Structural and functional insights into alphavirus polyprotein processing and pathogenesis

Gyehwa Shin, Rutgers State University
Samantha A. Yost, Rutgers State University
Matthew T. Miller, Rutgers State University
Elizabeth J. Elrod, Emory University
Arash Grakoui, Emory University
Joseph Marcotrigiano, Rutgers State University

Journal Title: Proceedings of the National Academy of Sciences
Volume: Volume 109, Number 41
Publisher: National Academy of Sciences | 2012-10-09, Pages 16534-16539
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.1073/pnas.1210418109
Permanent URL: https://pid.emory.edu/ark:/25593/s5cf2

Final published version: http://dx.doi.org/10.1073/pnas.1210418109

Copyright information:
© Shin et al. Freely available online through the PNAS open access option.

Accessed October 25, 2018 12:13 AM EDT
Structural and functional insights into alphavirus polyprotein processing and pathogenesis

Gyehwa Shin, Samantha A. Yost, Matthew T. Miller, Elizabeth J. Elrod, Arash Grakoui, and Joseph Marcotrigiano

*Center for Advanced Biotechnology and Medicine, Department of Chemistry and Chemical Biology, Rutgers University, Piscataway, NJ 08854; and ²Emory Vaccine Center, Division of Microbiology and Immunology, Department of Medicine, Division of Infectious Diseases, Emory University School of Medicine, Atlanta, GA 30322

Edited by Robert A. Lamb, Northwestern University, Evanston, IL, and approved August 31, 2012 (received for review June 19, 2012)

Alphaviruses, a group of positive-sense RNA viruses, are globally distributed arboviruses capable of causing rash, arthritis, encephalitis, and death in humans. The viral replication machinery consists of four nonstructural proteins (nsP1–4) produced as a single polyprotein. Processing of the polyprotein occurs in a highly regulated manner, with cleavage at the P2/3 junction influencing RNA template use during genome replication. Here, we report the structure of P23 in a precleavage form. The proteins form an extensive interface and nsP3 creates a ring structure that encircles nsP2. The P2/3 cleavage site is located at the base of a narrow cleft and is not readily accessible, suggesting a highly regulated cleavage. The nsP2 protease active site is over 40 Å away from the P2/3 cleavage site, supporting a trans cleavage mechanism. nsP3 contains a previously uncharacterized protein fold with a zinc-coordination site. Known mutations in nsP2 that result in formation of noncytopathic viruses or a temperature sensitive phenotype cluster at the nsP2/nsP3 interface. Structure-based mutations in nsP3 oppose the location of the nsP2 noncytopathic mutations prevent efficient cleavage of P23, affect RNA infectivity, and alter viral RNA production levels, highlighting the importance of the nsP2/nsP3 interaction in pathogenesis. A potential RNA-binding surface, spanning both nsP2 and nsP3, is proposed based on the location of inter- and intrabinding sites and adaptive mutations. These results offer unexpected insights into viral protein processing and pathogenesis that may be applicable to other polyprotein-encoding viruses such as HIV, hepatitis C virus (HCV), and Dengue virus.

Alphaviruses are small, enveloped RNA viruses that infect various vertebrates, including humans, horses, birds, and rodents (1). Transmission occurs via an invertebrate vector, usually mosquitoes. In humans, typical alphavirus infection can result in rash, arthritis, encephalitis, and death. Within the past decade, infection by Chikungunya virus (CHIKV), a member of the genus alphavirus, caused massive epidemics in Africa and Southeast Asia, with positive cases reported in Europe and the United States, raising the possibility of an emerging threat (2). Additionally, CHIKV often presents with nonclassic symptoms that go undiagnosed, making global pathogenicity difficult to monitor (3).

The alphavirus genome consists of single-stranded, positive-sense RNA of ~9–11 kb with a 5′ cap structure and 3′ polyadenosine tail. The genome contains two cistrons. The first, located in the 5′ two-thirds of the genome, encodes the viral replication machinery, termed nonstructural proteins (nsPs), whereas the structural proteins that form virus particles are encoded in the second cistron. The genomic RNA is used as an mRNA for the translation of the nsPs and as a template for the synthesis of complementary negative-sense RNA. The negative-sense strand is a template for subgenomic RNA containing the second cistron and progeny, genomic RNA. Because genomic and subgenomic RNAs are made in vast excess, the negative-sense RNA strand is considered a replication intermediate.

The alphavirus replication machinery is composed of four nonstructural proteins (nsP1 to -4), which are expressed as one of two polyproteins (P123 or P1234). P1234 is expressed as a read-through of an opal termination codon at the end of nsP3. These precursor polyproteins are cleaved by a protease within nsP2 (1, 4). After translation of P1234, cleavage at the P3/4 junction occurs either in cis or trans, followed by the P1/2 junction, which occurs in cis only (4, 5). Both P123+nsP4 and nsP1+P23+nsP4 preferentially synthesize negative strand viral RNA (6, 7). The final cleavage event between P23 produces fully mature nsPs and switches RNA template for synthesis of positive-sense genomic and subgenomic RNAs. The correlation between P23 cleavage and the switch from negative- to positive-sense RNA production is poorly understood.

In mammalian cells, alphavirus infection inhibits host gene expression, causes severe cytopathic effects (CPEs), disrupts the cellular response to IFN, and leads to cell death (8–11). Mature nsP2 contains an amino-terminal RNA helicase domain, a central protease domain that catalyzes all cleavages between the nonstructural proteins, and an inactive RNA methyltransferase-like (MT-like) domain (12). During infection, a portion of nsP2 localizes to the nucleus (13, 14) and plays a role in shutting off host-cell transcription and viral cytopathogenicity (9, 10). Two genetically distinct groups of alphaviruses termed New World and Old World have emerged because geographic isolation (8).

In Old World alphaviruses, such as Sindbis virus (SINV) and Semliki Forest virus (SFV), transcriptional shutoff is nsP2-dependent (15). Even in the context of a replicon lacking the structural proteins, the nsPs of SINV and SFV alone can efficiently shut off host transcription and induce severe CPE (16).

Mutations that cause alphaviruses to establish a persistent infection with minimal or no CPE have been identified (16–20). A key feature of these mutants is their inability to inhibit host-cell transcription (10, 15). The best-characterized noncytopathic mutations map to nsP2 proline 726 (P726) within the MT-like domain (10, 15). The P726 mutation has been shown to attenuate the cytotoxic effect caused by the expression of wild-type SINV nsP2 protein alone (10). Noncytopathic mutations of nsP2 P726 also reduce viral RNA replication levels and are unable to shut off host transcription and translation processes, showing


The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

Freely available online through the PNAS open access option.

Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org (PDB ID code 4GUA).

1G.S. and S.A.Y. contributed equally to this work.

2Present address: Science and Engineering Campus, Korea University, Seoul, Republic of Korea, 136-701.

3To whom correspondence should be addressed. E-mail: jmarco@cabm.rutgers.edu.

This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1210418109/-DCSupplemental.
the overall importance of nsP2 to viral replication, as well as host-cell interaction (11, 16).

The function(s) of nsP3 in alphavirus replication remains unresolved. Based on sequence conservation, nsP3 is organized into three domains: an amino-terminal macro or X domain (23), a central alphavirus-specific region, and a hypervariable-sequence carboxy terminus. Macro domains have been shown to bind ADP-ribose (24) with certain viral macro domains also retaining ADP-ribose-1′′-phosphatase activity (25, 26). Recent work in SFV shows that residues just after the macro domain of nsP3 play a role in positioning of the P23 cleavage site (27). Additionally, the carboxy-terminal region of nsP3 contains numerous phosphorylation sites targeted by yet unknown cellular kinases (28, 29).

Here, we present the structure of an uncleaved P23 precursor protein spanning the nsP2 protease domain through to the central, zinc-binding domain (ZBD) of nsP3 from SINV (Fig. L4). The P23 cleavage site is located in a narrow cleft formed between nsP2 and nsP3 that is inaccessible for proteolysis. Surprisingly, all of the previously reported nsP2 noncytopathic mutants lie at the interface between nsP2 and nsP3. Mutagenesis of residues in nsP3 opposite of nsP2 P726 showed inefficient cleavage of P23, delayed onset of pathogenesis and host gene shutoff, and altered viral RNA synthesis. Because P23-pro-zbd copurified with large amounts of RNA, we propose a potential RNA-binding surface that extends across both nsP2 and nsP3 and hypothesize that cleaving the P2/3 junction may alter the RNA-binding surface, thereby contributing to template switching by the replication machinery. Our results provide a better understanding of the mechanism of viral polyprotein processing and pathogenesis.

**Results**

**Alphavirus P23pro-zbd Structure.** The SINV P23 precursor protein extending from the nsP2 protease domain to the nsP3 central domain was shown to be fully intact by SDS/PAGE and mass spectrometry despite the presence of a competent nsP2 protease active site (Fig. S1 A and B). Sequence alignment of the central domain of nsP3 shows numerous absolutely conserved cysteine residues (Fig. S2), suggesting the presence of a metal ion-binding site. Various purified preparations of the middle region of nsP3 were analyzed by quantitative X-ray fluorescence for common transition metals found in metalloproteins (30, 31) and shown to have one zinc ion per molecule. Crystals of P23pro-zbd diffracted to 2.85 Å resolution and the structure was determined by molecular replacement with structures of nsP2pro (12) and nsP3 macro domain (32) (Table S1). The resulting electron density allowed for an unambiguous trace of the entire polypeptide chain, including the previously uncharacterized nsP3 central region containing the zinc-binding site. The location of the zinc ion was confirmed by anomalous difference maps (Table S1 and see Fig. S4B).

SINV P23pro-zbd shows a highly compact structure, with the four domains (protease and MT-like from nsP2; macro and ZBD from nsP3) arranged with each domain occupying a vertex of a rectangle (Fig. 1 B–D). The polypeptide chain progresses around the perimeter of the rectangle such that the nsP2 protease and nsP3 ZBD are adjacent to one another. The carboxyl terminus of the P23pro-zbd construct is pointing toward the protease active site (Fig. 1 B–D). A previously described SINV mutant (ts7) defective in RNA synthesis at nonpermissive temperatures contains a F312 to serine mutation in nsP3 (33). This residue is located in the hydrophobic core and surrounded by residues from the macro, ZBD, and MT-like domains (Fig. 1E). Mutation of this residue would likely destabilize the core and disrupt domain interaction, giving rise to the RNA" phenotype at the nonpermissive temperature.

In a previously described nsP2pro structure, the six carboxy-terminal amino acids were disordered (12). In the P23pro-zbd structure presented here, the entire polypeptide between nsP2 and nsP3 is ordered, permitting modeling of the P2/3 cleavage site. The P2/3 cleavage site is located at the base of a narrow cleft about 11−13 Å wide formed by the MT-like domain of nsP2 and the macro domain of nsP3 (Fig. 1 F and G). The cleavage site is 40 Å away from the nsP2 protease active site, supporting a trans cleavage mechanism (5, 34, 35). Although the 6 aa surrounding the scissile bond are solvent-exposed, attempts to position the protease domain onto the P2/3 cleavage site using the proposed model (36) resulted in numerous steric clashes.
ZBD in nsP3 Is Essential for Viral Replication. After the P2/3 cleavage site, the polypeptide chain continues into the nsP3 macro domain. The ADP ribose–binding site within the macro domain is solvent-exposed and points away from the other four domains. The polyprotein exits the last helix of the macro domain adjacent to the P2/3 cleavage site and continues into a stretch of nsP2 that lacks secondary structure but forms a twist (Fig. S3). The conserved nsP3 central domain contains an antiparallel α-helical bundle, two parallel β-strands, and a previously unknown zinc-coordination site (Fig. S4B). The zinc is coordinated by C263, C265, C288, and C306 of nsP3. C263 and C265 are located in the loop between the last two α-helices, whereas C288 and C306 are at the carboxyl termini of the two parallel β-strands (Fig. S3C). Each of the zinc coordinating cysteines is invariant among all alphavirus sequences (Fig. S2). Structural comparison of the nsP3 ZBD against all known folds in the Protein Data Bank using PDBe Fold (37) and Dali servers (38) failed to produce any statistically significant hits, and no metalloprotein folds were identified. The spatial arrangement of secondary structural elements that contribute to nsP3 zinc-binding site, i.e., two cysteines in a loop between alpha helices and the other two cysteines at the end of two parallel β-strands, represents a structural scaffold for zinc-ion coordination (39).

Site-directed mutagenesis was performed to examine the functionality of the four conserved cysteine residues involved in zinc coordination in nsP3 (Fig. S4A). All four SINV mutants (C263A, C265A, C288A, and C306A) failed to produce productive infection in BHK cells, as evaluated by infectious center and plaque assays (Fig. S4C). A conserved cysteine residue in nsP3 (C247) unrelated to the zinc-binding site served as a control and C247A mutant produced infectious particles to near wild-type levels. Zinc-coordination mutants failed to express the non-structural polyprotein and replicate virus, thereby supporting the structural role of the zinc ion (Fig. S4D).

The coordination site forms a distorted tetragon due to the presence of two highly conserved serine residues (S289 and S290) below the zinc atom (Fig. S2). Mutation of these serine residues did not alter viral replication levels. Furthermore, a double serine to alanine mutant (S289A+S290A) resulted in decreased viral RNA infectivity, but overall viral titers were comparable to wild type (Fig. S4C).

Importance of P23 Interface for Polyprotein Processing and Viral Pathogenesis. The interface between the macro and ZBD of nsP3 has a buried surface area of 570 Å², whereas the linker connecting the two domains is extended, making nsP3 into a ring-like structure with an inner diameter of 15–18 Å (Fig. 2A). Together, nsP2 and nsP3 share an extensive interface with 3,000 Å² of buried surface area. nsP3 encircles the MT-like domain of nsP2, with residue R781 of nsP2 protruding through the opening (Fig. 2C). The interface between nsP2 and nsP3 is charged, with the nsP2 surface being mostly basic, whereas nsP3 is generally acidic (Fig. 2B and C).

Several groups have identified mutations in the carboxyl-terminal portion of nsP2 that result in noncytopathic virus and persistent infection (16–20). These mutations reduce viral RNA replication levels with minimal effects on host gene expression and viral yields. All of the noncytopathic mutations map to the surface of nsP2 at the nsP3 interface (Fig. 2D). The mutations follow the encircling path made by the macro to ZBD linker. Mutagenesis of P726 in SINV to amino acids with large side chains (Phe, Tyr, Leu, Arg, or Gln) resulted in numerous steric clashes with nsP3, whereas Ser, Thr, Ala, and Val had fewer conflicts. Moreover, the P726G mutation could destabilize the Ω-β11 loop, causing it to interfere with P23 interaction. Attempts to purify P23Δpro-zbd with a P726S or P726L mutation resulted in poorly soluble and aggregated protein, supporting the importance of this position.

To investigate how destabilizing P23 interface may contribute to changes in viral pathogenesis and replication, site-directed mutagenesis to the corresponding surface of SINV nsP3 in close proximity to nsP2 P726 were created (Fig. 2E). The side chains of V162 and L165 of nsP3 are ~4.6 and 3.3 Å away from nsP2 P726 side chain, respectively (Fig. 2F). V162 and L165 of nsP3 were mutated to one of the following: Ala, Asp, Glu, Phe, Arg, or Trp, nsP2 P726 to Ser, Gly, or Leu mutants were used for comparison. RNA infectivity measured by infectious center assay on BHK cells were reduced in P726L, L165D, L165E, and L165R mutants compared with the wild-type SINV (Fig. 3A). P726L mutation in nsP2 reduced RNA infectivity to only 2% of wild-type levels, whereas the three nsP3 L165 mutants averaged an RNA infectivity at 8% of wild type. These four mutants also displayed a small size plaque phenotype in infectious center assay (Fig. S5 A–D). In addition,
reached wild-type levels of virus titers. All V162 mutants behaved similarly to wild type in both RNA infectivity and viral titer and were not pursued further. None of the nsP3 mutants displayed a noncytopathic phenotype in the TSG/puromycin N-acetyltransferase (PAC) replicon background (16), although onset of CPE in L165 to D, E, and R mutants was delayed.

Radiolabeling experiments were performed to evaluate both viral and host protein and RNA production over the course of infection and to analyze efficiency of host-cell transcriptional and translational shutoff caused by SINV nsP3 mutations. Wild-type SINV is able to halt host translation completely within 6 h postinfection, whereas P726S and P726G mutants were unable to do so even up to 12 h postinfection, as indicated by persistent actin production (Fig. S6). nsP2 P726L and the four selected nsP3 L165 mutants (Asp, Glu, Arg, Trp) were able to shut off host translation, although at a slightly delayed time postinfection compared with wild type (Fig. S6).

Tritium-labeled uridine was used to track viral and host RNA production over the course of infection. In the presence of actinomycin D, both genomic and subgenomic viral RNA production can be clearly observed in wild-type SINV starting as early as 2 h postinfection and remaining stable until 12 h (Fig. 3C). Levels of subgenomic and genomic RNA in P726G and P726S mutants are reduced compared with wild type. nsP2 P726L and the four nsP3 L165 mutants exhibit a slow initial rate of transcription, especially of genomic RNA; however, these mutants reach wild-type levels by 4 h postinfection. Thereafter, the levels of RNA replication in the L165 mutants continue to increase, and even surpass, wild-type RNA replication, especially subgenomic RNA (Fig. 3C). RNA labeling in the absence of actinomycin D showed complete inhibition of host cellular transcription by 8 h postinfection as demonstrated by decreasing 28S and 18S cellular rRNA (Fig. 3D). P726G and P726S mutants exhibited a lack of host-cell transcriptional shutoff with RNA persisting up to 12 h postinfection. The nsP2 P726L mutant and the four nsP3 L165 mutants showed a delay in host transcription inhibition but eventually achieved complete shutoff 12 h postinfection (Fig. 3D).

Given that residues R159 and E163 of SFV nsP3 have been shown to affect P2/3 cleavage (27), we assessed the effect of the SINV nsP3 mutations oncleavage activity. nsP3 linker region mutants L165D and L165E revealed drastic inefficiency in P23 cleavage that was not seen in L165W and L165R mutants (Fig. 3F and Figs. S7). The presence of uncleaved P23 in the nsP3 L165D and L165E mutants started 6 h postinfection and persisted up to 12 h, whereas P726 mutants, as well as nsP3 L165R and nsP3 L165W, showed substantially lower amounts of uncleaved P23. Western blot probed with anti-nsP3 antibodies were used to confirm the presence of uncleaved P23 (Fig. 3F). nsP3 appears as a smear, indicative of alternate phosphorylation states and possibly other posttranslational modifications (40). The nsP2 P726 mutants may also accumulate precleavage forms but at very low concentrations that do not build up over time, as in the case of the nsP3 linker mutants.

**Discussion**

The structure of alphavirus precursor polyprotein P23-pro-zbd presented here reveals the arrangement of four domains (protease, MT-like, macro, and ZBD) in a precleavage form. Based on the structure of P23-pro-zbd, several inferences can be made about the mechanism of alphavirus polyprotein cleavage. First, the carboxyl terminus of the P23-pro-zbd structure is pointing toward the nsP2 protease active site about 40–45 Å away. It is possible that the large, flexible carboxyl region of nsP3 could span the distance, placing the P3/4 junction in the protease active site for cis cleavage (34). Second, the 40 Å distance between the P3/2 cleavage site and the nsP2 protease active site supports the current trans model of cleavage (5, 34, 35). Third, the inaccessibility of the P23 cleavage site itself indicates that access is tightly regulated. It is possible the activator sequence present in the extreme...
Many important human viral pathogens encode a polyprotein that is cleaved by viral or cellular proteases. Given the wide prevalence of regulated polyprotein processing seen in a number of viral replication systems, it is not surprising that alphaviruses have adapted this system to ensure efficient virus replication.

The alphavirus genome contains conserved sequence elements (CSEs) that serve as promoters for production of positive- and negative-sense genomic RNA, as well as subgenomic RNA. Mutations in a 51-nt-long CSE in the nsP1-coding region of SINV disrupted the RNA replication structure without affecting the phenotype of the replication complex. Despite the large, bulky nature of the MT-like domain in the MT-like domain facilitates separation of nsP2 and nsP3 post-cleavage, which is triggered by binding viral proteins or RNA to the concave side of the nsP2 MT-like β-sheet.

Previously described nontypotypic mutations at the surface of nsP2 contacting the nsP3 linker region appear by molecular modeling to destabilize P23 interaction, decreasing efficiency of the RNA replication complex during virus infection. Altering the corresponding nsP3 residue (L165), which contacts nsP2 P726, did not produce virus with the same nontypotypic phenotype. In contrast to the nsP2 P726G or P726S mutants, nsP3 L165 mutants had enhanced RNA replication levels, indicating the apparent stabilization of the replication complex. Despite the large, bulky nature of a tryptophan substitution at L165, the virus remained similar in phenotype to wild type. The L165D and L165E mutants, however, exhibited the most extreme phenotype in terms of reduced RNA infectivity and inefficient P23 cleavage. The surface of nsP2 surrounding P726 is both basic and hydrophobic (Fig. 2F), suggesting that the P23 interface may be stabilized by the introduced acidic amino acid at L165 through interaction with the surface of nsP2. This stabilization could cause the interface to become “locked in,” preventing the flexibility needed for conformational change and, thus, disallowing nsP2 access to the P2/3 cleavage site.

The buildup of uncleaved P23 is evidence that the nsP3 linker region, which encircles nsP2, may function in positioning and recognition of the P2/3 cleavage site. Given the nsP3 linker residues and nsP2 P726 are within reasonable distance to the P2/3 cleavage site (21 and 17 Å from Cα of L165 or P726 to the cleavage site, respectively), it is plausible that mutation of the nsP3 linker region shifts local structural features, altering recognition of the cleavage site (Fig. 2F). A previously described E163R mutation in SFV nsP3 that altered P2/3 cleavage is completely solvent-exposed (Fig. 2E) (27). These results may indicate that the nsP3 linker residues surrounding nsP2 P726 are recognized by the nsP2 protease or another factor necessary for P2/3 cleavage.

Cleavage between P2/3 dictates template use by the alphavirus replication machinery. During purification, P23pro-zbd bound extensive amounts of bacterial RNA, which could only be removed by high concentrations of salt (Fig. S1A and C). The electrostatic potential of the P23pro-zbd structure identified a large basic surface that centers on the zinc-binding site of nsP3 and extends to nsP2 – morpholino)ethane sulfonic acid (MES), used in crystallization, were bound to P23pro-zbd along the basic surface, which may indicate potential binding sites for the phosphate groups of nucleic acids (Fig. 4 C–E). In fact, a molecule of MES occupies the ADP-ribose–binding site in each of the three macro domains in P23pro-zbd in the asymmetric unit, as well as in the macroH2A1.1 structure (Fig. 4E) (43). The alphavirus genome contains conserved sequence elements (CSEs) that serve as promoters for production of positive- and negative-sense genomic RNA, as well as subgenomic RNA. Mutations in a 51-nt-long CSE in the nsP1-coding region of SINV disrupted the RNA structure without affecting the coding sequence and demonstrated reduced replication in mosquito cells but not in mammalian cells (44). Adaptive mutations (E118K of nsP2 and T286I of nsP3) were shown to restore replication of the mutated CSE. T286 is 2 aa before one of the zinc-coordinating cysteine residues (C288) (Fig. 4F) and is located on the basic surface of the ZBD. T286 is 6.1 Å away from a sulfate ion that is coordinated by three highly conserved amino acids in nsP3: Y267, R273, and R276 (Fig. 4F). Taken together, these data strongly implicate this surface as a potential RNA-binding site.

Many important human viral pathogens encode a polyprotein that is cleaved by viral or cellular proteases. Given the wide prevalence of regulated polyprotein processing seen in a number of viral replication systems, it is not surprising that alphaviruses have adapted this system to ensure efficient virus replication.

Many important human viral pathogens encode a polyprotein that is cleaved by viral or cellular proteases. Given the wide prevalence of regulated polyprotein processing seen in a number of viral replication systems, it is not surprising that alphaviruses have adapted this system to ensure efficient virus replication.

Many important human viral pathogens encode a polyprotein that is cleaved by viral or cellular proteases. Given the wide prevalence of regulated polyprotein processing seen in a number of viral replication systems, it is not surprising that alphaviruses have adapted this system to ensure efficient virus replication.
of viral families, there is lack of structural information regarding the precursor forms. Currently, alphavirus P23pro-deletion and poliovirus 3CD (45) are the only structures of polyprotein precursor from a viral replication machinery. Both alphavirus P23pro-deletion and poliovirus 3CD structures contain the protease domain responsible for cleaving the precursor, which is thought to proceed by a trans mechanism. However, 3CD has no intramolecular contacts and a solvent accessible cleavage site that is regulated by intermolecular contacts, whereas P23pro-deletion has an extensive interface and an inaccessible cleavage site. The results presented here further expand our current understanding of viral polyprotein processing, replication, and cytopathogenesis, which may be applicable to other positive-sense RNA viruses.

Materials and Methods
Complete materials and methods are provided in SI Materials and Methods.

Structure Determination. P23pro-deletion of the SINV strain Toto110 was expressed in Escherichia coli as a fusion with GST. The protein was purified by sequential chromatography over GST affinity, hydroxyapatite, and heparin columns. Crystals of P23pro-deletion were grown by vapor diffusion against 2 M ammonium sulfate and 0.1 M MES (pH 6.5). Crystals were transferred to a cryoprotectant solution containing 2 M ammonium sulfate, 0.1 M MES (pH 6.5), 3% (vol/vol) xylitol, and 5% (vol/vol) trimethylamine-N-oxide and then flash-cooled in liquid nitrogen. Diffraction data were collected at the zinc, K absorption edge (1.28 Å) to 2.85 Å resolution at beam line X25 at Brookhaven National Synchrotron Light Source (Table S1). Phases were obtained by molecular replacement using the nsP29 (12) and nsP3 macro structures (32). The model contains 3 molecules of P23pro-deletion, 3 zinc ions, 31 sulfate ions, and 5 molecules of MES. Atomic coordinates have been deposited in the Protein Data Bank under PDB ID code 4GUA.

Virological Studies. Infectious center and plaque assays were performed similarly to those described in ref. 46. Protein and RNA radiolabeling were performed as described in ref. 9. Harvested protein from BHK-21 cells at time points postinfection were used in Western blotting experiments, and membranes were probed with anti-nsp2, anti-nsp3, and anti-GAPDH antibodies.

ACKNOWLEDGMENTS. We thank J. Bonanno for assistance with the data processing. C. Rice and M. MacDonald for providing BHK-21 cells, Sindbis virus DNA, and nsP3 antibody. J. Millong and P. Matteson for the GAPDH antibody; and A. Basant, J. Chiu, F. Jiang, A. Khan, C. Rice, A. Stock, V. Stollar, and J. Whidby for insightful discussions. S.A.V. was supported by the NIH-NIAID training Grant T32AI007403-19. This work was supported by National Institutes of Health Grants AI080659 (to J.M.) and DK083356 and AI070101 (to A.G.). New York Commission on Cancer Research Grant 10-1962-CCR-E (to J.M.) and Swiss National Science Research Grant 3100A001615 (to A.G.). We wish to express our profound gratitude to the late Dr. Aaron Shatkin for his unwavering support, sage advice, and insightful discussions.