SARS-CoV-2 Spike Protein Cleavage and Fusion

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This sequence of events was described by Jason McLellan TWiV 714 [1].

- 1. You can get cleavage at S1/S2 site (see Figure 1) while spike is being produced in the infected cell since there is furin present. Here the spike trimer is in the prefusion confirmation.
- 2. In this case on the surface of an infectous viron the spike protein has already been cleaved at S1/S2.

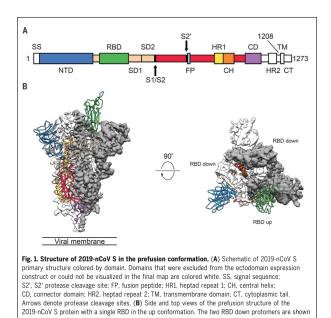


Figure 1: SARS-CoV-2 Spike Prefusion Confirmation [2]

as cryo-EM density in either white or gray and the RBD up protomer is shown in ribbons colored

corresponding to the schematic in (A).

- 3. Next S1 binds to ACE2 binding at the Receptor Binding Domains (RBDs), locking one, two or three of the RBDs in the "up" confirmation which is a thermodynamically unfavorable state.
- 4. The binding of S1 to ACE2 destabilizes the spike and causes S1 to shed and fall off S2.
- 5. S2 then undergoes a confirmational change and starts rearranging from its spring loaded state, extending towards the host cell membrane.
- 6. Cleavage at S2' (usually by TMPRSS2) liberates the fusion peptide from the new N-terminal domain of S2. S2 now is anchored in the host cell membrane and in the viral membrane.
- 7. S2 bends around and brings the host cell membrane into contact with the viral membrane. This is the post fusion state.

References

- [1] TWiV 714: The shape of spike with Jason McLellan. https://www.microbe.tv/twiv/twiv-714, 2021. [Online; accessed 01-Feburary-2021].
- [2] Daniel Wrapp et al. Cryo-EM structure of the 2019-nCoV spike in the prefusion conformation. https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7164637/pdf/367_1260.pdf, 2021. [Online; accessed 01-Feburary-2021].

¹The RBDs are locked in the "up" confirmation when bound by ACE2.