Molecular Modeling of the HR2 and Transmembrane Domains of the SARS-CoV-2 S Protein in the Prefusion State

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Abstract—SARS-CoV-2, the causative agent of COVID-19, remains the focus of research worldwide. SARS-CoV-2 entry into the cell starts with its S protein binding to the angiotensin-converting enzyme-2 (ACE2) expressed on the cell surface. The knowledge of the S protein’s spatial structure is indispensable for understanding the molecular principles of its work. The S protein structure has been almost fully described using experimental approaches with the only exception for the protein’s endodomain, the transmembrane domain, and the ectodomain parts adjacent to the latter. The paper reports molecular modelling of the S protein fragment corresponding to its coiled coil HR2 domain and fully palmitoylated transmembrane domain. Model stability in lipid bilayer was confirmed by all-atom and coarse-grained molecular dynamics simulations. It has been demonstrated that palmitoylation leads to a significant decrease in transmembrane domain mobility and local bilayer thickening, which may be relevant for protein trimerization.

Keywords: molecular modelling, SARS-CoV-2, S protein, lipid bilayer, palmitoylation, trimeric coiled coil

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We are currently experiencing the third wave of the COVID-19 pandemics caused by SARS CoV-2. Huge effort is being made worldwide towards the development of vaccines and drugs that would aid in overcoming the worldwide COVID-19 crisis. The virus surface S protein, which plays the key role in virus entry into the host cell by binding human angiotensin-converting enzyme 2 (ACE2), is the main target of most vaccines.

The S protein consists of the two subunits S1 and S2, which are produced by the cleavage at the Arg685-Ser686 site carried by the cellular furin protease. The S1 subunit contains the signal peptide, N-terminal domain, and receptor-binding domain that interact with the ACE2 receptor. The S2 subunit encompasses the fusion peptide, heptad repeats 1 and 2 (HR1 and HR2), transmembrane domain (TMD), and endodomain. The full-size membrane S protein, whose spatial structure remains unresolved so far, forms a homotrimeric complex and facilitates the fusion of the host cell and virus particle membranes, and consequently, virus entry into the cell [1].

A large amount of structural data is currently available for the S protein ectodomain, the primary immune system target, whereas we have much less information about its TMD and proximal regions localized close to the membrane [2]. It is assumed that the conformational mobility of the S protein is due to bending in a number of hinge regions, one of them being the region between HR2 and TMD [3]. Given that conformational mobility is crucial for the virus’s entry into the cell and possibly plays a role in the accessibility of the S protein surface epitopes to antibodies, constructing an accurate model of the S protein fragment, including TMD and neighboring domains, is critically important.

In the present work, we have built a structural model of the S protein fragment containing TMD and the adjacent HR2 domain located at the outer membrane surface and endodomain fragment at the inner surface, including the cysteine-rich domain (cys) domain. We used all-atom and coarse-grained molecular dynamics (MD) simulations in a complex membrane to optimize the obtained model and assess the effects of cysteine palmitoylation on S protein mobility in the membrane and its local lipid environment.

MATERIALS AND METHODS

All-atom model assembly and parametrization. The structural model of the SARS CoV-2 S protein fragment (amino acid residues 1157–1256 encompassing
the HR2 domain, TMD, and cyst domain) with the canonical sequence (Uniprot P0DT2C) was built using homology-based methods and two protein structures (PDB 2fxp [5] and 6e8w [6]) as templates with the aid of the Modeller software [4].

All-atom molecular dynamics protocol. The CHARMM-GUI web-server was used to preprocess the model before MD calculations [7] with palmitoylation of the cystein residues 1235, 1236, 1241, 1242, 1244, 1248, 1249, 1251, 1254, 1255 (Fig. 1a). Membrane lipid composition was as described previously [8]: cholesterol : palmitoyl-sphingomyelin (PSM) : palmitoyl-oleoyl-phosphatidylethanolamine (POPE) : dipalmitoyl-phosphatidylethanolamine (DPPE) : palmitoyl-oleoyl-phosphatidylcholine (POPC) : palmitoyl-oleoyl-phosphatidylcholine (DPPC) : dipalmitoyl-phosphatidylserine (POPS) : dipalmitoyl-phosphatidylcholine (DPPC) : dipalmitoyl-phosphatidylserine (DPPS) in a 15 : 10 : 9 : 6 : 3 : 3 : 2 : 2 ratio. The final ratio between lipid molecules after palmitoylation was 88 : 54 : 51 : 36 : 18 : 18 : 12 : 11, respectively. The total system size, including water molecules (TIP3P model) and Na + and Cl − ions (0.15 M), was 176 246 atoms. The cell size was 90 Å × 90 Å × 21 Å. System energy minimization and system equilibration was carried out according to the protocol recommended by the CHARMM-GUI web-server, which included the following steps: (1) system minimization using the steepest descend method, including 5000 steps, (2) six equilibration stages each 150–250 ps long with setting atoms velocities at the initial moment in accordance with the Maxwell distribution at 318 K and with gradual relaxation of the harmonic constraints on heavy atoms positions. MD was simulated using the Gromacs software [9] in the all-atom force field CHARMM36m [10] in the NPT ensemble at 318 K for 30 ns.

Coarse-grain model parametrization and coarse-grain molecular dynamics protocol. Initial models for coarse-grain MD calculations using the MARTINI 2.2 force field [11] were built using the martiniize.py script based on the obtained all-atom model. A script available at https://github.com/porekhov/palmitoylize was used for protein palmitoylation. Molecular topology for the palmitoylated cysteine residue was obtained from the work previously published by the Tieleman group [12]. Palmitoylated and nonpalmitoylated trimer models were embedded into the model bilayer with the same composition as that used for the all-atom calculation and solvated (standard MARTINI field water model and Na + and Cl − ions in the concentration of 0.15 M were used) using the insane.py scenario (script). MD protocol was chosen in accordance with the recommendations provided by the Marrink group [13]. The simulations were made in the NPT ensemble at 318 K. The integration interval was set as 20 fs. The trajectories 1 μs in length were obtained for both systems. Gromacs 2019.4 was used to carry out the simulations as it was in the previous case [9]. The g_lomepro software was used to calculate lipid bilayer thickness and lipid areas [14]. Python scripts implementing MDAnalysis libraries were used to calculate radial distribution functions.

RESULTS AND DISCUSSION

Model construction. The modelled S protein fragment contains several segments with differing degree of structural order (Fig. 1a). Modelling the HR2 domain is quite easy given the known structure of the same domain in the S protein of the closely related SARS-CoV coronavirus (100% identity for the residues 1157–1210) (PDB ID 2fxp, [5]). Monomer helices are oriented in such a way that the hydrophobic residues I1172, I1179, L1186, L1193, and L1200 are oriented towards the domain core.

At the same time, there are currently no high-identity templates available for the S protein TMD. For this reason, given the α-helical trimer structure and high number of aromatic residues present in the TMD N-terminus, we chose the human immunodeficiency virus (HIV) pg41 envelope protein TMD as a template (20% identity for the residues 1211–1240, with 33% identity for the residues 1211–1225 and 7% for the residues 1226–1240). Currently, a number of rather different structures have been obtained for this domain or one of its fragments (PDB ID 6e8w [6], 5yn [15], and 6b3u [16]) making it difficult to choose one of them as a template. Furthermore, the conformation of the residues 1205–1210 still remains unresolved since the corresponding residues in the template are disordered. α-helix may be one of the possible conformations. In this case, the HR2 domain and TMD would form a coiled coil [3]. However, such conformation appears to be sterically unfavorable because the numerous aromatic residues in the S protein are located at all sides of the α-helices hindering coiled-coil formation. As a result, we chose the 6e8w template, notwithstanding it having a structure unusual for a TMD [6] with bending α-helices and long 3–10-helices. However, it was possible with this template to locate the bulk side chains of the aromatic residues without overlapping and disturbing the helical symmetry of the coiled coil. In this case, Y1209, I1216, G1223, V1230, and M1237 residues are found at the coiled coil axis. The SARS-CoV-2 S protein TMD model obtained in this work (Fig. 1a; left) exhibited clear local similarity to the structure of the SARS-CoV S protein fragment (PDB ID 2run) obtained previously [17]. The deviation for the C atoms of the 11 central residues was not larger than 1 Å with most side chains having similar orientation.

Viral membrane protein palmitoylation has been studied for more than 40 years [18]. It is currently assumed that almost all cysteine residues in the C-terminal part of the S protein TMD and cyst-domain of coronaviruses, including SARS-CoV, are palmitoylated [19]. For this reason, the choice of the plexin
structure (PDB ID 5I5k) also enriched in cysteine residues linked by a high number of disulfide bonds as a template for modelling SARS-CoV-2 S protein cysteine domain structure made in Woo et al. (2020) [8] seems quite unreasonable. When building our model, we assumed all cysteine residues (ten in each subunit) to be palmitoylated. The final modelled structure is presented in Fig. 1a (left).

**All-atom molecular dynamics.** The CHARMM36m force field was chosen because of its being broadly used to study the dynamics of both proteins with low-ordered regions and protein–membrane systems [10]. Moreover, using this field allows one to more accurately reconstitute the conformation of experimentally determined structures, including short membrane proteins with non–ideal α-helical structure [20].

All-atom MD calculations showed that the structure of the studied protein undergoes some significant changes. It should be noted in this regard that, until the full structure of the protein is completely deciphered, the boundaries shown in Fig. 1a are, in a certain sense, tentative. For example, the TMHMM web service [21] predicts that the 1214–1234 residues (Fig. 1a) are located in the hydrophobic membrane layer. However, preliminary MD simulations showed that the location of bulk aromatic residues (W1214YWLGFL1220) in this membrane layer is sterically unfavorable, and, thus, this fragment “floats up” into the polar heads area of the outer lipid layer, simultaneously dragging some cysteine domain residues from the polar heads area of the inner layer into the hydrophobic membrane layer. At the same time, it is known...
that disordered secondary structure is untypical of the residues in the proteins’ transmembrane fragments, which suggests the helical structure for the residues downstream L1234 as well. The structures of the monomer (PDB ID 6b3u [16]) and trimers (PDB ID 5jyn [15] and 6e8w [6]) of the HIV envelope protein gp41 TMD with the helix length of more than 30 residues also speak in favor of the elongated helical TMD. Based on this comparison, the helical fragment in our model can extend up to the C1241 residue, with the total length of each transmembrane helix being 34 residues.

In the course of MD simulations, the N-tails generally retained their disordered structure (Fig. 1a; right) and demonstrated high mobility (Fig. 1b). However, partial ordering of the N-terminal residues with the formation of α-helices could be observed (Fig. 1c), which is consistent with the helical conformation of the residues at the C-termini of several S protein ectodomain structures (PDB ID 6x8r, etc.). On the contrary, the coiled coil state of the HR2 domain appeared to be very stable, with highest mobility having been observed at its boundaries. The hinge fragment totally changed its conformation, with relatively slight thermal fluctuations of the atoms. The transmembrane domain generally retained its helical structure, including the conformationally strained 3–10-helices observed in the template (PDB ID 6e8w). Large fluctuations are not typically characteristic of transmembrane helices, with the model proposed in this paper being no exception. Finally, despite palmitoylation and close interaction with the membrane, the cys-domain showed rather high mobility on average, which was largely due to the relatively mobile closing K1256 and F1256 residues. Here, it should also be noted that palmitoyl residues attached to the C1235 and C1236 residues exhibited different behavior compared to those attached to the other cysteine residues. In our model, the indicated cysteine residues belong to TMDs and are visibly embedded into the membrane. This results in that the acyl chains attached to them are able, in terms of geometry, to penetrate into the membranes outer lipid layer, while the acyl chains of the remaining palmitoyl residues are incapable of doing so and fluctuate exclusively within the inner lipid layer. In this connection, it is interesting to note that mass spectrometry has previously shown [22] that cysteine residues at the C-terminus of the influenza A virus hemagglutinin TMD undergo not palmitoylation as such but rather stearylation. Our observations suggest that the position of cysteine residues at the C-terminus of the S-protein TMD also allows their stearylation, while the addition of a longer aliphatic chain will result in local disordering of the membrane environment. Thus, the MD results show that, under natural conditions, stearylation is the most likely posttranslational modification of the cysteine residues in the S-protein TMD.
the conformational conversion of 3-10-helices into α-helices.

**Coarse-grain molecular dynamics.** To elucidate the effects of the cysteine residues' palmitoylation on the local TMD lipid environment, additional studies of palmitoylated and nonpalmitoylated proteins embedded in a membrane of similar composition were performed using a coarse-grained approximation that allowed increasing the simulation time to 1 μs.

The local membrane bilayer thickness was calculated for both systems (Fig. 2a). It can be seen that the presence of palmitoyl residues leads to a local thickening and distortion of the membrane around the protein (black area around the white area corresponding to the protein; Fig. 2a). The averaged local area per lipid with the embedded palmitoylated (left) and nonpalmitoylated (right) trimer is shown in Fig. 2b. The lipids around the protein are packed tighter than in the
remaining part of the membrane. In addition, we additionally calculated the radial distribution functions of the lipids present in the membrane relative to the TMD axis of mass in a membrane plane in two systems (Figs. 2e–2h). It can be seen from the obtained results that protein palmitoylation has hardly any effect on the membrane distribution of cholesterol (Fig. 2e) and some other lipids (DPSS, DPPC, DPFE, and PSM, plots not shown), while the radial distribution function values for POPE and POPS (Figs. 2f–2h) near the nonpalmitoylated protein are higher than those near the palmitoylated protein, i.e., the membrane in the TMD region becomes enriched in POPE and POPS. For POPC, the opposite effect is observed consisting in a significantly higher probability of detecting this lipid near a palmitoylated protein compared to a nonpalmitoylated one. Therefore, palmitoylation leads to a significant redistribution of lipid types in a realistic multicomponent bilayer model. Together with the observed membrane thickening and a decrease in the area per lipid near the palmitoylated protein, this may indicate the formation of a raft-like lipid microdomain in its vicinity. These observations are also in good agreement with the palmitoylation effects reported by Charollais and Van Der Goot (2009) [25]: palmitoyls can increase the effective hydrophobic length of the protein TMD by increasing the thickness of the adjacent lipid bilayer and facilitating association with ordered lipids.

In the course of MD simulation, the HR2 domain of the S-protein trimer tilts relative to the TMD (Figs. 2b–2d) in the same way as it does in the case of the all-atom simulation, which correlates well with the maximum angle distribution of approximately 20° and a significant portion of conformations showing the angle between the domains as low as 0° (i.e., a completely “straight” conformation). In addition, the presence of palmitoyl residues slightly reduces the amplitude of the S-protein bend. This may be partly due to the fact that a denser and thicker membrane layer is formed around the TMD protein decreasing its mobility.

The calculated lateral diffusion coefficients for palmitoylated and nonpalmitoylated protein $D_{\text{palmitoyl}} = (0.0205 \pm 0.0004) \times 10^{-5}$ and $D_{\text{nonpalmitoyl}} = (0.0025 \pm 0.0001) \times 10^{-5}$ cm$^2$/s indicate that palmitoylation decreases the protein mobility rate in the lateral membrane plane by an order of magnitude. It may be suggested that this decrease facilitates S-protein TMD trimer stabilization.

To summarize, the present study reports the first 3D model of the S-protein TMD and HR-2 domains that would consider cysteine palmitoylation. The protein model was embedded into the membrane of complex composition and optimized. Model stability was evaluated by all-atom and coarse-grain MD simulations. It was shown that the conformational mobility of the HR2-TMD fragment is achieved via bending in the hinge region located at the boundary between the two domains. We also showed that TMD palmitoylation leads to local changes in the bilayer (its thickening and compaction as well as lipid redistribution) similar to those taking place during raft-like microdomain formation. The model obtained in the present work may be used for interpreting structural data in further studies.

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**COMPLIANCE WITH ETHICAL STANDARDS**

Conflict of interests. The authors declare that they have no conflicts of interest.

Statement on the welfare of humans or animals. This article does not contain any studies involving humans or animals performed by any of the authors.

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