

NMR Observation of Sulfhydryl Signals in SARS-CoV-2 Main Protease Aids Structural Studies

Angus J. Robertson,^[a, b] Jinfa Ying,^[a] and Ad Bax^{*[a]}

The 68-kDa homodimeric 3C-like protease of SARS-CoV-2, M^{pro} (3CL^{pro}/Nsp5), is a key antiviral drug target. NMR spectroscopy of this large system proved challenging and resonance assignments have remained incomplete. Here we present the near-complete (>97%) backbone assignments of a C145A variant of M^{pro} (M^{pro}_{C145A}) both with, and without, the N-terminal autocleavage substrate sequence, in its native homodimeric state. We also present SILLY (Selective Inversion of thiol and Ligand for NOESY), a simple yet effective pseudo-3D NMR experiment that utilizes NOEs to identify interactions between Cys-thiol or

aliphatic protons, and their spatially proximate backbone amides in a perdeuterated protein background. High protection against hydrogen exchange is observed for 10 of the 11 thiol groups in M^{pro}_{C145A}, even those that are partially accessible to solvent. A combination of SILLY methods and high-resolution triple-resonance NMR experiments reveals site-specific interactions between M^{pro}, its substrate peptides, and other ligands, which present opportunities for competitive binding studies in future drug design efforts.

Introduction

Thiol (-SH) groups of Cys residues in proteins can play important structural and functional roles but are often omitted from NMR studies because the proton lacks large J couplings to stable spin-1/2 isotopes that are required to record triple-resonance NMR spectra. We demonstrate that the uniquely upfield thiol chemical shifts, combined with considerable protection from solvent exchange, make these signals readily observable in a perdeuterated protein, where they provide valuable information for validating both assignments and structural models. The approach is illustrated for the main protease of the SARS-CoV-2 virus, termed M^{pro}, 3CL^{pro} or Nsp5, a chymotrypsin-like cysteine protease that plays an essential role in the viral replication process by excising 11 non-structural proteins (Nsp's), including itself, from the virus' two polyprotein chains. The H41-C145 catalytic dyad cleaves at the recognition sequence LQ↓S, and following the success of protease inhibitors to fight other viral diseases such as AIDS and hepatitis-C, the first FDA-approved specific M^{pro} inhibitor is also showing high clinical effectiveness.^[1] M^{pro} is only catalytically active as a

homodimer,^[2] with one solvent exposed active site (an oxyanion hole) per monomer. Inactivation of the monomeric enzyme is mediated by the conversion of a loop (S138-C145) to a short 3₁₀-helix, which collapses the oxyanion hole,^[3] a feature that was shown to be reversible on re-binding of substrate peptide with implications for M^{pro} liberation from the polypeptide chain.^[4] The recognition sequence of M^{pro} is nominally five residues, including four residues prior to the cleavage site (occupying subsites S4...S1) and the first residue past the cleavage site (P1' in subsite S1').^[5] The sequence of the SARS-CoV-2 variant of M^{pro} is 96% identical to that of SARS-CoV, with a high fitness cost to viral particles that develop M^{pro} inhibitor-resistance.^[6] The recent myriad of structure determinations by X-ray crystallography^[7] and corroboration by NMR^[8] indicate that none of the 12 conservative sequence differences significantly impact the backbone structure of M^{pro}.

In a search for leads in M^{pro}-targeted drug development, a recent NMR study of M^{pro} identified 38 hits.^[8a] Resonance assignment of large homodimeric proteins such as M^{pro} proved challenging and prior backbone amide assignments were limited to ~63%.^[8a] To prevent auto-proteolysis of the enzyme during prolonged exposure to the high concentrations (0.2–1.1 mM) and temperature (25–35 °C) needed for NMR measurements, we focus on two M^{pro} active site mutants, H41Q (M^{pro}_{H41Q}) and C145A (M^{pro}_{C145A}). Furthermore, NMR study of this 68-kDa homodimeric protein requires deuteration of the non-exchangeable hydrogens in combination with TROSY-based observation of the back-exchanged amide protons.^[9] Here, we take advantage of the 11 M^{pro}_{C145A} Cys sulfhydryl protons (¹H') that also exchange with solvent, as well as M^{pro}-substrate interactions, to validate and extend the M^{pro} assignments to essential completeness (>97%) and demonstrate that the interaction with its N-terminal autocleavage substrate extends well beyond the canonical five substrate-binding subsites S4–S1'.

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Results and Discussion

The M^{Pro} NMR assignment process proved considerably more challenging than expected (Figure 1; Supporting Information Experimental Section). Not only was a precarious partial-refolding step necessary to fully re-protonate the backbone amide groups after expression in D_2O (Figure S1, see Supporting Information text), but resonance broadening from conformational and solvent exchange (particularly for residues near the active site; Figure S2,S3), as well as conformational heterogeneity involving the presence of both *cis* and *trans* isomers of P184 (Figure S4) posed major challenges to software^[10] that usually makes the resonance assignment step straightforward. Nevertheless, nearly complete backbone assignments of M^{Pro}_{C145A} , M^{Pro}_{H41Q} and the complex with its N-terminal autocleavage substrate analogue, M^{Pro}_{C145A} :SAVLQSGFRK, were achieved (BMRB: 51455, 51456, Tables S1–S4, Supporting Information Figures S5–S7).

For dimethylsulfoxide (DMSO) and weakly binding ligands such as boceprevir^[11] (Figures S8 and S9),^[12] the site of interaction on the protein is commonly identified by observing chemical shift perturbations (CSPs) of protein resonances,^[13] which can be correlated with ligand observations.^[14] The absence of CSPs is commonly used to demonstrate the lack of interaction - as applies for ivermectin, which was proposed to act as a M^{Pro} inhibitor based on computational modeling^[15] (Figure S10). For ligands where the exchange rate between free and bound species is comparable to (or slower than) the difference in protein chemical shift between the apo and

ligated states, CSPs can be difficult to ascertain, particularly for larger proteins with crowded spectra. In this case, Nuclear Overhauser effects (NOEs) between ligand and protein can be used instead. For perdeuterated proteins, the sensitivity of this NOE-based approach can be strongly enhanced by using a band-selective pulse to simultaneously invert all upfield resonances (+3.5 to −0.7 ppm), representing most of the ligand's aliphatic hydrogens, and monitoring NOE transfer of their magnetization to protein backbone amide protons.^[16] Recovery of longitudinal 1H magnetization in large, perdeuterated proteins such as M^{Pro} is dominated by efficient 1H – 1H cross relaxation with hydroxyl and amide protons that rapidly exchange with the water signal, which is kept along the z-axis by the TROSY pulse scheme,^[9b] permitting short delays between scans (Table S6). The band-selective pulse also inverts protein sulfhydryl resonances (random coil chemical shift *ca* 1.9 ppm), prior to readout by the highly sensitive 2D 1H – ^{15}N -TROSY-HSQC pulse sequence, and therefore include NOE interactions to Cys- 1H resonances.^[9b] We refer to this method as “Selective Inversion of thiol and Ligand for NOESY” (SILLY), and the difference of spectra collected with- and without- the band-selective inversion pulse yields the final SILLY-TROSY spectrum. When applying the SILLY method to M^{Pro}_{C145A} in the presence of a 10-fold excess of the substrate analogue tetrapeptide N-acetyl-Val-DTyr-Leu-Gln (VylQ), the SILLY-TROSY spectrum shows many amides whose CSPs are challenging to resolve without NOE-editing (Figure 2). Because VylQ binds the protein with moderate affinity ($K_d \approx 80 \mu M$; Figure S11, Table S5) and the exchange between free and bound ligand is on the

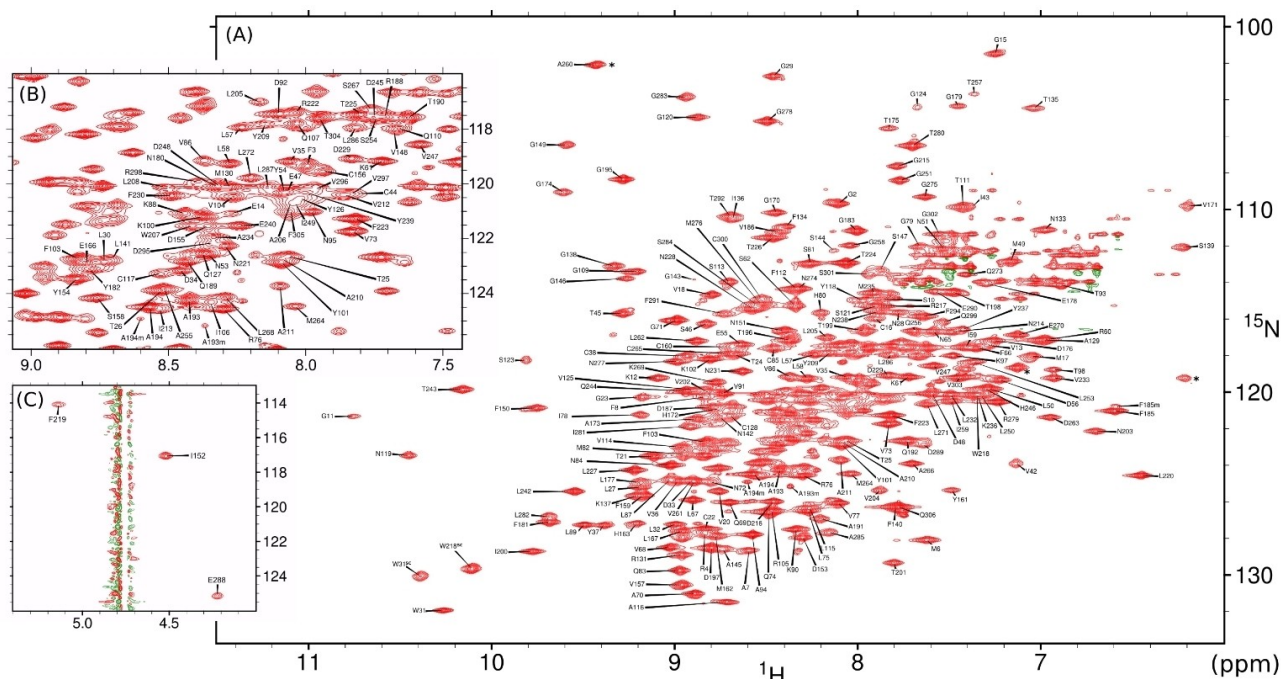


Figure 1. $^1H/^{15}N$ -TROSY-HSQC spectrum (900 MHz, 25 °C) of re-protonated M^{Pro}_{C145A} in 10 mM sodium phosphate pH 7.0, supplemented with 3% D_2O and 0.5 mM tris(2-carboxyethyl)phosphine hydrochloride (TCEP). (A) Full spectrum with aliased peaks marked with asterisks. (B) Expanded view of most crowded region of (A) with the contour threshold raised two-fold for clarity, and (C) three residues near the H_2O resonance, observed by omitting the typical water flip-back pulses from the $^1H/^{15}N$ -TROSY-HSQC pulse scheme.

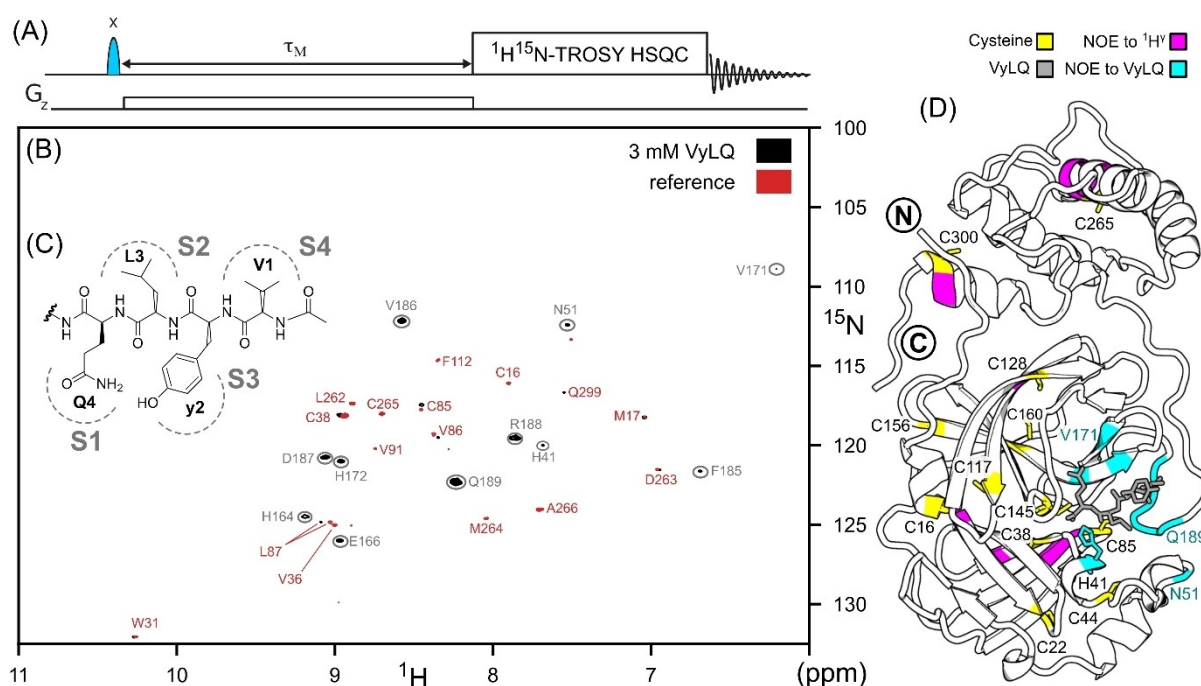


Figure 2. SILLY-TROSY spectra (800 MHz, 25 °C) used to investigate the binding of VyLQ to $\text{M}^{\text{pro}}_{\text{C145A}}$ in 10 mM NaPi pH 7, 0.5 mM TCEP, 3% D_2O . (A) Schematic view of the SILLY element preceding a standard TROSY-HSQC pulse scheme, with the band-selective I-BURP2 inversion pulse^[20] applied either on- (1.4 ppm) or off- (100 ppm) resonance in an interleaved manner, and a weak (0.35 G/cm) z-gradient applied during the NOE mixing period, τ_M . Care should be taken to eliminate water perturbation by the 'on-resonance' inversion pulse because even small (~2%) perturbations of the water signal will lead to bleed-through artifacts in the final spectrum and complicate analysis. The difference of spectra collected with the SILLY-element applied off/on resonance yields the final SILLY-TROSY spectrum. (B) Superimposed 800-MHz SILLY-TROSY spectra recorded for 0.3 mM (reprotonated) $^2\text{H}/^{15}\text{N}$ - $\text{M}^{\text{pro}}_{\text{C145A}}$ in either the absence (red) or presence (black) of 3 mM VyLQ, revealing M^{pro} backbone amides that have NOEs ($\tau_M = 0.5$ s) to $^1\text{H}^\gamma$ thiol (red labels) or aliphatic protons of the VyLQ peptide (grey circles). The long mixing time favors detection of NOEs to excess ligands in exchange between free and bound states. Shorter mixing times favor detection of NOEs to thiol protons (Figure S12). (C) VyLQ substrate analogue, with its corresponding subsite locations. (D) Residues with intra- (yellow; except C22 and C145) or inter- (magenta) residue NOEs to Cys $^1\text{H}^\gamma$ groups are indicated on the top ranked AF-M model of the $\text{M}^{\text{pro}}_{\text{C145A}}$:VyLQ complex (Figure S15), with both the H41/C145 catalytic dyad and the substrate peptide shown as sticks, and the N- and C-termini of M^{pro} indicated with encircled N and C characters, respectively. The tetrapeptide is presented as grey sticks, with intermolecular NOEs from VyLQ to $\text{M}^{\text{pro}}_{\text{C145A}}$ colored cyan on the AF-M model.

intermediate NMR time scale, an excess of ligand and a relatively long mixing time of 500 ms were used to maximize transfer of ligand magnetization to the protein, analogous to transferred NOE experiments.^[17] In contrast, magnetization from thiol protons is not replenished by exchange with free ligand and decays rapidly with increasing mixing times due to cross relaxation. NOEs from thiol to amide protons are therefore more intense at shorter mixing times such as 70 ms, where interactions are observed for the majority of Cys residues and their immediate neighbors in $\text{M}^{\text{pro}}_{\text{C145A}}$ (Figure S12). Such signals potentially provide valuable "anchors" during the early stage of sequential assignment and validation markers at a later stage, in particular for regions where sequential connectivities are sparse. Recording of the spectrum in the absence of ligand is essential to distinguish SILLY-TROSY resonances originating from thiol and unusually upfield shifted hydroxyl protons from ligand protons. In cases where perdeuteration is less complete than achieved in our work (ca 98%), weak signals resulting from NOEs to residual aliphatic sidechains may also appear in the SILLY-TROSY spectrum.

A more detailed view of the NOE interactions with the thiol signals can be obtained from three-dimensional (3D) ^{15}N -separated NOE spectra, recorded for an $\text{M}^{\text{pro}}_{\text{C145A}}$ sample that

contained the peptide N-acetyl-SAVLQSGFRK (SAVLQSGFRK). This auto-substrate peptide contains the M^{pro} N-terminal cleavage site required to excise itself from the SARS-CoV-2 precursor pp1a, where $\text{S6}_{\text{peptide}}$ corresponds to the N-terminal residue of mature M^{pro} . Affinity of M^{pro} for this peptide is high ($K_d \leq 4 \pm 2 \mu\text{M}$) and NOE interactions extend from the peptide's N-terminus to $\text{R9}_{\text{peptide}}$ (Figure S13) validating both a PDB X-ray structure,^[4] and AlphaFold-Multimer (AF-M)^[18] models of this complex (Figures S14, S15). Strips taken through such spectra at the $^{15}\text{N}/^1\text{H}$ frequencies of selected amide groups then reveal the thiol- $^1\text{H}^\gamma$ chemical shifts (Figure 3). Such resonances are valuable markers for validating resonance assignments, in particular for proteins such as M^{pro} for which good structural models are available.^[19] For example, $\text{C265-}^1\text{H}^\gamma$ shows NOEs to residues A260-K269, spanning nearly its entire α -helix (Figure 3A). Whereas most thiol NOEs are observed to amides proximate in sequence to each Cys, long-range NOEs are also observed. For example, NOEs from $\text{C300-}^1\text{H}^\gamma$ in the protein's C-terminal α -helix show proximity to N-terminal residues G2 and F3 that are essential to M^{pro} dimerization and catalytic activity (Figure 3B). Other examples include L87, which pairs with C38 in an antiparallel β -sheet and shows a strong NOE to the C38 thiol, and F112 which shows a strong NOE to the thiol of its β -sheet

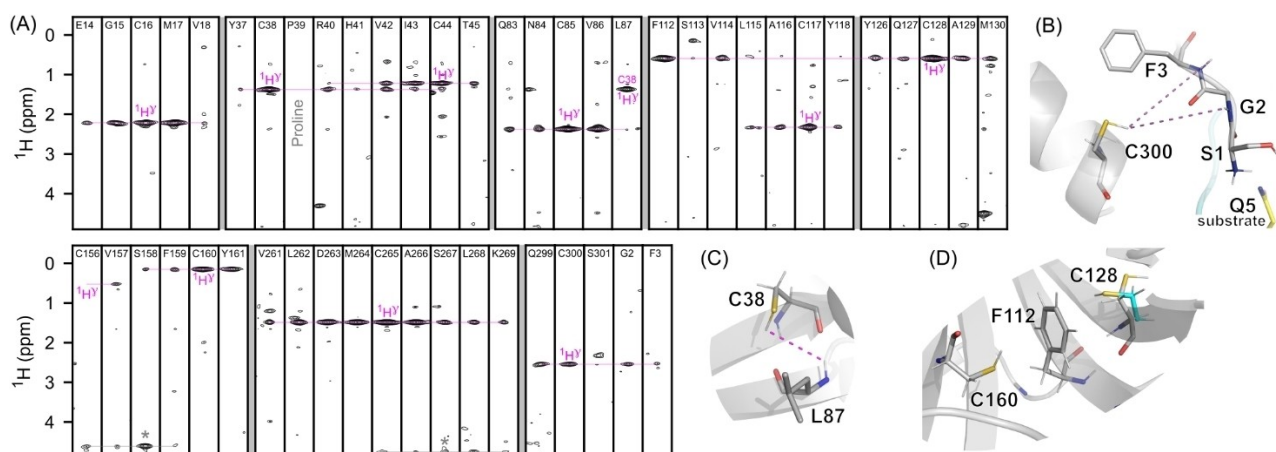


Figure 3. Observation of Cys- $^1\text{H}_\gamma$ in the $\text{M}^{\text{pro}}_{\text{C145A}}:\text{SAVLQSGFRK}$ complex. (A) Narrow strips taken from the 800-MHz ^{15}N -separated 3D NOESY-TROSY spectrum (800 MHz, 25°C), recorded with a 200 ms NOE mixing time, that show cross peaks from Cys- $^1\text{H}_\gamma$ protons (magenta annotations) to backbone $^1\text{H}^\text{N}$ atoms of residues indicated at the top of each strip. The full set of strips is included in Figure S17. (B,C,D) Examples of observed long range interactions between thiol protons and backbone amides or aromatic groups in the AF-M model of $\text{M}^{\text{pro}}:\text{SAVLQSGFRK}$, with long-range NOEs between thiol- $^1\text{H}_\gamma$ and amide- $^1\text{H}^\text{N}$ atoms shown as dashed magenta lines. In (B), substrate residue Q5 (yellow carbons) occupies the S1 subsite of the companion monomer, while in (D) the alternate conformer of C128 (cyan) taken from PDB 7MNG is required to reproduce the *ca* -1.3 ppm ring current shift, rather than the AF-M model (grey).

partner C128 (Figure 3C, 3D). Many more such interactions can be seen in the full set of strips (Figures S16, S17) and are invaluable to validate the backbone resonance assignments, typically based on triple-resonance 3D experiments such as HNCA (Figure S18).

The observation of sharp resonances for 10 of the 11 Cys sulfhydryl resonances in $\text{M}^{\text{pro}}_{\text{C145A}}$ is perhaps surprising considering that thiol protons exchange with solvent at rates that are catalyzed by both OH^- and phosphate and the intrinsic hydrogen exchange (HX) rate under the conditions of our sample is larger than 10^4 s^{-1} .^[21] Many of the thiols are buried in the protein interior and appear engaged in structure-stabilizing hydrogen bonds. With an S-H bond length of 1.34 \AA and an $\text{C}^\beta\text{-S}^\gamma\text{-H}^\gamma$ bond angle of $\sim 95^\circ\text{--}100^\circ$,^[22] thiols can act as donor for one H-bond and acceptor for two. The sulfhydryl can also engage in interactions between an aromatic π donor-orbital and the S-H σ^* acceptor-orbital,^[22] as appears to apply for C128- H^γ interacting with F112- C^ϵ (Figure 3D). Because the proton in such interactions is situated above the ring carbon, the ring current contribution to its chemical shift (*ca* -1.3 ppm) is smaller than if positioned directly above the center of the aromatic ring, and very sensitive to its precise location. For several of the other M^{pro} thiols, the pattern of thiol-stabilizing interactions is less clear cut. For example, with a *gauche*⁺ χ_1 rotamer, the C160 thiol appears to donate a third H-bond to G149-O, which already accepts two backbone H-bonds from the C160 and Y161 amide groups. However, a strong NOE interaction from C160- H^γ to Y161- H^N in the 70-ms mixing time SILLY-TROSY spectrum (Figure S12) is consistent with this geometry.

The observation of strong cross peaks to thiol resonances in the 3D NOESY spectrum (Figure 3A, S17), which employed a 200 ms NOE mixing time, indicates that solvent exchange of these labile protons cannot be much faster than 5 s^{-1} . The only thiol resonance missing from the 3D spectrum (Figure S17) is

that of partially solvent-exposed C22, whose intensity may also be adversely impacted by its *trans* χ_1 rotamer, resulting in a larger intrasidue $\text{H}^\text{N}\text{-H}^\gamma$ distance; rapid exchange with solvent of G23- H^N , which is close to C22- S^γ in the X-ray structure,^[19] prevents observation of this NOE. Other surface-exposed sulfhydryls include C85, C156, and C160 which show strong NOE interactions to one or more backbone amides, indicating that intramolecular H-bonds strongly protect these thiol protons from exchange with solvent.

The H^γ chemical shifts, seen in the 3D NOESY spectrum, allow multiple amides to be linked to individual thiol resonances, providing information not available from the SILLY-TROSY 2D spectrum. However, the sensitivity of the 2D spectrum is considerably higher, as reflected in a more than 20-fold shorter measurement time.

Conclusion

The importance of the local structure around Cys thiol groups, which due to their reactive chemistry can play important functional roles, is well recognized.^[23] The positions of hydrogen atoms are typically inaccessible in X-ray diffraction data and instead their presence is commonly modeled based on nearby H-bond acceptor atoms, which can often be ambiguous. The clear appearance of sulfhydryl signals in NOE spectra of perdeuterated proteins provides a straightforward avenue for studying these reactive groups in more detail, while simultaneously aiding protein resonance assignment. For M^{pro} , many of the thiols are close to aromatic moieties, leading to upfield ring current shifts (e.g. C128-F112; C160-F150; C265-F230) or, when located near the plane of such rings, to decreased chemical shielding (C85-F181; C300-F3), so that chemical shifts provide another sensitive tool for probing local structure, in addition to isotope effects.^[23] The nearly complete backbone assignment of

M^{pro} together with the introduction of two active site binding peptides with affinities of *ca* 4 and 80 μ M can facilitate competitive binding NMR assays in further drug development.

Experimental Section

The materials and methods used in our study can be found in the associated Supporting Information.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The backbone chemical shifts for M^{pro}_{C145A} (BMRB: 51455) and for M^{pro}_{C145A}:SAVLQSGFRK (BMRB: 51456) complexes have been deposited in the BioMagResBank (<http://www.bmrwisc.edu/>), and additional assignments are presented in the Supporting Information.

Keywords: AlphaFold-Multimers · ligand binding · NMR spectroscopy · M^{pro}/3CLpro · SARS-CoV-2

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