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Structure and Cleavage Specificity of the Chymotrypsin-Like Serine Protease (3CLSP/nsp4) of Porcine Reproductive and Respiratory Syndrome Virus (PRRSV)

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Biogenesis and replication of the porcine reproductive and respiratory syndrome virus (PRRSV) include the crucial step of replicative polyprotein processing by self-encoded proteases. Whole genome bioinformatics analysis suggests that nonstructural protein 4 (nsp4) is a 3C-like serine protease (3CLSP), responsible for most of the nonstructural protein processing. The gene encoding this protease was cloned and expressed in Escherichia coli in order to confirm this prediction. The purified protein was crystallized, and the structure was solved at 1.9 Å resolution. In addition, the crystal structure of the Ser118Ala mutant was determined at 2.0 Å resolution. The monomeric enzyme folds into three domains, similar to that of the homologous protease of equine arteritis virus, which, like PRRSV, is a member of the family Arteriviridae in the order of Nidovirales. The active site of the PRRSV 3CLSP is located between domains I and II and harbors a canonical catalytic triad comprising Ser118, His39, and Asp64. The structure also shows an atypical oxyanion hole and a partially collapsed S1 specificity pocket. The proteolytic activity of the purified protein was assessed in vitro. Three sites joining nonstructural protein domains in the PRRSV replicative polyprotein are confirmed to be processed by the enzyme. Two of them, the nsp3/nsp4 and nsp11/nsp12 junctions, are shown to be cleaved in trans, while cis cleavage is demonstrated for the nsp4/nsp5 linker. Thus, we provide structural evidence as well as enzymatic proof of the nsp4 protein being a functional 3CLSP. We also show that the enzyme has a strong preference for glutamic acid at the P1 position of the substrate.

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Abbreviations used: nsp, nonstructural protein; 3CLSP, 3C-like serine protease; EAV, equine arteritis virus; SARS-CoV, severe acute respiratory syndrome coronavirus; 3C*pro, 3C-like protease; GST, glutathione S-transferase; MR, molecular replacement; SeMet, selenomethionine; PDB, Protein Data Bank; SGPE, Streptomyces griseus proteinase E; MS, mass spectrometry; MW, molecular weight; ESI, electrospray ionization.
Introduction

Porcine reproductive and respiratory syndrome virus (PRRSV) was first discovered in North America in 1987\(^1\)\(^2\) and almost simultaneously (in 1990) in Europe.\(^3\) It manifests itself clinically with severe reproductive failure in sows and respiratory distress in neonatal pigs.\(^2\) In recent years, the virus has become endemic throughout the world, leading to significant economic losses in pig production.\(^4\) In the autumn of 2006, unparalleled outbreaks of atypical PRRS occurred in more than 10 provinces (or autonomous cities) of China, affecting more than 2 million pigs with nearly 400,000 fatal cases. Adult and growing pigs were also affected, which was different from classical PRRS.\(^5\) In 2007, an almost identical atypical PRRSV was found to be epidemic in Vietnam and China.\(^6\) Presently, only inactivated or attenuated vaccines are available to combat PRRS, but their efficacy is less than satisfactory due to adaptive evolution of PRRSV,\(^7\)\(^8\) especially in the cases of the atypical PRRS outbreaks in China and Vietnam.\(^5\)\(^6\) This urgent issue prompted us to exploit the possibility of drug discovery for treatment of the highly pathogenic PRRSV infection.

Like other RNA viruses in the order Nidovirales, such as equine arteritis virus (EAV), human coronavirus 229E, and severe acute respiratory syndrome coronavirus (SARS-CoV), PRRSV has a single-stranded positive-sense RNA genome that is about 15.3 kb in size, containing a large 5' termini replicase gene and a downstream set of structural protein genes.\(^9\) The replicase gene is translated into two large polyprotein precursors, 1a and 1ab (with molecular masses of about 269.1 and 430.3 kDa for pp1a and pp1ab, respectively), with the latter being synthesized following a ribosomal frame-shifting event.\(^10\) After the rapid autocatalytic release of nonstructural protein 1 (nsp1) and nsp2, the remainder of the polyproteins are cleaved into 10 mature nonstructural proteins (nsp3–nsp12) by the 3C-like serine protease (3CL\(\text{SP}\)).\(^9\)\(^11\) The functional importance of the 3CL protease in the viral life cycle makes it an attractive target for the development of drugs directed against PRRSV.

Here, we confirm that PRRSV nsp4, which we demonstrate to extend from residue G1780 to E1983 in the viral polyprotein of the virus strain JXA1, is a 3CL\(\text{SP}\) from both the structural and enzymatic points of view. The structure of the enzyme reveals two chymotrypsin-like β-barrel domains and an extra C-terminal α/β domain similar to that observed in EAV nsp4.\(^12\) A canonical catalytic triad that is composed of Ser118, His39, and Asp64 is located in the open cleft between the two β-barrel domains. An S1 subsite consisting of His133 and very likely Ser136 and an oxyanion hole without the signature helix-like turn are seen in our structure. Our results also clearly show that the recombinant enzyme displays proteolytic activity in vitro. Besides, based on the results of the cleavage assay, three nsp junction sites in the PRRSV replicase precursor are confirmed, thereby shedding light on the processing pathway of pp1a and pp1ab in the virus.

Results

Overall description of the recombinant PRRSV 3CL\(\text{SP}\) used in this study

As a member of the Arteriviridae, PRRSV shows substantial similarities in genome organization to EAV—the prototype virus of this family.\(^13\)–\(^15\) Accordingly, whole genome bioinformatics analysis predicts that the PRRSV nsp4 domain should be the main protease responsible for processing most of polyproteins pp1a and pp1ab.\(^15\) Previously, we reported on the successful expression of a gene construct encoding this putative serine protease in Escherichia coli as well as the crystallization of the protein.\(^16\) As shown in Fig. 1a, after removal of the glutathione S-transferase (GST) tag, nine additional amino acid residues were still left at the N-terminus of the protein. The in vitro peptide cleavage assay in this study was performed using this recombinant enzyme with the nine extra residues present, while for the cis cleavage assay, even more additional N-terminal residues were present (Fig. 1b). In both cases, the recombinant protein displayed proteolytic activity in vitro, thereby confirming the identity of nsp4 as the main protease in PRRSV replication (see results below).
Structure determination and quality of the refined structure

Although the sequence identity between PRRSV and EAV nsp4s is more than 34%, we failed to solve the structure of PRRSV 3CLSP by molecular replacement (MR) using the structure of EAV nsp4 as a search model. The presence of two methionine residues in the recombinant PRRSV 3CLSP suggested that selenomethionine (SeMet)-based multiple-wavelength anomalous dispersion17 could be used to solve the phase problem. The unit-cell dimensions of the crystals (\(a = 112.4\) Å, \(b = 49.2\) Å, \(c = 42.9\) Å, \(\beta = 110.3^\circ\); space group \(C2\)) indicate that there is only one molecule per asymmetric unit, and we successfully located both of the two crystallographically independent selenium sites. Based on the obtained phases, a final model with good stereochemistry was built (for details, see Materials and Methods).

As shown in Table 1, the structure was refined to 1.9 Å resolution, with \(R_{\text{cryst}} = 19.6\%\) and \(R_{\text{free}} = 24.6\%\); 89.5% of the residues lie in the most favored regions of the Ramachandran plot and 10.5% lie in the additionally allowed regions.18 The quality of the electron density map of the recombinant protease at 1.9 Å is excellent, with an overall temperature factor of 22.2 Å\(^2\), and allowed us to model the positions of the main chain, most side-chain atoms, and 208 solvent molecules unambiguously. At the polypeptide chain termini, electron density is visible starting from residue 3 and ending at residue 199, but loop 136–140 is not seen in the maps due to disorder.

The overall structure of PRRSV 3CLSP and comparison with EAV nsp4

The PRRSV 3CLSP is folded into three domains, including two antiparallel \(\beta\)-barrels and an extra \(\alpha/\beta\) domain (Fig. 2a and b). The N-terminal \(\beta\)-barrel (domain I) consists of six \(\beta\)-strands (aI to fI) and a short \(\alpha\)-helix (helix A) that closes the barrel like a lid, whereas the middle \(\beta\)-barrel (domain II) comprises seven \(\beta\)-strands (aII to gII) and is connected to domain I via a very long loop (amino acids 69–89) (Fig. 2a). The catalytic site is located at the opening of the cleft between domains I and II and consists of residues Ser118, His39, and Asp64 (Fig. 2a), which are totally conserved among members of the Arteriviridae (Fig. 2c). The extra C-terminal domain (domain III) comprises residues 157–199 and consists of two pairs of short antiparallel \(\beta\)-sheets (strands aIII–cIII and bIII–dIII) and two \(\alpha\)-helices. The crossing angle is \(\sim 90^\circ\) between helix B and helix C (Fig. 2a). A long N-terminal loop extends over domain II to the vicinity of domain III. The orientation of this loop is stabilized by the three hydrogen bonds Arg4 N…Glu100 OE1, Thr5 N…Va199 O, and Thr5 OG1…Va199 N (Fig. 2a). The phenyl ring of Phe3 is directed into the hydrophobic interface between domains II and III and, therefore, also helps to stabilize the orientation of the N-terminal loop (Fig. 5a, left image).
Table 1. Data collection and refinement statistics

| Data collection                  | Native protein | SI18A mutant |
|----------------------------------|----------------|--------------|
| Space group                      | C2             | C2           |
| Cell dimensions                  |                |              |
| a, b, c (Å)                      | 112.4, 49.2, 42.9 | 113.3, 48.6, 42.9 |
| α, β, γ (°)                      | 90.0, 110.3, 90.0 | 90.0, 110.4, 90.0 |
| Resolution range (Å)             | 32.3           | 2.0        |
| Wavelength (Å)                   | 1.5418         | 1.5418      |
| Resolution range (Å)             | 0.9793         | 0.9000      |
| Rmerge                          | 0.045 (0.167)  | 0.213 (0.459) |
| Average I/σ(I)                   | 22.1 (4.3)     | 12.5 (1.4)  |
| Completeness (%)a                | 92.5 (91.6)    | 95.7 (77.6) |
| Redundancy                       | 4.5 (4.3)      | 4.5 (3.4)   |
| Rmerge/Reff (%)b                 | 19.6/24.6      | 18.2/21.5   |
| Ramachandran plot (%)c           |                |              |
| Most favored regions             | 89.5           | 90.2        |
| Additionally allowed regions     | 10.5           | 9.8         |
| Generously allowed regions       | 0              | 0           |
| Disallowed regions               | 0              | 0           |
| r.m.s.d. from ideality           | 0              | 0           |
| Bonds (Å)                       | 1.572          | 1.529       |
| Angles (°)                       | 6.832          | 6.832       |
| Average R-factor (Å²)            | 0.017          | 0.018       |
| Protein                         | 28.120         | 28.120      |
| Water                            | 1.472          | 1.529       |

* Values for the outermost resolution shell are given in parentheses.

b Rmerge = ΣΣ[Io–(I)]/ΣΣ[Io], where Io is the observed intensity and (I) is the average intensity from multiple measurements.

c Reff = Σ[|Fo–Fc|]/Σ|Fo|, where Fo and Fc are the structure-factor amplitudes from the data and the model, respectively. Reff is the R-factor for a subset (%5) of reflections that was selected prior to refinement calculations and was not included in the refinement.

d Ramachandran plots were generated by using the program PROCHECK.

A Dali search of the Protein Data Bank (PDB) database gave a variety of proteins with significant structural similarities to our structure, among which the EAV nsp4 expectedly had the highest Z score (21.7). As the prototype structure of nsp4s in Arteriviridae, the EAV 3CLSP also folds into three domains. Superposing domains I and II of PRRSV 3CLSP onto those of the EAV nsp4 (copy A of the four copies in the asymmetric unit) yielded an r.m.s.d. of 2.04 Å for 122 equivalent (out of 134 compared) Cα pairs (Fig. 3a), while domain III alone exhibited an r.m.s.d. of 0.52 Å for 30 (out of 38) Cα pairs (Fig. 3d), indicating that the two enzymes share great structural similarity at the tertiary level. However, the steric arrangement of the domains in the two proteases is quite different. An ~8Å shift of domain III relative to domains I and II was observed in our structure, when compared to EAV nsp4. This shift causes the whole domain III to move away from the cleft where the catalytic triad is located (Fig. 3a). Clear electron density for the loop Asn153-Ile157 that connects domains II and III, as shown in Fig. 3c, leaves no doubt about the chain tracing of the loop and verifies the 8Å shift of domain III, compared to EAV nsp4. The shift may also account for our failure to solve the structure by MR using EAV nsp4 as the search model.

Further discrepancies were also observed in two loop regions. One is the loop connecting strands eII and dII (Cys111-Val121 in PRRSV nsp4). In EAV nsp4, this loop is much like a tadpole with its head lying next to the catalytic nucleophile Ser120 (Fig. 3b). However, the corresponding loop in our structure is approximately “J” shaped with a protuberance pointing to domain III (Fig. 3a and b) on which the hydrophobic side chain of Phe112 resides. This residue is conserved in PRRSV and lactate dehydrogenase-elevating virus 3CLSP but not in EAV nsp4, where it is substituted by Trp (Fig. 2c). The phenyl ring of Phe112 protrudes into the hydrophobic interface between domains II and III (Fig. 5a, left image) and thereby causes the “J”-shaped conformation of the dII–dII loop.

Another loop of great variance is the one connecting strands eII and III. In PRRSV 3CLSP, this loop is eight residues in length, from Gly135 to Gly142, and is highly flexible as proved by the total lack of electron density for residues 136–140. However, in EAV nsp4, the corresponding region is mainly a β-strand with a much shorter loop, indicating that the conformation of this part is relatively rigid (Fig. 3b).

Interestingly these two regions showing significant variance are actually believed to cover, respectively, the S1 specificity pocket and the oxyanion hole in EAV nsp4, two highly important features involved in substrate recognition and catalysis by 3CL proteases.

The S1 specificity pocket

In 3CL proteases, the S1 specificity pocket accommodates the side chain of the substrate P1 residue and therefore determines the enzyme’s
Fig. 2. Crystal structure of PRRSV nsp4. (a) Cartoon representation of the overall structure of PRRSV nsp4. The N-terminal (aI–fI) and middle (aII–gII) β-barrels are colored green and cyan, respectively, with the red broken line indicating the loop 136–140, which is totally devoid of visible electron density. The C-terminal extra domain is shown in yellow. Members of the catalytic triad and the three hydrogen bonds that are believed to stabilize the orientation of the long N-terminus are featured. The long loop connecting domains I and II is highlighted in red. (b) Topological presentation of the same structure. (c) Multiple sequence alignment of the 3CLSPs of the members in the family Arteriviridae. The catalytic triad is indicated by red arrows, and the three residues that are believed to constitute the S1 specificity pocket in EAV nsp4 are marked with blue stars. The Phe residue, which is thought to lead to the atypical oxyanion hole in PRRSV 3CLSP, is labeled by a dark triangle. Cys111 and Cys115, which have been referred to in the text, are marked with black squares. LDVC and LDVP, lactate dehydrogenase-elevating virus neurovirulent type C and strain Plagemann, respectively; PRRSVAS and PRRSVES, PRRSV American strain and European strain, respectively.
preference for specific amino acid residues at this position. To identify the possible residues that may be involved in the formation of the S1 pocket in PRRSV 3CLSP, we used the structure of Streptomyces griseus proteinase E (SGPE) in complex with a tetrapeptide as the reference structure for superposition. SGPE is another chymotrypsin-like protease that has a substrate preference for Glu at the P1 position. Both the EAV nsp4 (copy A) and our structure were aligned. As shown in Fig. 4a, the S1 specificity pocket of both the SGPE and EAV nsp4 is composed of three residues, with a conserved His (His134 in EAV nsp4 and His213 in SGPE) lying at the bottom of the pocket and one Ser (EAV nsp4: Ser136; SGPE: Ser120) lining one side and another Thr/Ser (Thr115 in EAV nsp4, Ser192 in SGPE) on the other side. However, only the histidine (His133) could be found in our structure. The first serine involved in the S1 subsite is conserved in PRRSV 3CLSP (Ser136) (Fig. 2c), but we cannot define its position due to the absence of electron density for the loop 136–140 as mentioned before. Nevertheless, there is no residue in PRRSV nsp4 to match the Thr/Ser on the other side of the pocket. Thr113 of PRRSV nsp4, which is also highly conserved among

![Structural alignments between EAV (magenta) and PRRSV (light blue) nsp4s.](image)

**Fig. 3.** Structural alignments between EAV (magenta) and PRRSV (light blue) nsp4s. (a) Superposition of the two β-barrel domains of PRRSV 3CLSP with those of EAV nsp4. A shift of about 8 Å observed in the C-terminal extra domain is highlighted. The well-aligned catalytic triad of the two proteases is shown in stick representation. The two loop regions that display great discrepancies between the two enzymes are also featured. Residue Phe112, which makes hydrophobic interactions with domain III in PRRSV nsp4, is highlighted by line representation. (b) Magnified view of the two loop regions referred to in the text. (c) Stereo view of the electron density map of the loop (N153-I157) connecting domains II and III in PRRSV nsp4. This 2|Fo|−|Fc| map is contoured at 1.0σ above the mean. (d) Alignment of domain III alone of PRRSV and EAV nsp4 to show their well superposition.
members of *Arteriviridae* (Fig. 2c), is too far away to contact the substrate P1 residue (Fig. 4a).

An atypical oxyanion hole without the signature helix-like turn

Both EAV nsp4 and SGPE form a canonical oxyanion hole that is characterized by a helix-like turn preceding the nucleophilic serine. As shown in Fig. 4b, the backbone amides of Ser195 and Gly193 in SGPE can form hydrogen bonds with and therefore stabilize the carbonyl oxyanion in the tetrahedral intermediate during catalysis. Although, in EAV nsp4, only copy B has the normal active oxyanion hole geometry while the other three copies only form collapsed ones, this difference is mainly caused by flipping of the peptide bond between Ser117 and Gly118 rather than a big change of the orientation of the signature turn.12 However, in PRRSV nsp4, this oxyanion-specific turn is missing. Instead, a much more stretched loop leaves only the amide group of the catalytic nucleophile (Ser118) in contact with the oxyanion. We term this rare property an atypical oxyanion hole.

A strong hydrophobic interaction between domains II and III

Compared to that of the EAV nsp4, an even stronger hydrophobic interaction between domains II and III is observed in our structure. With Phe3, Val103, Phe108, Phe110, Phe112, Ile157, Leu162, Phe165, Phe166, Ile182, L196, and L197, the interface harbors 12 hydrophobic residues, whereas in EAV nsp4, this interface only includes eight such residues (Fig. 5a). This hydrophobic core holds domains II and III together and likely forces the loop connecting the two domains to adopt its observed orientation, leading to the 8-Å shift as described above.

Another unexpected feature of the extra domain is the presence of two patches of solvent-exposed hydrophobic residues, implying possible interactions of PRRSV nsp4 with other proteins. One patch, which is also present in EAV nsp4 (Fig. 5b, right image),12 consists of Leu173, Ile178, and Ile183 (Fig. 5b, left image). Another patch, comprising Val171, Val176, Ala195, Ala198, and Ala199, is even more extended and locates on a surface that is almost perpendicular to the former one (Fig. 5c, left image, and 5b, left image). It is noteworthy that this second hydrophobic patch is totally absent in EAV nsp4 (Fig. 5c, right image).

Trans cleavage of domain junctions in pp1a/pp1ab

Peptides corresponding to eight predicted junctions in PRRSV pp1a or pp1ab (Fig. 1a; Table 2) were synthesized, tested for trans cleavage, and further analyzed by high-performance liquid chromatography (HPLC) in order to see whether the recombinant
nsp4 is proteolytically active in vitro. In our assay, we showed that peptides corresponding to two junction sites can be cleaved by the enzyme. For each predicted cleavage site, the scissile peptide bond was between Glu and Gly.

Just like other reports on peptide-based cleavage assays, our enzyme also shows a remarkable difference in efficiency towards different peptides. Of the nine potential substrates investigated, only three can be cleaved to the extent of being detectable on an HPLC system (see Table 2). Peptide SP8-1, derived from the junction sequence between nsp11 and 12, is most susceptible to this enzyme. After incubation with the recombinant enzyme, two new peaks with retention times of 16.9 and 17.4 min, respectively, were observed after analyzing a reaction volume of 40 μl on a C18 column (Fig. 6b). The uncleaved peptide gave a peak at 18.5 min, consistent with that for the peptide-alone control (Fig. 6a). The two product peaks were collected and analyzed by mass spectrometry (MS). The results showed that the calculated molecular masses corresponding to the peaks were 1241.6293 and 802.3875, respectively, matching their theoretical monoisotopic molecular weight (MW) perfectly (the theoretical MW for fragment KDKTAYFQLE is 1241.6291, and for GRHFTW, it is 802.3874). A typical representation of the MS result is shown in Fig. 6f. When the catalytic serine residue was replaced by alanine (Ser118Ala mutant), no detectable peaks of cleaved peptides were observed in the HPLC profile (Fig. 6c). This confirmed the crucial function of Ser118 as the nucleophile in catalysis.

Another peptide that was shown to be processed is SP1, which corresponds to the nsp3/nsp4 junction. However, the proteolytic efficiency for SP1 was much lower than that for SP8-1. Only a small portion (~20%) of the peptide was cleaved after a 15-h incubation (data not shown).

The kinetic parameters of the protein were determined in vitro using a fluorescent assay system to quantitatively test the proteolytic activity of the recombinant enzyme. The results showed that the enzyme exhibits a $K_m$ of about 2.24 mM and a $k_{cat}$ of about 0.435 min$^{-1}$ for the substrate Dabsyl-KDTAYFQLE↓GRHFE-Edans.
Table 2. Comparative cleavage of peptides corresponding to polyprotein junctions in PRRSV JXA1

| Peptide | Polyprotein junction | Sequence(P–P′) |
|---------|----------------------|----------------|
| SP1     | nsp3/nsp4            | P10 S Q L G S L E G A F R T Q K P + |
| SP2     | nsp4/nsp5            | P9 L A A K P E L E G G L S T V Q L _ |
| SP3     | nsp5/nsp6            | P8 F F L R Y F A E G K L R E G V S F |
| SP4     | nsp6/nsp7            | P7 Q S C G M S H E S L T G A L A M - |
| SP5     | nsp7/nsp8            | P6 E N R R A Q W E A A K L S V E Q _ |
| SP6     | nsp9/nsp10           | P5 E K L R S N H E G K K S R M C G - |
| SP7     | nsp10/nsp11          | P4 K R A I C A D L E G S S S P L F K - |
| SP8     | nsp11/nsp12          | P3 K D K T A Y F Q L E G R H F T W Y Q +++ |
| SP8-1   | nsp11/nsp12          | P2 K D K T A Y F Q L E G R H F T W _ |
| SP8-M   | nsp11/nsp12          | P1 K D K T A Y F Q L E G R H F T W + / - |

Synthetic 16-mer peptides that represent the eight predicted junction sequences in the replicative polyprotein were incubated with 25 μM recombinant PRRSV nsp4 as described in Materials and Methods and analyzed by HPLC for the extent of cleavage. After incubation at 37 °C for 15 h, if more than 50% of the peptide was found to be cleaved as is determined by the decrease in area of the peak representing the parental peptide, ‘+++’ is given; if 10–20% of the parental peptide is cleaved, we perceive it as ‘+’; if less than 10% is cleaved but still detectable amounts of products are observed as marked by the appearance of new peaks in the HPLC profile, we mark it ‘+/−’; ‘−’ means no visible product peaks in the HPLC profile could be observed at all. All the cleaved products were verified by MS and/or comparative with HPLC traces of peptide standards. SP8-1 and SP8 correspond to the same junction site in pp1b. SP8-1 was synthesized and analyzed for the purpose of separating the two product peaks from each other.

The glutamic acid at the P1 position of the peptide substrate is crucial for effective cleavage and cannot be replaced by glutamine

3C-like proteases of RNA viruses are specific for either Gln or Glu at the P1 position of the substrate. Among the nidoviruses, coronavirus 3Cpro cleavage sites always have a Gln at the P1 position.9,23,24. However, in the same order Arteriviridae, the 3CLSPs of Arteriviridae, which are represented by the prototype EAV nsp4, show a clear preference for glutamic acid over glutamine at the P1 position of the substrate.7 In EAV, of the eight cleavage sites of the replicative polyprotein processed by the main protease, only one junction contains a glutamine at the P1 position (Q2837/S joining nsp10/11), while all other cleavage sites have glutamic acid in this position.11,25,26. Based on the similarity in genome organization between PRRSV and EAV, the putative cleavage sites processed by 3CLSP were predicted in PRRSV, all of which harbor a Glu at the P1 position.15 Therefore, it is a reasonable assumption that the 3CLSP of PRRSV also shows preference for Glu over Gln at the P1 position of the substrate.19,35. It is also consistent with the fact that it contains a hexa-His tag at both termini while the wild-type protein only harbors the hexa-Histidine tag to confirm this conclusion. Two different amounts (2 and 1 μg) of both proteins were loaded. At both amounts, the wild-type protein showed no signal for the Myc epitope, in contrast to a strong Myc signal given by the mutant protein (Fig. 7c). Both proteins displayed an unambiguous His-tag signal, which was much stronger in the case of the mutant protein, consistent with the fact that it contains a hexa-His tag at both termini while the wild-type protein only has the N-terminal tag left (Fig. 7b). These results clearly showed that his-nsp4[nsp4/nsp5]-Myc-his removed its C-terminal tail via cis cleavage after expression.

The purified wild-type protein was sent to the National Center of Biochemical Analysis, Academy
Fig. 6. Proteolytic activity of recombinant PRRSV nsp4 in an in vitro peptide cleavage assay using either SP8-1 or SP8-M as substrate. Two hundred fifty micromolar of either peptide was incubated either with 25 μM nsp4 or 25 μM S118A mutant protein or without any enzyme added at 37 °C for 15 h and analyzed by HPLC (for details, see Materials and Methods). Two additional peaks were observed for the peptides incubated with nsp4. The retention time and the identity of each peptide are given. (a) SP8-1 alone. (b) SP8-1 plus nsp4. (c) SP8-1 plus nsp4 (S118A). (d) SP8-M alone. (e) SP8-M plus nsp4. (f) A typical MS result of the product peptide with a retention time of 16.89 min in the HPLC profile for SP8-1 incubated with nsp4. Its calculated MW is determined using the formula shown in the figure. 1.0074 represents the accurate MW of the hydrogen atom.
of Military Medical Sciences, for C-terminal sequencing to further ascertain the exact peptide bond cut by the enzyme. After digestion of the protein using trypsin, a peptide fragment with observed mass of 2117.06 was found and selected for further sequencing. 

![Image](image_url)

Fig. 7. The cis cleavage activity of PRRSV nsp4. Both his-nsp4-[nsp4/nsp5]-Myc-his and his-nsp4(S118A)-[nsp4/nsp5]-Myc-his proteins bear the nsp4/nsp5 junction sequence fused to a Myc epitope at the C-terminus. The N- and C-terminal hexa-His tags were introduced by the pET-28b vector. After induction by IPTG at 37 °C for 5 h, the cells were harvested, lysed, and then purified by affinity chromatography on a Ni-NTA column followed by gel filtration. Coomassie staining as well as Western blot analysis results are shown. (a) The 12% SDS-PAGE separation profile of both the wild-type and the mutant fusion proteins. The uninduced cell sample, cell lysate after induction, and the purified protein are indicated. (b) Detection of the wild-type and the mutant fusion proteins by the anti-His antibody. (c) Detection of both proteins using an anti-Myc antibody. (d) C-terminal sequencing result of the purified his-nsp4-[nsp4/nsp5]-Myc-his protein. After digestion of the protein by trypsin, a peptide fragment of 2117.06 Da was chosen to sequence by nano-ESI-MS/MS. The deduced sequence (read from right to left) is identical with that of the C-terminal portion of the his-nsp4-[nsp4/nsp5]-Myc-his protein after cleavage at the E1983/G junction, confirming the cleavage.

The structure of the S118A mutant of 3CLSP

The 2.0-Å structure of the S118A mutant protein was solved by MR using the wild-type structure as the search model and refined to $R_{cryst}=18.2\%$ and $R_{free}=21.3$ (see Table 1). The overall structure of the
The mutant protein is almost identical with that of the native 3CLSP, with an r.m.s.d. of only 0.19 Å. The structure contains residues 3–199 but, like the wild-type structure, lacks electron density for the loop 136–140. As shown in Fig. 8a, there is clear electron density for the catalytic center, demonstrating that substitution of Ser118 by Ala was successful. At the same time, the substitution did not change the microenvironment surrounding residue 118, as seen from the unchanged conformations of the (truncated) catalytic triad, the S1 subsite residue His133, and the oxyanion hole (see Fig. 8b). The consistent conformations of these parts between the native and the mutant proteins make the mutant structure an appropriate model for enzyme–substrate interaction studies.

Discussion

PRRSV nsp4 has been predicted and accepted as the main proteinase, with a pivotal role in the viral life cycle, but experimental proof had been lacking. Our work, for the first time, reports the crystal structure and in vitro proteolytic activity of this enzyme.

The S1 specificity pocket that accommodates the side chain of the substrate’s P1 residue plays a vital role in substrate recognition and catalysis. In all the reported 3CLpro’s with known structures, a conserved histidine is found sitting at the bottom of the pocket.12,22,27–33 Mutational analysis found that any replacement of this residue both in human and feline coronavirus Mpro completely abolished their proteolytic activities.34,35 As a member of the 3C-like protease family, it is not surprising that in PRRSV nsp4, His133 is localized appropriately to fulfill this function. However, besides this conserved His, the S1 subsite of both SGPE and EAV nsp4 includes two additional Ser/Thr residues as shown in Fig. 4a,12,20 for which no counterparts can be found in our structure. We suspect that Ser136 must also be involved in P1 side-chain recognition based on two facts. First, this Ser is absolutely conserved in arterivirus nsp4 sequences. In EAV nsp4, the same Ser has been proposed to contact the side chain of the P1 glutamic acid (Fig. 4a). Furthermore, although we cannot define the position of Ser136 due to total lack of electron density, the two residues directly preceding it are clearly defined. Therefore, Ser136 is confined to a limited steric space close to the modeled P1 residue. A hydrogen bond between this disordered residue and the P1 amino acid could help stabilize both residues upon substrate binding. We also note that loop 136–140 lines one wall of the S1 subsite and is highly flexible. It may function like a lid that can partially cover the pocket after the entry of the P1 side chain and therefore help clinch the substrate to allow the occurrence of nucleophilic attack onto the scissile bond. This may also, to some extent, compensate the lack of a third partner in the S1 specificity pocket of PRRSV nsp4 (see Results).

Snijder et al. reported that mutation of this third residue, Thr115 in EAV nsp4, to other amino acids can have various effects on catalysis, indicating its important role as part of the S1 subsite.11

It is noteworthy that our enzyme also shows weak proteolytic activity against SP8-M with a Gln at the P1 position (Fig. 6e; Table 2). This might reflect the flexibility of the substrate binding of the enzyme that can also accommodate Gln in addition to Glu.

Fig. 8. The structure of the S118A mutant 3CLSP. (a) Stereo view of the electron density map of the mutant residue Ala118 as well as the other two residues of the catalytic triad. The 1.0 σ contoured map confirmed the successful substitution of Ser118 by Ala. (b) Comparison of the native structure (light blue) with that of the mutant protein (magenta). The catalytic triad is shown as sticks. His133 is also shown because of its crucial role in substrate recognition. The missing loop and the oxyanion hole region are also highlighted. The consistent conformations of these parts between the native and the mutant proteins make the mutant structure an appropriate model for enzyme–substrate interaction studies.
Recognition of both residues has been reported in other 3C proteinases, for example, 3C from members of the 
Picornaviridae. Taking the similarity of the side chains of Glu and Gln into account, this might not be surprising. Furthermore, the flexibility of Ser136 might also be attributed to this partial compatibility.

We also report a rare form of oxyanion hole in our structure, which is totally lacking the signature helix-like turn. In serine proteinases, the catalytic cleavage of the substrate peptide bond is assumed to undergo a transformation from the ground planar state to a transition state that is characterized by the formation of a tetrahedral intermediate. The backbone NH groups (in the case of subtilisin also including the side chain of Asn153) can hydrogen-bond with the resultant carboxyl oxyanion of the substrate P1 residue during cleavage at the scissile bond and are therefore very important in catalysis.

In EAV nsp4 and SARS-CoV M Pro, two forms of the oxyanion hole were observed, an active form and a collapsed one. Nevertheless, in both forms, the signature turn remains almost the same. In EAV nsp4, the different forms of the oxyanion hole are believed to arise mainly from the flipping of the peptide bond between Ser117 and Gly118, which is common in pairs of homologous protein structures. However, as shown in Fig. 4b, the stretched form observed in our structure (lacking the helix-like turn) can only support the formation of one hydrogen bond with the oxyanion to stabilize the negative charge. We suspect that the hydrophobic strength of Phe112 might be the main cause for this atypical oxyanion hole. This is supported by our recent mutational work of Phe112Ala. The mutant enzyme shows some higher enzymatic activity, implying the role that Phe112 played in stabilizing the negative charge.

Either addition of extra residues to the N-terminus or deletion of the N-terminal residues is deleterious to enzyme activity. There are also many reports showing that the homodimeric form of the SARS-CoV 3CL Pro is the catalytically active form and the unique C-terminal extra domain plays a key role in controlling the dimer–monomer equilibrium. In our recent report, we showed that the recombinant PRRSV nsp4 with nine extra amino acid residues at the N-terminus or deletion of the N-terminal residues is deleterious to enzyme activity. It is very likely that the homodimeric form of the SARS-CoV 3CL Pro is the catalytically active form and the unique C-terminal extra domain plays a key role in controlling the dimer–monomer equilibrium.

Materials and Methods

Construction of the recombinant plasmids

The construction of the recombinant plasmid encoding wild-type 3CLSP has been described previously. Briefly, the coding sequence of the PRRSV nsp4 was amplified by
Table 3. Primers used in this study

| Primer | Restriction site | Nucleic acid sequence (5’–3’)
|--------|------------------|-----------------------------|
| 3CL-F2 | Small            | cccgggGCCTTGAGAACACTAAAA     |
| 3CL-R2 | Xhol             | CGCagTTTCCAGTTGCGG            |
| 3CLM-F1| None             | TGGCCATGCTAGC                  |
| 3CLM-R1| None             | GGGATCC                        |
| 3CL-EA-F1| NdeI            | CGCctatagGGCCTTGAGAACACTAAAA  |
| 3CL-EA-R1| XhoI            | CGCgcatgGTGCTAGC                |
| 3CL-EA-R1| XhoI            | AAGTTGAACTGTGGAAAGGCTCCTTCCAGTTCGGGTTTGGC |

Primers used in this study. Noncapitalized letters indicate the sequence of the restriction site inserted. Bold letters indicate nucleotide changes introduced. Italic letters represent the sequence encoding the Myc epitope. Underlined letters represent the coding sequence of the eight N-terminal residues of nsp5.

reverse transcription-PCR reaction from the genomic RNA extracts of PRRSV isolate (GenBank accession no. EF112445) and then inserted into Smal and XhoI sites of pGEX-6P-1 to yield plasmid pGEX-3CL. The S118A (Ser118 substituted by Ala) mutant construct was achieved by overlapping extension PCR. Using pGEX-3CL as template, with primer combinations of 3CL-F2/3CLM-R1 or 3CLM-F1/3CL-R2, two segments were obtained first and then they were mixed together at equal molar ratio as new templates in a second round of PCR by primer pairs 3CL-F2/3CL-R2. The resultant fragment that carries the S118A substitution was inserted into pGEX-6P-1 to generate pGEX-3CLM-S118A.

With primer pairs 3CL-EA-F1/3CL-EA-R1, using either pGEX-3CL or pGEX-3CLM-S118A as template, DNA fragments encoding translational fusion proteins his-nsp4-[nsp4/nsp5]-Myc-his were amplified respectively. The fragments encoding translational fusion proteins his-nsp4-[nsp4/nsp5]-Myc-his and his-nsp4(S118A)-[nsp4/nsp5]-Myc-his were amplified respectively. The fragments were introduced into the pET-28b plasmid to create pET-3CL-EA and pET-3CLM-S118A-EA (Fig. 1b).

All the constructs were verified by direct DNA sequencing, and the sequences of the primers used in this study are listed in Table 3.

Protein purification and crystallization

The detailed production and purification procedure for the wild-type 3CLSP have been described previously. For the mutant enzyme, the same method was adopted. In brief, 1 μl of the plasmid was transformed into E. coli BL21 (DE3) competent cells for protein expression. The resultant GST fusion protein was purified by glutathione-Sepharose chromatography, cleaved with PreScission Protease, and the recombinant protein was further purified by size-exclusion chromatography.

SeMet-labeled 3CLSP was produced in BL21 (DE3) E. coli grown in SeMet minimal medium (0.65% YNB, 5% glucose, and 1 mM MgSO4 in M9 medium) supplemented with 1 μl of the plasmid was transformed into E. coli BL21 (DE3) E. coli grown in SeMet minimal medium (0.65% YNB, 5% glucose, and 1 mM MgSO4 in M9 medium) supplemented with 1 μl of tris at 60 mg L−1; lysine, threonine, and phenylalanine at 100 mg L−1; and leucine, isoleucine, and valine at 50 mg L−1. The expressed protein was purified as described above.

The purified proteins were concentrated, and their concentration was determined using BCA™ Protein Assay Kit (Pierce) according to the manufacturer’s instructions.

Crystallization conditions were optimized, and the best crystals for both the wild-type and SeMet-labeled 3CLSP were obtained by vapor diffusion in hanging drops consisting of 1 μl of reservoir solution (0.1 M sodium citrate, pH 5.3, and 0.7 M monobasic ammonium phosphate) and 1 μl of concentrated protein solution (12 mg ml−1 in 10 mM Tris–HCl and 10 mM NaCl, pH 8.0), followed by incubation at 18 °C for 7 days.

For the S118A mutant protein, the crystals were obtained under conditions similar to those for the wild-type 3CLSP, except that the pH value was elevated to 5.7 and the concentration of the mutant protein was 18 mg/ml. Crystals appeared after 10 days at 18 °C.

Data collection, phasing, and refinement

Crystals were flash-cooled in liquid nitrogen in cryoprotectant solution containing 15% glycerol and 50% reservoir solution. Multiwavelength X-ray diffraction data to 1.9 Å were collected from crystals of SeMet-labeled protein on beamline 3W1B of the Beijing Synchrotron Radiation Facility and processed with the HKL2000 suite of programs. Two selenium sites were located from data collected at peak and remote wave-lengths using SOLVE. An initial model was built using REEMAC5 followed by model refinement using REFMAC5 (CCP4 suite).

X-ray diffraction data collected from crystals of the native protein on an in-house Rigaku MicroMax007 rotating-anode X-ray generator were phased using MR procedures in MOLREP as implemented in CCP4 Version 6.02. The initially determined structure was used as a search model. A series of iterative cycles of manual rebuilding were carried out in Coot and refinement with REFMAC5 (CCP4 suite).

The S118A mutant structure was solved by MR method using the wild-type structure as the search model and refined to 2.0 Å.

The final statistics are summarized in Table 1. During the course of model building and refinement, the stereochemistry of the structures was checked by PROCHECK. All figures were generated using PyMOL and ESPript.

In vitro peptide trans cleavage assay

The 16-mer substrate peptides (representing eight putative cleavage sites in PRRSV pp1a or pp1ab) for the in vitro 3CL protease peptide cleavage assay were purchased from Beijing Scilight Biotechnology Ltd. Co. Each peptide was dissolved in 100% dimethyl sulfoxide at a concentration of 50 μM and stored at −80 °C. The nsp4 activity assay was performed in a reaction volume of 200 μl, using 5 μl nsp4 (or S118A mutant nsp4) (25 mg/ml) and 1 μl peptide (50 μM) in the reaction buffer (20 mM Tris, pH 7.4, 150 mM NaCl, 1 mM ethylenediaminetetraacetic acid, 1 mM DTT, and 10% glycerol) to yield a final concentration of 25 μM enzyme and 250 μM peptide in the assay. Cleavage reactions were routinely incubated at 37 °C for 15 h. The reactions were terminated by the addition of an equal volume of 2% trifluoroacetic acid, and samples were frozen in liquid nitrogen immediately.
performed in a buffer consisting of 20 mM Tris buffer, fluorescence spectrophotometer. The experiments were conducted in a buffer consisting of 20 mM Tris-HCl (pH 7.3), 100 mM NaCl, 1 mM ethylenediaminetetraacetic acid, and 5 mM DTT. Kinetic parameters \( k_{\text{cat}} \) and \( k_{\text{m}} \) were determined by initial-rate measurements at 25 °C. The reaction was initiated by adding proteinase (final concentration, 0.5 μM) to a solution containing different final concentrations of the fluorogenic peptide (10–40 μM). The initial rates were converted to enzyme activities (micro-molar substrate cleaved per second). Kinetic constants were derived by fitting the data to the Michaelis–Menten equation with the nonlinear regression analysis program Sigma Plot.

**Analysis of the cis proteolytic activity of 3CLSP**

To investigate the cis cleavage activity of 3CLSP, we designed two constructs bearing the nsp4/nsp5 scissile bond fused to a Myc epitope at the C-terminus of 3CLSP or 3CLSP (S118A), resulting in the expression vectors pET-3CL-EA and pET-3CL-S118A-EA. The N-terminal and C-terminal hexa-His tags were introduced into the recombinant proteins via vector pET-28B (1b). The recombinant proteins were expressed and purified by using a Ni-NTA affinity chromatography column and a Superdex 75 HiLoad 16/60 column (GE Healthcare). The uninduced cell sample, cell lysate after induction, and the purified protein were separated by 12% SDS-PAGE for Coomassie brilliant blue staining.

For Western blot analysis, 2 or 1 μg of the purified his-nsp4-[nsp4/nsp5]-Myc-his and his-nsp4(S118A)-[nsp4/nsp5]-Myc-his were loaded onto 12% SDS-PAGE for separation and then transferred onto Hybond nitrocellulose membranes (Amersham Biosciences). Membranes were blocked overnight in 10% milk powder dissolved in Tris-buffered saline. Blotted proteins were detected with the rabbit polyclonal anti-His primary antibody (Santa Cruz) or the mouse monoclonal anti-Myc antibody (Santa Cruz). A horseradish-peroxidase-labeled anti-rabbit or anti-mouse secondary antibody (Santa Cruz) was used. After each incubation step, membranes were washed four times with 0.05% Tween-20 in Tris-buffered saline. Detection was performed by using SuperSignal West Pico Chemiluminescent substrate solution (Pierce Biotechnology).

**C-terminal sequencing of his-nsp4-[nsp4/nsp5]-Myc-his protein**

The protein was prepared in 50 mM NH₄HCO₃ at a concentration of 1 mg/ml followed by digestion using trypsin (Roche) at 37 °C for 16 h with the mass ratio of trypsin to the protein set at 1:30. The digested products were further reduced by the addition of DTT to the digestion volume at 10 mM followed by incubation at 56 °C for 1 h. The resultant products were analyzed by electrospray ionization (ESI)-MS/MS. Then, a peptide fragment of 2117.06 Da was chosen for sequencing by ESI-MS/MS on a Q-TOF2 hybrid quadrupole/time-of-flight mass spectrometer (Micromass, UK).

**PDB accession numbers**

The coordinates and associated structure factors have been deposited in the Research Collaboratory for Structural Bioinformatics PDB (PDB code 3FAN for the native protein; PDB code 3FAO for the S118A mutant structure).

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