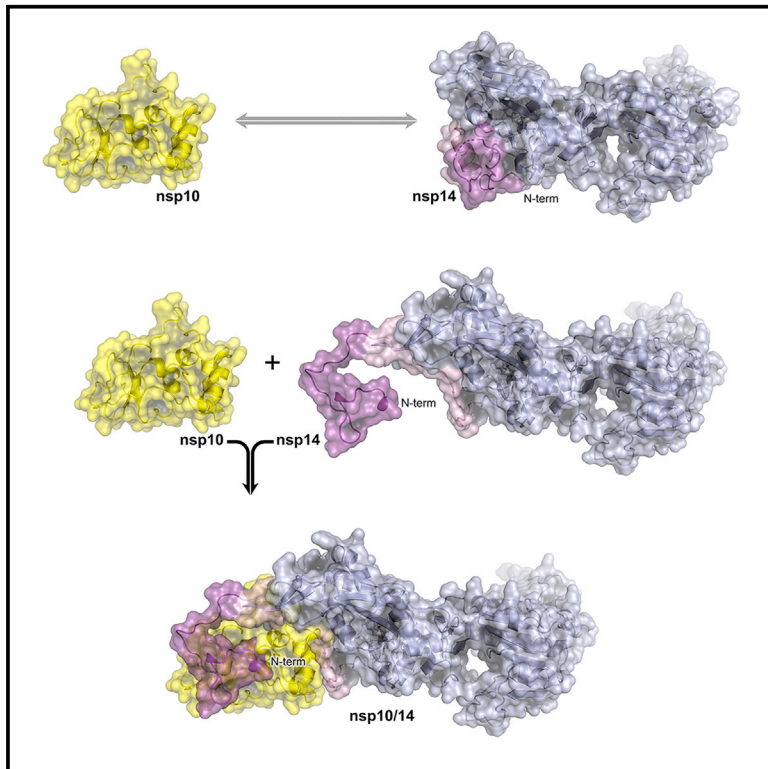


Structure

Refolding of lid subdomain of SARS-CoV-2 nsp14 upon nsp10 interaction releases exonuclease activity

Graphical abstract



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In brief

Czarna et al. use X-ray crystallography to determine the structure of SARS-CoV-2 nsp14 methyltransferase with the bound co-factor SAH. The N-terminal “lid” is undefined, revealing a high flexibility of the nsp14 in the absence of its partner nsp10.

Highlights

- Structure of SARS-CoV-2 nsp14 in absence of nsp10 is presented
- Refolding of the “lid” domain upon nsp10 binding promotes exonuclease activity
- Structure facilitates the development of inhibitors for SARS-CoV-2 methyltransferase



Article

Refolding of lid subdomain of SARS-CoV-2 nsp14 upon nsp10 interaction releases exonuclease activity

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SUMMARY

During RNA replication, coronaviruses require proofreading to maintain the integrity of their large genomes. Nsp14 associates with viral polymerase complex to excise the mismatched nucleotides. Aside from the exonuclease activity, nsp14 methyltransferase domain mediates cap methylation, facilitating translation initiation and protecting viral RNA from recognition by the innate immune sensors. The nsp14 exonuclease activity is modulated by a protein co-factor nsp10. While the nsp10/nsp14 complex structure is available, the mechanistic basis for nsp10-mediated modulation remains unclear in the absence of the nsp14 structure. Here, we provide a crystal structure of nsp14 in an apo-form. Comparative analysis of the apo- and nsp10-bound structures explain the modulatory role of the co-factor protein and reveal the allosteric nsp14 control mechanism essential for drug discovery. Further, the flexibility of the N-terminal lid of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) nsp14 structure presented in this study rationalizes the recently proposed idea of nsp14/nsp10/nsp16 ternary complex.

INTRODUCTION

Upon release into the cytoplasm of the host cell, the coronavirus genome is translated into a single non-functional polyprotein 1a/1ab. Its autoproteolytic processing releases a number of nonstructural proteins (nsps) responsible for viral replication. The activity of a particular nsp is relatively well characterized, but the mechanistic understanding of their interplay is still insufficient (Pyrc et al., 2004; Yadav et al., 2021; Sola et al., 2015).

Nsp14 consists of two domains, each with a distinct catalytic activity. The C-terminal domain is a SAM-dependent methyltransferase, while the N-terminal ExoN domain exhibits 3'-5' exonuclease activity. Recognition of unmodified RNA molecules constitutes a universal antiviral strategy embedded in the host innate immune defenses. To avoid detection, coronaviral RNA is capped by viral enzymes, which additionally increase its stabil-

ity and facilitates translation initiation. Capping involves a series of enzymatic reactions. In the penultimate step, the C-terminal domain of nsp14 catalyzes N7-methylation of a GpppA intermediate. Subsequent methylation by nsp16 results in a functional cap (⁷MeGpppA_{2'OMe}) (Sevajol et al., 2014; Bouvet et al., 2010, 2012; Eckerle et al., 2010; Yan et al., 2021; Chen et al., 2009).

The large coronaviral genome requires replication fidelity to maintain its functionality. While the nsp12-centered polymerase complex is highly processive, it is also error prone. An unaudited replication would result in an excessive accumulation of mutations and the generation of defective progeny virions. Nsp14 associates with the replication complex and removes incorrectly incorporated nucleotides from the 3' end of the newly formed RNA strand. Nsp14 is associated with nsp10, a protein co-factor that modulates nsp14 exonuclease activity in the replication complex (Smith and Denison, 2013; Eckerle et al., 2010).



Table 1. Data collection and refinement statistics

Data collection statistics	
PDB ID	7R2V
Wavelength (Å)	1.00004
Resolution range	86.59–2.53 (2.89–2.53)
Space group	P 1 21 1
Unit cell	67.18, 100.31, 90.87 (Å) 90, 107.66, 90 (°)
Data-scaling software	autoPROC, AIMLESS, (STARANISO)
Unique reflections	19,046 (5,869)
Rmeas (%)	18.8 (99.9)
Rpim (%)	8.7 (40.0)
Multiplicity	5 (6.2)
CC1/2	0.989 (0.664)
Mean I/σ(I)	6.8 (1.6)
Completeness	
Spherical (%)	99.8 (7.7)
Ellipsoidal (%)	99.8 (65)
Refinement	
Rwork/Rfree/test set (%)	0.196/0.254/4.97
Wilson B factor (Å)	57.062
Clashscore	3
Raachandran outliers (%)	0.5
Sidechain outliers (%)	5.1
RSRZ outliers (%)	3.4

Values provided in parentheses are for the highest resolution shell. The $I/\sigma(I) > 1.2$ was automatically used by the STARANISO server for the highest resolution shell limit. The data was further limited based on analysis of density maps. Rmeas, multiplicity-independent R-factor introduced by Diederichs that indicates the real precision of the measurement, independent of the multiplicity of the reflection; Rpim, precision of the averaged merged intensity measurements; Rwork, working R-factor of the refined structure; Rfree, a residual function calculated during structure refinement in the same way as the conventional R factor, but applied to a small subset of reflections that are not used in the refinement of the structural model. The purpose is to monitor the progress of refinement and to check that the R factor is not being artificially reduced by the introduction of too many parameters; RSRZ, the real-space R-value (RSR) is a measure of the quality of fit between a part of an atomic model (in this case, one residue) and the data in real space. RSR Z-score (RSRZ) is a normalisation of RSR specific to a residue type and a resolution bin.

The structural and biochemical basis of nsp14 interaction with nsp10 has been elucidated (Baddock et al., 2022; Liu et al., 2021; Lin et al., 2021). We have recently demonstrated that nsp14, nsp10, and nsp16 form a ternary complex, further modulating nsp14 catalytic activity (Matsuda et al., 2022). Despite the detailed structural characterization of the complexes, the mechanistic basis of the modulation of nsp14 exonuclease activity remains elusive. The absence of structural information on nsp14 alone hinders the mechanistic understanding of the consequences of binding with nsp10 and nsp10/nsp16.

Here, we provide the crystal structure of nsp14 in the apo-form (without nsp10), which, together with prior data on nsp10/nsp14 complex, provide insight into the modulatory role of nsp10 on nsp14 exonuclease activity by a set of complex allosteric modi-

fications. Furthermore, the presented structure strongly supports the nsp14/nsp10/nsp16 heterotrimer formation model.

RESULT

Full-length severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) nsp14 exonuclease/methyltransferase was expressed in *Escherichia coli* and purified. The crystal structure of S-adenosylhomocysteine (SAH)-bound nsp14 was determined at 2.5 Å resolution (Table 1). The crystals belonged to P2₁ space group and contained 2 molecules in the asymmetric unit. Nsp14 is characterized by a bimodular structure corresponding to the two distinct catalytic activities (Figure 1A). The C-terminal methyltransferase domain of nsp14 (amino acids 287–524) is well defined by the electron density for most of its part save for amino acids 454–465. At the same time, the N-terminal exonuclease domain (encompassing amino acids 1–286) is largely undefined by the electron density, indicating significant flexibility (Figure S1). The flexible regions are between amino acids 1–24, 40–44, 93–103, and 123–154 and are of importance for further discussion. The undefined parts are identical in both chains.

The methyltransferase domain of nsp14 is characterized by a fold atypical to methyltransferases. The structure is built around a central β sheet composed of 5, instead of 7, canonical β strands. Further, the methyltransferase domain of nsp14 contains an atypical peripheral zinc finger stabilized by another element that distinguishes nsp14 from the typical methyltransferase fold, a supplementary C-terminal α-helix. The methyltransferase fold within our structure closely resembles that previously seen in the nsp10/nsp14 complex of SARS-CoV-1 (PDB: 5C8S) with a root-mean-square deviation (RMSD) of 0.57 Å for 197 equiv C_α atoms. A reaction product, SAH, is clearly defined by the electron density in our structure (Figure 1B). The remaining part of the active site has an empty, tubular shape (the methylated reaction product is absent). Interestingly, the active site faces a large, unobstructed water channel. With unobstructed access to the active site, our structure is particularly valuable in terms of inhibitor development (fragment screening and inhibitor soaking).

The SAM/SAH-binding region in our structure (331-DxGxPxA-337) diverges from the classical methyltransferase motif I (E/DxGxGxG; Martin and Mcmillan, 2002) but is consistent with that described in nsp14 of SARS-CoV-1 (Ma et al., 2015). The density in the MTase active site unambiguously shows SAH despite the fact that SAM was added to the protein solution (Figure S2). The purine ring of SAH is positioned via sulfur/π interaction with Cys387; hydrophobic interactions contributed by side chains of Ala353, Val389, Phe367, and direct hydrogen bonds contributed by main chain of Tyr368. A water-mediated hydrogen bond contributed by Asn388 is additionally observed in model B. Ribose is stabilized by hydrogen bonds contributed by the side chain of Asp352. Mainchain carbonyl oxygens of Gly333 and Trp385 contribute hydrogen bonds with the cysteine amine, whereas its carboxyl group forms a salt bridge with Arg310. The identified SAH positioning residues are consistent with the conclusions of prior mutagenesis studies (Ogando et al., 2021). The majority of reported nanomolar MTase inhibitors are SAM competitive (Devkota et al., 2021). By delineating

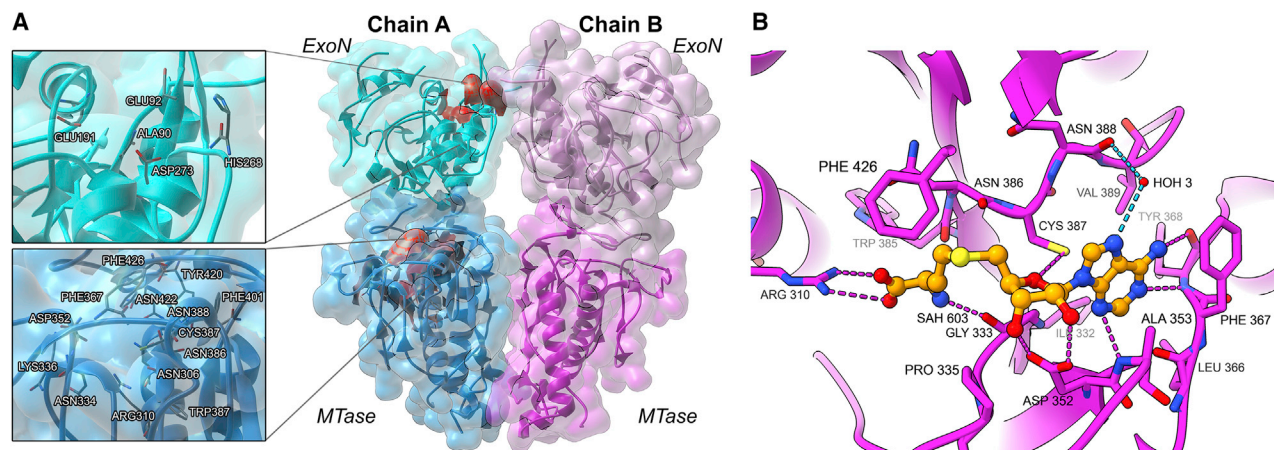


Figure 1. Crystal structure of apo-nsp14

(A) Arrangement of the molecules in the asymmetric unit. Model (A) blue and model (B) pink. MTase and ExoN domains within the bimodular structure of nsp14 are distinguished by a primary color shade. Active-sites residues are indicated in red in one molecule, and the detailed arrangements of the active sites are shown in inserts.

(B) Interactions of SAH (orange, stick model) at the active site of nsp14 MTase domain (pink, ribbon). Hydrogen bonds are depicted as dotted lines; waters are indicated as red spheres.

See also Figure S1.

the detailed molecular interactions in the SAH pocket, our study facilitates the development of SAH mimetic inhibitors of SARS-CoV-2 MTase.

A structure of nsp14 from SARS-CoV-2 has been recently deposited at the Protein Data Bank (PDB: 7QGI), which contains an empty active site (no substrates or reaction products). Comparison with our structure demonstrates that the SAM/SAH binding site in 7QGI is occluded by a loop connecting $\beta 2'$ and $\beta 3'$ strands of a central β sheet (Figure S3). The place where the loop in question is located in SAH-bound structures (nsp10/nsp14 complex from SARS-CoV-1 and this study) is occluded by an adjacent molecule in 7QGI. This may suggest that either the blocked active site in 7QGI is an artifact of crystal packing or, alternatively, it may represent one of the multiple forms found in solution selected by the particular crystal packing. When Ma et al. (2015) analyzed SARS-CoV-1 nsp10/nsp14 complexes in SAH-bound and empty forms, the loop in question had the same conformation in both SAH-bound and empty structures (and comparable to that found in our structure), suggesting indirectly that the open conformation is the prevalent form.

The exonuclease domain of nsp14 is built around a five-stranded, twisted β sheet flanked by α helices. Such an arrangement of the core structure elements follows the overall design of DEDD family exonucleases but contains unique features: the N-terminal region (1–84) and the first zinc finger. The striking features of the exonuclease domain in our structure became evident only upon comparison with the structure of nsp10/nsp14 complex. The N terminus of nsp14 spanning amino acids 1–70 is folded and oriented differently in compared structures (Figures 2A and 2B). The N terminus is well defined by electron density in the nsp10/nsp14 complex but contains few ordered secondary structures. Amino acids 1–56 constitute the first of the two interaction regions with nsp10 (nomenclature of structural features after Ma et al. (2015)). The second interaction region comprises residues 60–67 and 200–202. In apo-nsp14

structure, residues 1–24 are not defined by the electron density. Region 25–70 is completely refolded in the apo-nsp14 structure compared with the nsp10/nsp14 complex. For example, residues Thr50–Met58 form an α helix in the apo-nsp14 structure, whereas residues 51–54 form a $\beta 1$ strand within $\beta 1$, $\beta 5$, and $\beta 6$ sheets in the nsp10/nsp14 complex. The remaining residues (25–49, 59–67) contribute completely different intermolecular interactions within the apo-nsp14 structure compared with the nsp10/nsp14 complex. In the absence of nsp10, the N-terminal region of nsp14 covers the second nsp10 interaction region, burying, among others, residues 200–202. Thereby, the N terminus of nsp14 acts as a lid, covering the nsp10 binding site in the absence of the protein co-factor. The lid rearranges its structure in the presence of nsp10 to create a significant interaction surface with nsp10. The above conclusion is corroborated by comparing our structure and 7QGI, which show similar arrangements of lid domains irrespective of crystal packing and different from that found in nsp10/nsp14 complex.

Nsp10 enhances the 3' exonucleolytic activity of nsp14 but does not significantly affect the methyltransferase activity. Nsp10's effect on nsp14 was rationalized by the nsp10/nsp14 crystal structure, where nsp10 contributes significant contacts with the ExoN domain of nsp14 but does not interact with the N7-methyltransferase domain (Ma et al., 2015). The extensive surface area of interaction suggested that nsp10 maintains the structural stability of ExoN, but the nature and extent of instability of apo-nsp14 were not known in the absence of the apo-nsp14 structure. In the current study, we show that the ExoN domain of nsp14 is partially unfolded in the absence of nsp10. The proximal part of the $\beta 2$ - $\beta 3$ sheet adjacent to catalytic Glu92 is undefined by electron density (flexible, unstructured) in apo-nsp14. The same is true for the entire region between $\alpha 3$ and $\beta 4$. Residues constituting $\alpha 2$ and $\beta 5$ - $\beta 6$ in the nsp10/nsp14 complex structure are not defined by the electron density in the apo-nsp14 structure. Significantly, in apo-nsp14, the

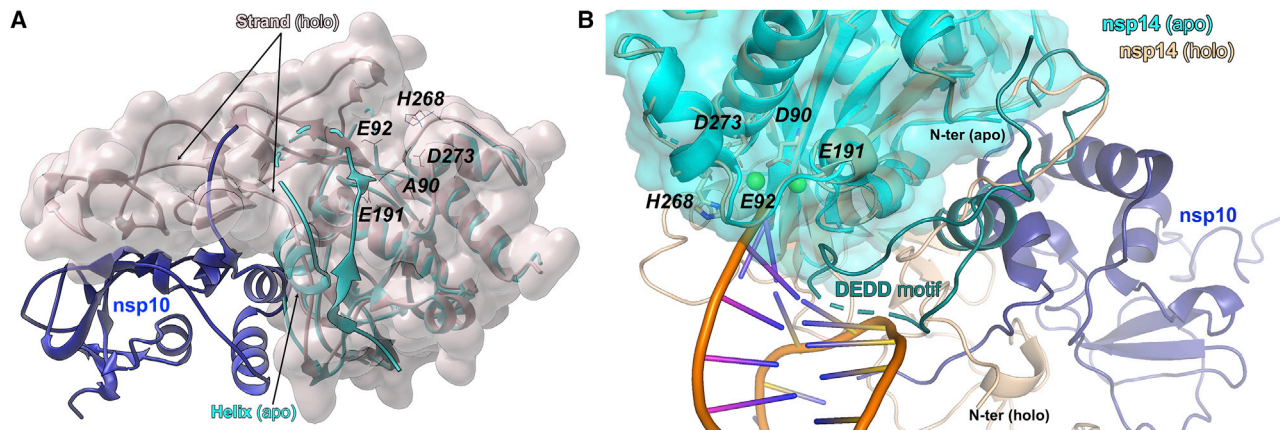


Figure 2. Nsp10 induced refolding of nsp14 ExoN lid subdomain

(A) Nsp10 (navy blue, ribbon)/Nsp14 (gray, surface) complex overlaid on apo-nsp14 (cyan). The N-terminal lid of nsp14 refolds upon binding of nsp10 as exemplified by Thr50–Met58 region assuming a helical structure in apo-nsp14 and forming a strand of a β sheet indicated in nsp10/nsp14 complex. In the apo structure, the lid occludes nsp10-binding site at the surface of nsp14.

(B) The RNA interaction site of nsp10 (navy blue)/nsp14 (beige) complex (7N0B) is sterically occluded by lid subdomain in apo-nsp14 structure (cyan), providing a rationale for the negligible exonuclease activity of nsp14 in the absence of nsp10. RNA (orange). The apo-nsp14 loop, occluding the RNA-binding site, is denoted in dark cyan (dotted line).

See also Figures S2 and S3.

catalytic metal ion-binding site within the DEDD motif is partially occluded by the side chain of His188. Further, the RNA-binding site is partially occluded by the lid fragment (Thr35–Lys47), which additionally buries the nsp10-binding site. It has been shown that Lys9 of the nsp14 is critical for exonuclease activity (Moeller et al., 2022). This residue is not visible in our electron density. Thus, the refolded N-terminal part of the protein is not structurally oriented for RNA interaction. This further confirms the role of 1–50 span refolding in exonuclease activity control. Nsp10-induced refolding of the lid removes the steric occlusion and induces a catalytically capable conformation of nsp14 observed in the nsp10/nsp14 complex (Figure 2B).

We have demonstrated earlier that nsp14, nsp10, and nsp16 form a ternary complex centered around nsp10 (Matsuda et al., 2022). However, such complex is not supported by the structural data available on nsp10/nsp14 and nsp10/nsp16 binary complexes. When the binary complexes are overlaid on nsp10, the lid of nsp14 clashes with nsp16, demonstrating that ternary-complex formation would require a structural rearrangement within the lid. The current study demonstrates that the lid of nsp14 is capable of structural morphing, indirectly supporting the mechanism of triplex formation.

DISCUSSION

The current pandemic highlighted the importance of understanding coronavirus physiology. Prior research on other human and animal coronaviruses facilitated an unprecedented pace of vaccine and antiviral development, but the constant genetic drift and the threat of new zoonotic transmission warrant continued effort.

Nsp14 constitutes a valid target in SARS-CoV-2, but the structural information was incomplete. This study provided the structural basis for a mechanistic explanation of nsp10-mediated modulation of nsp14 3' exonuclease activity, defined the structure of nsp14 N7-methyltransferase in the absence of RNA sub-

strate, and offered crystallization conditions suitable for soaking of small molecules at the active site of nsp14 N7-methyltransferase. Thereby, the study contributes to our understanding of coronavirus physiology and facilitates the effort of anti-coronavirus drug development.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.str.2022.04.014>.

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AUTHOR CONTRIBUTIONS

Conceptualization, A.C. and G.D.; methodology, L.K., A.M., and A.K.; validation, P.G.W., G.D., and K.P.; investigation, J.P., M.P., and P.W.; visualization A.C., A.M., C.R., I.G. and G.P.; writing – original draft, G.D. and K.P.; writing – review & editing, A.C., J.P., C.R., I.G., S.O., and F.C.; formal analysis, J.P. and A.C.; project administration, A.C. and P.G.W., supervision, K.P.; funding acquisition, K.P. and G.P.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Bacterial and virus strains		
<i>E. coli</i> BL21 Rosetta2 (DE3)	Milipore	71402
Chemicals, peptides, and recombinant proteins		
Terrific Broth medium	Bioshop	TER409
Ampicillin	Sigma-Aldrich	A9393
isopropyl-D-1-thiogalactopyranoside	Sigma-Aldrich	I5502
Tris-HCl	Bioshop	TRS003
NaCl	Bioshop	SOD004
MgCl ₂	Bioshop	MAG520
Glycerol	Bioshop	GLY004
β-mercaptoethanol	Sigma-Aldrich	M6250
Imidazole	Bioshop	IMD508
protease inhibitors cocktail	Roche	11697498001
Benzonase	Milipore	101654 SAFC
tobacco etch virus (TEV) protease	In-house	n.a.
S-Adenosyl methionine	New England BioLabs	B9003S
Deposited data		
Structure of nsp14 from SARS-CoV-2 in complex with SAH	this paper	PDB ID: 7R2V
SARS-CoV-1 nsp14	(Ma et al., 2015)	PDB ID: 5C8S
SARS-CoV-2 nsp14	(Imprachim et al., 2022)	PDB ID: 7QGI
SARS-CoV-2 nsp10-nsp14 (WT)-RNA complex	(Liu et al., 2021)	PDB ID: 7N0B
Recombinant DNA		
Uniprot P0DTD1 encoding amino acids 5926 – 6452	GeneArt	n.a.
pETDuet-1	Milipore	71146
Software and algorithms		
The STARANISO Server	Global Phasing Limited	https://staraniso.globalphasing.org/cgi-bin/staraniso.cgi
CCP4 v7.1	(Winn et al., 2011)	https://www.ccp4.ac.uk
CheckMyBlob server	(Kowiel et al., 2018)	https://checkmyblob.bioreproducibility.org
PyMol Molecular Graphics	System Schrodinger, LLC	https://pymol.org/2/

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact: K.P.: k.p.pyrc@uj.edu.pl

Materials availability

Plasmids produced in this study are available from the authors.

Data and code availability

- The atomic coordinates and structure factors of SARS-CoV-2 nsp14 have been deposited to PDB and are publicly available as of the date of publication. Accession numbers and DOI are listed in the [key resources table](#) as well as in [Table 1](#).
- The paper does not report original code.
- Any additional information required to reanalyze the data reported in this paper is available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Recombinant SARS-CoV-2 nsp14 was purified from the expression in *E. coli* BL21 Rosetta2 (DE3) strain.

METHOD DETAILS

Expression construct

Sequence encoding amino acids 5926 – 6452 of SARS CoV-2 polyprotein 1ab (Uniprot P0DTD1), including D90A and D92A substitutions (Bouvet et al., 2012), flanked by BamHI and NotI cleavage sites and containing a TEV protease cleavage site was codon-optimized for expression in *E. coli* and obtained by synthesis at GeneArt (sequence available in Table S1). The sequence was subcloned into pETDuet-1 expression vector.

Protein expression and purification

The plasmid was transformed into *E. coli* BL21 Rosetta2 (DE3) strain. Expression was carried out in Terrific Broth medium (Bioshop) supplemented with 100 µg/mL ampicillin (Sigma-Aldrich). Bacterial cultures were incubated at 37°C until the OD₆₀₀ value reached 1.2–1.4. The cultures were cooled to 4°C for 20 min, and the recombinant protein expression was induced with 0.5 mM isopropyl-D-1-thiogalactopyranoside (IPTG; Sigma-Aldrich). The bacteria were incubated at 18°C for additional 18 h. Bacterial cells were harvested by centrifugation at 7,000 × g for 30 min at 4°C and stored at –20 °C before the purification.

Bacterial pellets were resuspended in lysis buffer (50 mM Tris-HCl pH 8.5, 500 mM NaCl, 5 mM MgCl₂, 10% v/v glycerol, 5 mM β-mercaptoethanol, 10 mM imidazole), supplemented with benzonase (Millipore) and protease inhibitors cocktail (Roche), and lysed by sonication at 80% amplitude for 20 min at 14°C with 3 s on/2 s off cycles. The fraction of the supernatant containing soluble protein was separated from debris by centrifugation at 25,000 × g for 40 min at 4 °C. The clarified supernatant was incubated overnight at 4 °C with HisPur Ni-NTA Resin (Thermo Scientific) pre-equilibrated with the lysis buffer. Resin was sequentially washed with lysis buffer, then with wash buffer 1 (50 mM Tris-HCl pH 8.5, 500 mM NaCl, 5 mM MgCl₂, 10% v/v glycerol, 5 mM β-mercaptoethanol, 20 mM imidazole) and wash buffer 2 (50 mM Tris-HCl pH 8.5, 500 mM NaCl, 5 mM MgCl₂, 10% v/v glycerol, 5 mM β-mercaptoethanol, 30 mM imidazole). The immobilized protein was eluted with elution buffer (50 mM Tris-HCl pH 8.5, 250 mM NaCl, 5 mM MgCl₂, 10% v/v glycerol, 5 mM β-mercaptoethanol, 250 mM imidazole). Fractions containing the protein of interest (as determined by SDS-PAGE) were pooled and supplemented with tobacco etch virus (TEV) protease in a 1:10 molar ratio. TEV cleavage was combined with overnight dialysis into a cleavage buffer (50 mM Tris-HCl pH 8.5, 200 mM NaCl, 5 mM MgCl₂, 10% v/v glycerol, 5 mM β-mercaptoethanol, 10 mM imidazole). Reverse chromatography on HisPur Ni-NTA resin pre-equilibrated with the dialysis buffer allowed the cleaved His-tag removal. The protein was further purified by size exclusion chromatography on HiLoad 26/600 Superdex 200 prep grade column (GE Healthcare) equilibrated with the working buffer (50 mM Tris-HCl pH 8.5, 150 mM NaCl, 5 mM MgCl₂, 1 mM TCEP). The purity of the sample was assessed by SDS-PAGE.

Crystallization and data collection

The purified protein was concentrated to 11.5 mg/mL. The sample was supplemented with 2mM SAM. Crystallization screening was performed at 4°C using the sitting drop vapour diffusion method. Crystals were obtained from a solution containing 15% v/v 2-propanol, 0.2 M imidazole pH 7.6 and 500 mM polypropylene glycol 400 (JBScreen Plus HTS additive screen; Jena Bioscience) and 1:1 protein to mother liquor drops. Further optimization was performed around initial conditions where the crystals generally appeared after 4 days. The crystals were cryoprotected in 30% ethylene glycol in the mother liquor and flash-cooled in liquid nitrogen. The diffraction data were collected at Swiss Light Source (SLS, Villigen, Switzerland) beamline X06DA - PXIII.

QUANTIFICATION AND STATISTICAL ANALYSIS

The data were indexed and integrated using XDS (Kabsch, 2010). The data were scaled and merged using STARANISO webserver. A molecular replacement solution was found using Phaser (Mccoy, 2007) and the structure of SARS-CoV-1 nsp14 (PDB code: 5C8S) as a search model. SAH was identified in the electron density map using CheckMyBlob server (<https://checkmyblob.bioreproducibility.org>). Restraints were obtained from ligand builder in Coot (Emsley, 2017). The initial model was subjected to several iterations of manual and automated refinement cycles using COOT and REFMAC5, respectively (Emsley and Cowtan, 2004; Murshudov et al., 2011). Throughout the refinement, 5% of the reflections were used for cross-validation analysis, and the behaviour of R_{free} was employed to monitor the refinement strategy. The data collection and refinement statistics are shown in Table 1.

Structure, Volume 30

Supplemental Information

**Refolding of lid subdomain
of SARS-CoV-2 nsp14 upon nsp10
interaction releases exonuclease activity**

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Supplemental information

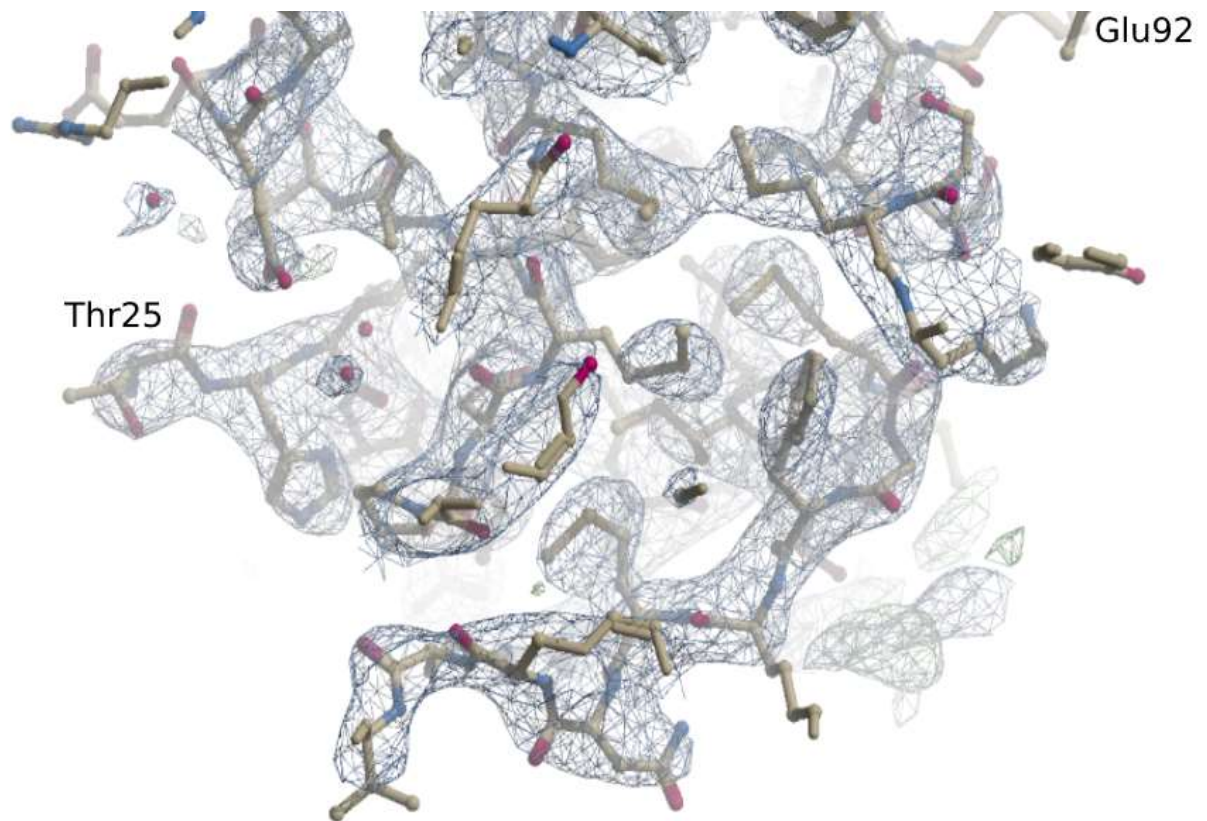


Figure S1: Related to Figure 1. Overview of the electron density defining the SARS-CoV-2 nsp14 N terminal lid region. The 2F_o-F_c map (blue mesh) is contoured at 1σ. The final model is depicted in stick representation (beige). The density allows for unambiguous determination of residues from 25 to 39 and 45-92. Other residues of the lid region were poorly defined or undefined by the electron density and were not included in the model.

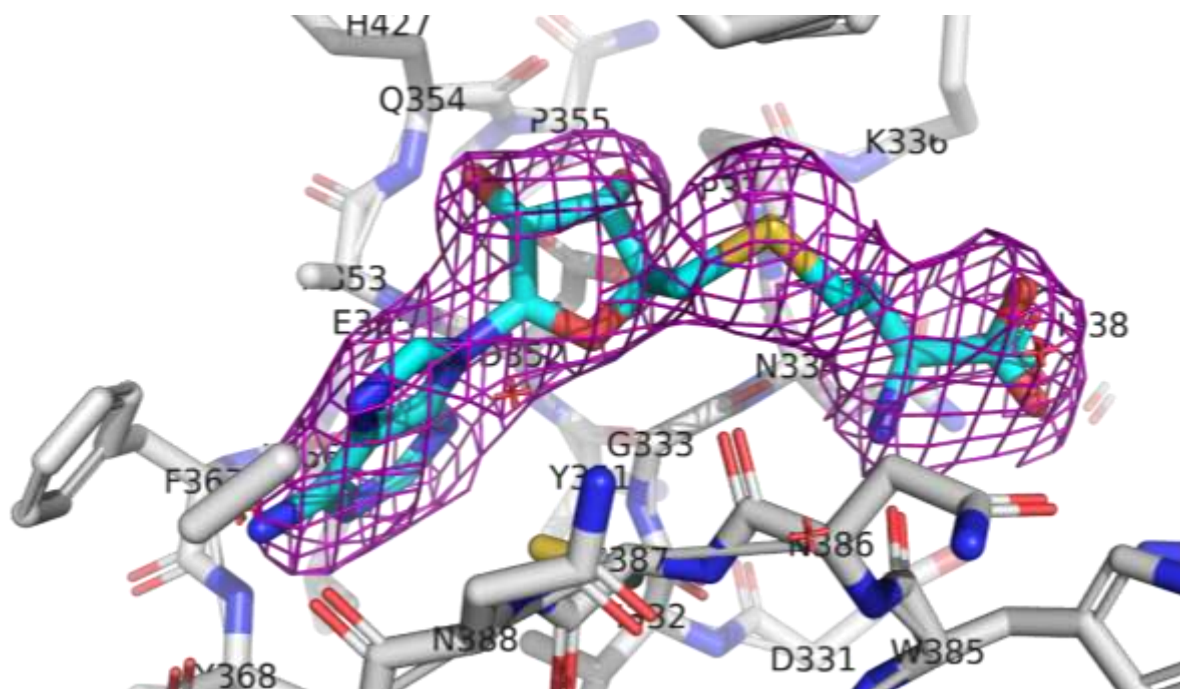


Figure S2: Related to Figure 2. Overview of the SAM/SAH binding site. The bound cofactor (SAH) is shown in stick model (blue). The electron density describing the cofactor is shown as the omit map ($2F_o - F_c$) with contour level 1σ (magenta mesh). No density describing the methyl moiety at sulphur atom is evident allowing to identify SAH (and not SAM) as a cofactor. Protein model shown in stick representation (gray).

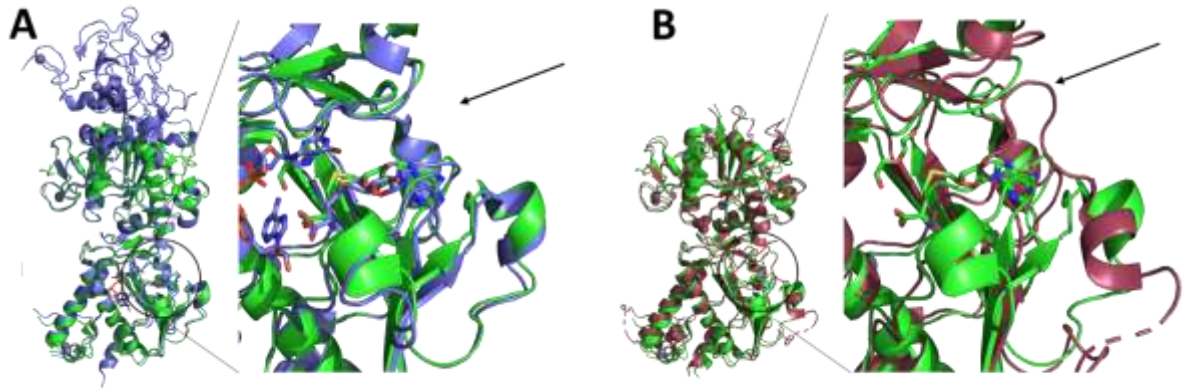


Figure S3: Related to Figure 2. Overlay of selected nsp14 structures. (A) Overlay of SARS-CoV-2 nsp14 structure containing SAH (green; this study) and SARS-CoV-1 nsp14/10 complex containing SAH and GpppA (violet; PDB ID:5C8S). No occlusion of the active site is seen. **(B)** Overlay of SARS-CoV-2 nsp14 structure containing SAH (green; this study) and SARS-CoV-2 nsp14 with no cofactor (brown; PDB ID: 7QGI). Significant differences in the orientations of loop connecting $\beta 2'$ and $\beta 3'$ strands blocking the active site in the structure not containing the cofactor. **(A, B)** Insets show close-ups of SAM binding pocket (indicated by an arrow). The structures of PDB ID: 7R2V and PDB ID:5C8S overlap ideally with RMSD of 0.802 for 362 atoms showing no occlusion to the active site for both nsp10-free and nsp10-bound structures (green and violet structures in S2 A, respectively). The structures of PDB ID: 7R2V and PDB ID:7QGI have a significant difference with a loop connecting $\beta 2'$ and $\beta 3'$ strands blocking the active site (indicated with the arrow) and RMSD of 1.440 for 399 atoms.

Table S1. Related to STAR★Methods. Nucleotide and amino acid sequences of SARS-CoV-2 nsp14 used in this study.

Nucleotide sequence:
(ggc)atggctgaaaatgtaacaggactctttaaagattgtagtaaggtaatcactgggttacatcctacacaggcacctacacac ctcagtgttgactaaattcaaaactgaaggttatgtgttgacatacctggcatacctaaggacatgacctatagaagactcatc tctatgatgggttttaaaatgaattatcaagtaatggttaccctaacatgtttatcaccgcgaagaagctataagacatgtacgtg catggattggcttcgctgtagctgggtgctatgctactagagaagctgttggtaccaattacctttacagctaggttttctacaggt gttaacctagttgctgacctacaggttatgttgatacacctaataatacagattttccagagttagtgtctaaaccaccgctggag atcaattaaacacctcataccacttatgtacaaaggacttccttggatgtagtgcgtataaagattgtacaaatgtaagtgaca cacttaaaatctctctgacagagtcgtatgtcttatgggcacatggctttgagttgacatctatgaagtattttgtgaaaatagg acctgagcgcacctgttgctatgtgatagcgtgccacatgctttccactgcttcagacacttatgcctgttggcatcattctattg gatttgattacgtctataatccgtttatgattgatgtcaacaatggggtttacaggtaacctacaagcaacctgatctgtattgt caagccatggtaatgacatgtagctagttgtgatgcaatcatgactaggtgtctagctgtccacgagtgctttgtaagcgtgtg actggactattgaatacctataattgggtgatgaactgaagattaatcgggctttagaaaaggttcaacacatggttgttaaagctg cattattagcagacaaattcccagttcttcagacattggtaaccctaaagctattaagtgtgtacctcaagctgatgtagaatgga agttctatgatgcacagcctgtagtgacaaagctataaaatagaagaattattctattcttatgccacacattctgacaaattcac agatgggtgatgcctatgttgaattgcaatgctgatagatatctgctaattccattgtttgtagattgacactagagtgctatcta acctaaactgcctgggttgatgggtggcagtttgatgtaataaacatgcattccacacaccagctttgataaaagtgctttgtt aatttaaaacaattaccattttctattactctgacagtcctatgtgagtcctatggaaaacaagtagtgcagatatagattatgtac cactaaagtctgctacgtgtataacacgttgcaatttaggtggctgtctgtagacatcatgctaagtagtacagattgtatctcga tgcttataacatgatgatctcagctggctttagcttgggtttacaacaatttgatactataaccttggaaacacttttacaagac ttcag
Amino acid sequence
(G)MAENV TGLFKDCSKVITGLHPTQAPTHLSVDTKFKTEGLCVDIPGIPKDM TYRRLISMMGFKMNYQ VNGYPNMFITREEAIRHVRAWIGFAVAGCHATREAVGTNLPLQLGFSTGVNLVAVPTGYVDTPNNTDF SRVSAKPPPGDQFKHLIPLMYKGLPWNVVRKIVQMLS DTLKNLS DRVVFVLWAHGFELTSMKYFVKIG PERTCCLDRRATCFSTASD TYACWHHSIGFDYVYNPFMIDVQQWGF TGNLQSNHDLYCQVHGN AH VASCDAIMTRCLAVHECFVKRVDW TIEYPIIGDELKINAACR KVQH MVVKAALLADKFPVLHDIGNPKAI KCV PQADVEWKFYDAQPCSDKAYKIEELFYSYATHSDKFTDGVCLFWNCNVDRYPANSIVCRFDTRVLS NLNLP GCDGGS LYVNKHAFHTPAFDKSAFVNLKQLPFFYSDSPCESHGKQVVSDIDYVPLKSATCITRC NLGGAVCRHHANEYRLYLDA YNMMISAGFSLWVYKQFDY NLWNTFTRLQ

The initial glycine residue in parentheses is a part of TEV recognition site and remains appended to nsp14 sequence in the purified nsp14. Table with nucleotide and amino acid sequence for SARS-CoV-2 nsp14 used in in this study.