E) Same as panel B but with the addition of equimolar aptamer 47tr. (F) Same as panel A but with the addition of equimolar aptamer 47tr. The insets in panels A and B are higher-magnification views. Negative-stain TEM; scale bars, 500 Å.

![FIG 3](image2) Light-scattering measurements of fibril formation over a time course of up to 30 min at room temperature. Symbols: solid circles, PV 3Dpol alone (3 μM); open diamonds, FMDV 3Dpol elongation assay (EA) with 3 μM 3Dpol(C) and the method described for Fig. 1C; solid diamonds, FMDV 3pol(C) alone; shaded triangles, FMDV 3Dpol EA (as above) in the presence of equimolar aptamer 47tr; cross, EA using FMDV 3Dpol active-site mutant D388A. OD$_{350}$ nm, optical density at 350 nanometers.

### TABLE 1 Formation of 3Dpol fibrils under different experimental conditions

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<tr>
<th>Assay or component</th>
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<td>Full elongation assay</td>
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<tr>
<td>Full elongation assay</td>
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<tr>
<td>With catalytically inactive 3Dpol</td>
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<td>Without 3Dpol</td>
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<td>Without primer</td>
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<tr>
<td>With aptamer 47tr (1–5 μM)</td>
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<tr>
<td>With RNase (25 μg/ml)</td>
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<tr>
<td>With proteinase K (500 μg/ml)</td>
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<tr>
<td>With EDTA (10 mM)</td>
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<td>With DTT (5 mM)</td>
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concentration of components, and a clustering of binding sites could increase the efficiency of replication. It is also tempting to speculate that the higher-order structures have a role in preventing the formation of double-stranded RNA. During replication, a negative-stranded cRNA is first produced, which then acts as a template for the synthesis of positive strands. If the RNA is “coated” in polymerase molecules during this process, forming fibril-like structures, then the formation of a double-stranded negative/positive RNA complex could be prevented. This would allow replication to proceed without the necessity to melt an RNA duplex, which would be energetically demanding and therefore inefficient. It must be noted that in vivo, a number of other viral and host proteins (e.g., RNA helicase [10]) are also involved in replication. Our studies are therefore continuing with the use of an FMDV replicon (14) in order to probe the formation of higher-order structures in cells where all replication components are present.

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REFERENCES