

Published in final edited form as:

Science. 2010 October 22; 330(6003): 505–508. doi:10.1126/science.1191714.

Mechanisms of Proton Conduction and Gating in Influenza M2 Proton Channels from Solid-State NMR

Fanghao Hu, Wenbin Luo, and Mei Hong*

Department of Chemistry, Iowa State University, Ames, IA 50011, USA

Abstract

The M2 protein of influenza viruses forms an acid-activated tetrameric proton channel. We used solid-state nuclear magnetic resonance spectroscopy to determine the structure and functional dynamics of the pH-sensing and proton-selective histidine-37 in M2 bound to a cholesterol-containing virus-envelope-mimetic membrane so as to better understand the proton conduction mechanism. In the high-pH closed state, the four histidines form an edge-face π -stacked structure, preventing the formation of a hydrogen-bonded water chain to conduct protons. In the low-pH conducting state, the imidazoliums hydrogen-bond extensively with water and undergo microsecond ring reorientations with an energy barrier greater than 59 kilojoules per mole. This barrier is consistent with the temperature dependence of proton conductivity, suggesting that histidine-37 dynamically shuttles protons into the virion. We propose a proton conduction mechanism in which ring-flip–assisted imidazole deprotonation is the rate-limiting step.

Proton transport in synthetic materials is mediated either solely by hydrogen-bonded (H-bonded) water, as in hydrated ionic polymers (1), or solely by titratable heterocycles, such as imidazoles tethered to the backbone of anhydrous polymers (2). In comparison, the conduction mechanism of biological proton channels in cell membranes is more complex because both water and titratable protein sidechains are usually present (3). The influenza M2 protein forms a tetrameric proton channel that is important for the virus life cycle (4). Activated below pH 6, the M2 channel conducts 10 to 10,000 protons per second (5, 6). The pH-sensing and proton-selective residue is a single histidine, His³⁷, in the transmembrane (TM) domain (7). ¹⁵N chemical shifts of His³⁷ in 1,2-dimyristoyl-*sn*-glycero-3-phosphocholine (DMPC)/dimyristoyl phosphatidylglycerol (DMPG) bilayers indicated that the four histidines titrate with pK_as of 8.2, 8.2, 6.3, and <5.0 (where K_a is the acid dissociation constant); thus, the third protonation event is responsible for channel activation (8). However, the precise role of His³⁷ in proton conduction is still debated. Two models have been proposed. In the “shutter” model, the pore at His³⁷ is enlarged through electrostatic repulsion among the imidazoliums, permitting a continuous H-bonded water

*To whom correspondence should be addressed. mhong@iastate.edu.

Supporting Online Material

www.sciencemag.org/cgi/content/full/330/6003/505/DC1

Materials and Methods

Figs. S1 to S9

Tables S1 and S2

References

chain over which protons hop by means of the Grotthuss mechanism (9, 10). The rate-limiting step is proton-hopping across three or four charged imidazoliums, with a calculated energy barrier of 29 to 42 kJ/mol (9). In the “shuttle” model, His³⁷ actively participates in proton relay through protonation and deprotonation. Tautomerization or ring flips reestablish the original conformation required for the next proton relay (11). The rate-limiting step in this model is the His³⁷ conformational change.

Although high-resolution structures of the M2 TM domain (M2TM) in detergents at high and low pH have been reported (12, 13), the His³⁷ sidechain conformations differed in these structures, and sidechain dynamics and water interactions were not probed. Further, detergent molecules can perturb the packing of weakly bound membrane protein complexes; thus, the structures may not accurately reflect the chemistry of the imidazoles in the lipid membrane.

To elucidate the proton conduction mechanism of M2, we used solid-state nuclear magnetic resonance (NMR) to determine the structure and dynamics of His³⁷ in M2TM reconstituted into a cholesterol-rich virus-envelope-mimetic lipid membrane (14, 15). Extensive data yielded the His³⁷ protonation state, tautomerization, rotameric conformation, sidechain dynamics, and hydrogen bonding from pH 8.5 to 4.5. Here, we focus on pH 8.5 for the closed channel and pH 4.5 for the conducting channel. M2TM exhibits acid-activated and amantadine-sensitive proton currents similar to the intact protein (16) and fully assembles into four-helix bundles (17) in the viral membrane with immobilized backbones (14), which allowed His³⁷ sidechain motion to be elucidated.

Histidine ¹⁵N and ¹³C chemical shifts are exquisitely sensitive to the protonation state and tautomeric structure of imidazoles. Deprotonation increases the ¹⁵N chemical shift by ~80 parts per million (ppm) (18), and C γ /C δ 2 chemical shifts also systematically depend on the imidazole structure (19). Two-dimensional (2D) ¹³C-¹³C and ¹⁵N-¹³C correlation spectra of His³⁷-labeled M2TM revealed only neutral imidazoles at pH 8.5. The N ϵ 2-protonated τ -tautomer and N δ 1-protonated π -tautomer exist at a ~3:1 ratio (Fig. 1), with slow or no exchange at ambient temperature (fig. S1). Inter-tautomer C δ 2(τ)-C γ (π) and C δ 2(τ)-C δ 2(π) cross peaks (Fig. 1A) indicate that both tautomers exist in each channel. The ~25% fraction of the π -tautomer is much higher than in small imidazole-containing compounds (18), suggesting stabilization of the protonated N δ 1(π) through hydrogen bonding (20).

At pH 4.5, both N δ 1 and N ϵ 2 exhibit protonated chemical shifts (170 to 180 ppm); no unprotonated signal was observed at 250 ppm (fig. S1), indicating that the neutral species is below the detection limit (< 5%). The charged imidazoliums showed much larger linewidths than the neutral species (table S1), indicating broader conformational distribution of the protein at low pH.

We probed interhelical packing of the His³⁷ tetrad through χ ₁- and χ ₂-dependent backbone-sidechain distances. The C α -N δ 1 distance constrains the χ ₂ torsion angle, whereas the C δ 2-N α distance constrains both χ ₁ and χ ₂ angles. At both pH, the C α -N δ 1 distance was 3.9 Å (Fig. 2 and fig. S2), which ruled out the +60° and -60° χ ₂ rotamers and was consistent only with the 180° rotamer. Similar experiments yielded a C δ 2-N α distance of 4.4 to 4.9 Å (fig.

S3), which ruled out the $\chi_1 = -60^\circ$ rotamer. Thus, the high-pH τ -tautomer adopts the *tt* rotamer, which is consistent with interhelical His³⁷-Trp⁴¹ (21) and Trp⁴¹-Trp⁴¹ distances (17). At pH 4.5, the C α -N δ 1 (3.9 Å), C α -N ϵ 2 (4.4 Å), and C δ 2-N α (>4.5 Å) distances similarly indicated the *tt* conformation (Fig. 2B and fig. S3).

Placing the His³⁷ rotamer into the different backbone structures at low pH and high pH revealed substantially different packing of the imidazole tetrad (Fig. 2, C and D). In the closed channel (22), the major τ -tautomers pack in an edge-face fashion in which each C ϵ 1-H ϵ 1 bond points to the center of the neighboring ring. The packing is tight, with a nearest-neighbor C ϵ 1-N ϵ 2 distance of ~4.9 Å, which is consistent with inter-tautomer cross peaks and suggests aromatic CH- π interaction (23). The high density of π -electrons should repel water oxygens and orient them in opposite directions across the tetrad, thus disabling proton conduction. The tetrad dimension is still possible for metal-ion coordination (24), which may explain Cu²⁺ inhibition of M2 (25). The minor π -tautomer can readily maintain the fourfold symmetry by adopting *tt* or *t0* rotamer. In comparison, in the low-pH structure (13) the imidazoliums show no edge-face stacking and leave a much wider pore.

The considerable packing difference suggests that His³⁷ sidechains may be immobilized at high pH but dynamic at low pH. To test this hypothesis, we measured one-bond C γ -N δ 1 and C δ 2-H δ 2 dipolar couplings at physiological temperature. The viral membrane immobilized the protein backbone, giving N α -H and C α -H α order parameters of 1.0 (figs. S4 and S5) (14), thus isolating potential sidechain motion. Fast motions scale the couplings by an order parameter (*S*) that reflects the motional amplitude. At pH 8.5, we obtained rigid-limit C γ -N δ 1 and C ϵ 1-N δ 1 couplings (1.15 kHz and 1.39 Å) and a rigid-limit C δ 2-H δ 2 coupling (23.9 kHz and 1.08 Å) (Fig. 3, A and B), confirming immobilization of the neutral imidazoles by edge-face stacking. However, at pH 4.5 the C γ -N δ 1 and C δ 2-H δ 2 couplings are scaled by a factor of 0.85 and 0.80, respectively (Fig. 3C), indicating sidechain motion. The availability of two-order parameters constrained the geometry of the imidazolium motion. The most likely motional axis is the C β -C γ bond (26). Uniaxial rotation is ruled out because it predicts a very small $S_{C\gamma-N\delta 1}$ of 0.06 because of the 57° angle of the C γ -N δ 1 bond to the C β -C γ axis. The well-known 180° ring flip motion is also ruled out because it has little effect on the C δ 2-H δ 2 coupling ($S_{C\delta 2-H\delta 2} = 0.94$) (table S2). The near-invariance of the C δ 2-H δ 2 coupling to 180° ring flips also rules out a scenario in which some imidazoliums undergo ring flips whereas others remain static (fig. S6). Instead, analysis of the *S* dependence on χ_2 angles shows that only two-site jumps with a χ_2 change of 45° satisfies both the C γ -N δ 1 and C δ 2-H order parameters (fig. S7). Given the average χ_2 of 180° at low temperature, the most likely instantaneous χ_2 angles are about 160° and -155° (Fig. 3D).

The restricted nature of the imidazolium ring reorientation may result from the symmetric low pH across the bilayer in the NMR samples because the imidazoles may not need to substantially reorient to be deprotonated and reprotonated. When a proton concentration gradient exists, such as in the virus membrane, full ring flips may occur. The motion must be much faster than 10⁴ s⁻¹ to average the C δ 2-H δ 2 coupling. Temperature-dependent C δ 2-H δ 2 couplings from 308 to 243 K indicated that the imidazolium was frozen by 263 K but fully mobile at 293 K (fig. S8). Using a lower-limit of 50 kHz for the 293 K rate and an

upper-limit of 3 kHz for the 263 K rate, we obtained an energy barrier of >59 kJ/mol, which is consistent with the 50 to 120 kJ/mol reported for imidazole motions in synthetic proton conductors (27). M2 proton conductivities differ by 14-fold between 18° and 37°C at pH 5.7 (5), indicating an energy barrier of 104 kJ/mol. Thus, the barrier of imidazolium motion is consistent with the functional data, whereas the barrier for water-mediated proton hopping (29 to 42 kJ/mol) is not (9), suggesting that His³⁷ ring reorientation is directly involved in proton transport, as in the “shuttle” model. Imidazole motion was also observed at the physiological pH of 6.0 and 5.2, at which the channel first opened and both charged and neutral histidines were present (fig. S8) (7, 8). Thus, His³⁷ motion appears to be an intrinsic property of the spacious conducting channel, although its precise amplitudes and rates may vary with pH.

Water is still necessary for delivering protons to the imidazoles before they can be relayed to the virus interior. Thus, water-His³⁷ hydrogen bonding is implied in the shuttle model. We probed H-bond formation by measuring imidazole N-H and C-H dipolar couplings at 243 K, at which the sidechain was frozen. H-bond formation stretches the N-H and C-H bonds from their covalent lengths (1.03 Å and 1.10 Å), thus weakening dipolar couplings (28). At pH 8.5, the protonated Nδ1(π) showed a significantly stretched N-H bond of 1.08 Å, indicating hydrogen bonding and explaining the π-tautomer stabilization. Even the unprotonated Nδ1(τ) showed a sizeable coupling of 2.1 kHz, suggesting a nearest-proton distance of 1.8 Å and a weak Nδ1...H-O H-bond. In contrast, the protonated Nε2(τ) exhibited an unstretched bond length of 1.03 Å (11.1 kHz) (Fig. 4A), despite the presence of a small amount of water in the H37-W41 region on the basis of Nε2(τ)-water cross peaks in 2D ¹⁵N-¹H correlation spectra (fig. S9). At pH 4.5, the combined Nδ1/Nε2 peak showed a reduced N-H coupling of 8.8 kHz, indicating a stretched bond of 1.11 Å. Given the spaciousness of the low-pH pore, the H-bond acceptors cannot be another imidazolium. Trp⁴¹-His³⁷ aromatic interaction may partly contribute to Nε2-H bond stretching (10, 29), but we propose the most likely cause for N-H bond elongation at low pH is H-bond with frozen water, which is more abundant in the low-pH channel than the high-pH channel, as shown with spin diffusion NMR (30) and molecular dynamics simulations (9, 31).

Similar C-H coupling measurements revealed that the Cδ2-Hδ2 bond was unstretched (1.08 Å) at either pH, whereas the Cε1-Hε1 bond was stretched to 1.20 Å at pH 8.5 and 1.19 Å at pH 4.5 (Fig. 4B). The latter may be attributed to CH-π interactions at high pH and H-bond with water at low pH. The imidazole Cε1-Hε1 bond is known to be prone to elongation because of its acidic nature (32), although the large magnitude of stretching observed here is surprising and requires further investigation.

These bond lengths reveal an extensive H-bond network that covers three sides of the imidazolium at low pH (Fig. 4C), creating a continuous H-bonded chain. Similar to the histidine in the catalytic triad of serine proteases, Cε1 hydrogen bonding may facilitate Nε2 deprotonation by evenly distributing the positive charge and increasing Nε2 electronegativity (26, 33). At high pH, the H-bond network is incomplete, excluding Nε2, which we attribute to the opposing water orientation and possible His³⁷-Trp⁴¹ interactions (Fig. 4D).

Taken together, these data suggest the following mechanism for proton gating and conduction by M2 (Fig. 4, D and E). At high pH_{out} , the neutral imidazoles form tightly packed electron-rich CH- π stacks, preventing the formation of a H-bonded water chain. The outward-facing N δ 1(π) H-bonds with water, whereas the inward-facing N ϵ 2 does not. Lowering pH_{out} protonates N δ 1, resulting in several imidazoliums per channel, which repel each other and cause backbone conformational changes that widen the pore (34, 35). More water permeates this region (30), establishing a H-bonded chain that includes His³⁷. The larger pore frees the imidazoliums to undergo microsecond ring reorientations. We propose a proton conduction mechanism in which imidazolium deprotonation is facilitated by C ϵ 1-H ϵ 1 hydrogen bonding and continuous ring flips achieve the dual purpose of properly aligning the charged imidazolium with the C-terminal water molecules so as to cause proton transfer and then pointing the unprotonated nitrogen to the low-pH extracellular side to be reprotonated. Our data indicate that the highest energy barrier of this process is the imidazolium motion, which may account for the temperature dependence of M2 proton conductance, possibly in combination with an additional small barrier for proton transfer (5). This dynamically assisted proton transfer model is consistent with the observed deuterium isotope effect, whose magnitude also suggested a mixed H-bonded chain with dissimilar elements (6). Thus, the present data strongly suggest that His³⁷ is actively involved in proton conduction by M2. The structural information obtained here is largely invisible to conventional high-resolution techniques and demonstrates the ability of solid-state NMR to elucidate functionally important membrane protein dynamics and chemistry.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

References and Notes

1. Mauritz KA, Moore RB. *Chem Rev.* 2004; 104:4535. [PubMed: 15669162]
2. Bureekaew S, et al. *Nat Mater.* 2009; 8:831. [PubMed: 19734885]
3. Nagle JF, Morowitz HJ. *Proc Natl Acad Sci USA.* 1978; 75:298. [PubMed: 272644]
4. Cady SD, Luo WB, Hu F, Hong M. *Biochemistry.* 2009; 48:7356. [PubMed: 19601584]
5. Lin TI, Schroeder C. *J Virol.* 2001; 75:3647. [PubMed: 11264354]
6. Mould JA, et al. *J Biol Chem.* 2000; 275:8592. [PubMed: 10722698]
7. Wang C, Lamb RA, Pinto LH. *Biophys J.* 1995; 69:1363. [PubMed: 8534806]
8. Hu J, et al. *Proc Natl Acad Sci USA.* 2006; 103:6865. [PubMed: 16632600]
9. Chen H, Wu Y, Voth GA. *Biophys J.* 2007; 93:3470. [PubMed: 17693473]
10. Okada A, Miura T, Takeuchi H. *Biochemistry.* 2001; 40:6053. [PubMed: 11352741]
11. Pinto LH, et al. *Proc Natl Acad Sci USA.* 1997; 94:11301. [PubMed: 9326604]
12. Schnell JR, Chou JJ. *Nature.* 2008; 451:591. [PubMed: 18235503]
13. Stouffer AL, et al. *Nature.* 2008; 451:596. [PubMed: 18235504]
14. Luo W, Cady SD, Hong M. *Biochemistry.* 2009; 48:6361. [PubMed: 19489611]
15. Materials and methods are available as supporting material on *Science* Online.
16. Ma C, et al. *Proc Natl Acad Sci USA.* 2009; 106:12283. [PubMed: 19590009]
17. Luo W, Mani R, Hong M. *J Phys Chem.* 2007; 111:10825.
18. Munowitz M, Bachovchin WW, Herzfeld J, Dobson CM, Griffin RG. *J Am Chem Soc.* 1982; 104:1192.

19. Henry B, Tekely P, Delpuech JJ. *J Am Chem Soc.* 2002; 124:2025. [PubMed: 11866617]
20. Bachovchin WW, Roberts JD. *J Am Chem Soc.* 1978; 100:8041.
21. Nishimura K, Kim S, Zhang L, Cross TA. *Biochemistry.* 2002; 41:13170. [PubMed: 12403618]
22. Cady SD, et al. *Nature.* 2010; 463:689. [PubMed: 20130653]
23. Waters ML. *Curr Opin Chem Biol.* 2002; 6:736. [PubMed: 12470725]
24. Brinen LS, Willett WS, Craik CS, Fletterick RJ. *Biochemistry.* 1996; 35:5999. [PubMed: 8634241]
25. Gandhi CS, et al. *J Biol Chem.* 1999; 274:5474. [PubMed: 10026160]
26. Ash EL, et al. *Proc Natl Acad Sci USA.* 2000; 97:10371. [PubMed: 10984533]
27. Fischbach I, Spiess HW, Saalwachter K, Goward GR. *J Phys Chem.* 2004; 108:18500.
28. Song XJ, Rienstra CM, McDermott AE. *Magn Reson Chem.* 2001; 39(S1):S30.
29. Tang Y, Zaitseva F, Lamb RA, Pinto LH. *J Biol Chem.* 2002; 277:39880. [PubMed: 12183461]
30. Luo W, Hong M. *J Am Chem Soc.* 2010; 132:2378. [PubMed: 20112896]
31. Yi M, Cross TA, Zhou HX. *J Phys Chem B.* 2008; 112:7977. [PubMed: 18476738]
32. Scheiner S, Kar T, Pattanayak J. *J Am Chem Soc.* 2002; 124:13257. [PubMed: 12405854]
33. Derewenda ZS, Derewenda U, Kobos PM. *J Mol Biol.* 1994; 241:83. [PubMed: 8051710]
34. Khurana E, et al. *Proc Natl Acad Sci USA.* 2009; 106:1069. [PubMed: 19144924]
35. Yi M, Cross TA, Zhou HX. *Proc Natl Acad Sci USA.* 2009; 106:13311. [PubMed: 19633188]
36. We thank S Li for discussions This work was supported by NSF grant MCB-543473 and NIH grant GM088204.

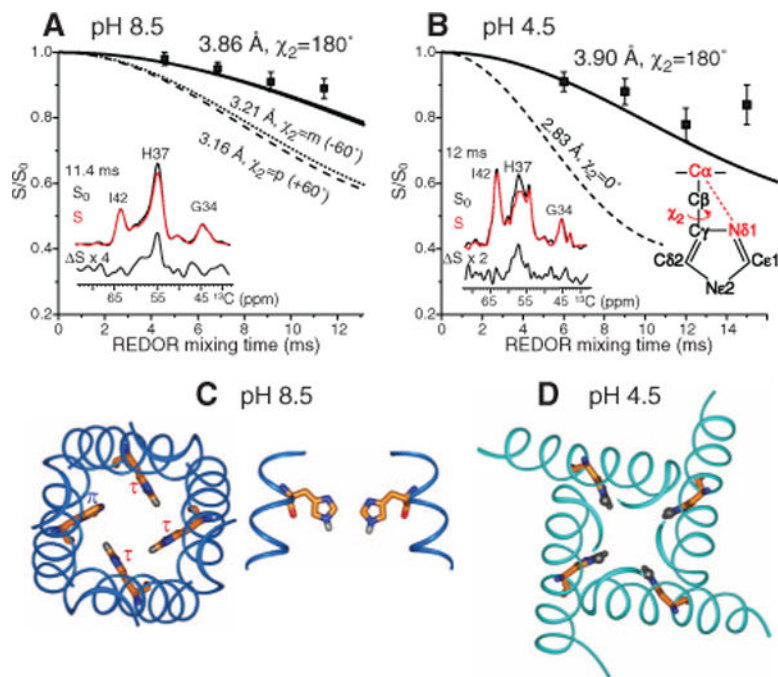


Fig. 2. His³⁷ rotameric conformation from C α -N $\delta 1$ distances. **(A)** pH 8.5 data, with representative rotational-echo double-resonance control (S_0), dephased (S), and difference (ΔS) spectra. The 3.9 Å distance indicates $\chi_2 = 180^\circ$. **(B)** pH 4.5 data, showing a similar distance and χ_2 angle. **(C)** Top and side views of the His³⁷ tetrad in the *tt* rotamer in the high-pH structure [Protein Data Bank (PDB) number 2KQT] (22). **(D)** Top view of the His³⁷ tetrad in the *tt* rotamer in the low-pH structure (PDB number 3C9J) (13).

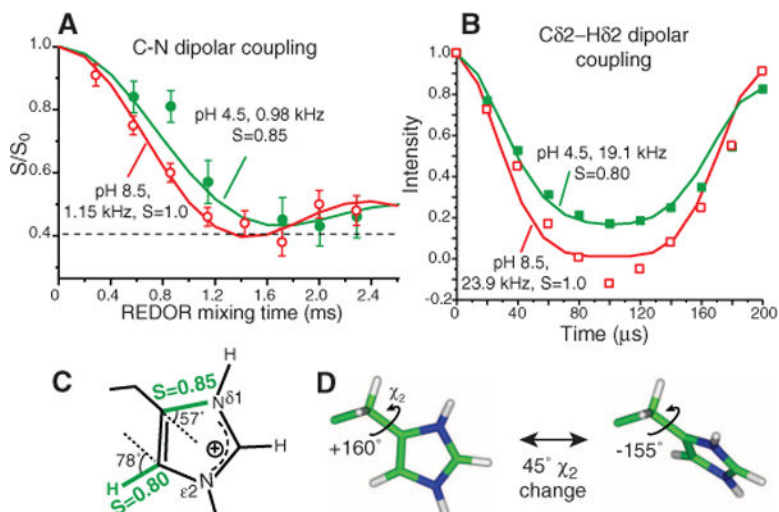
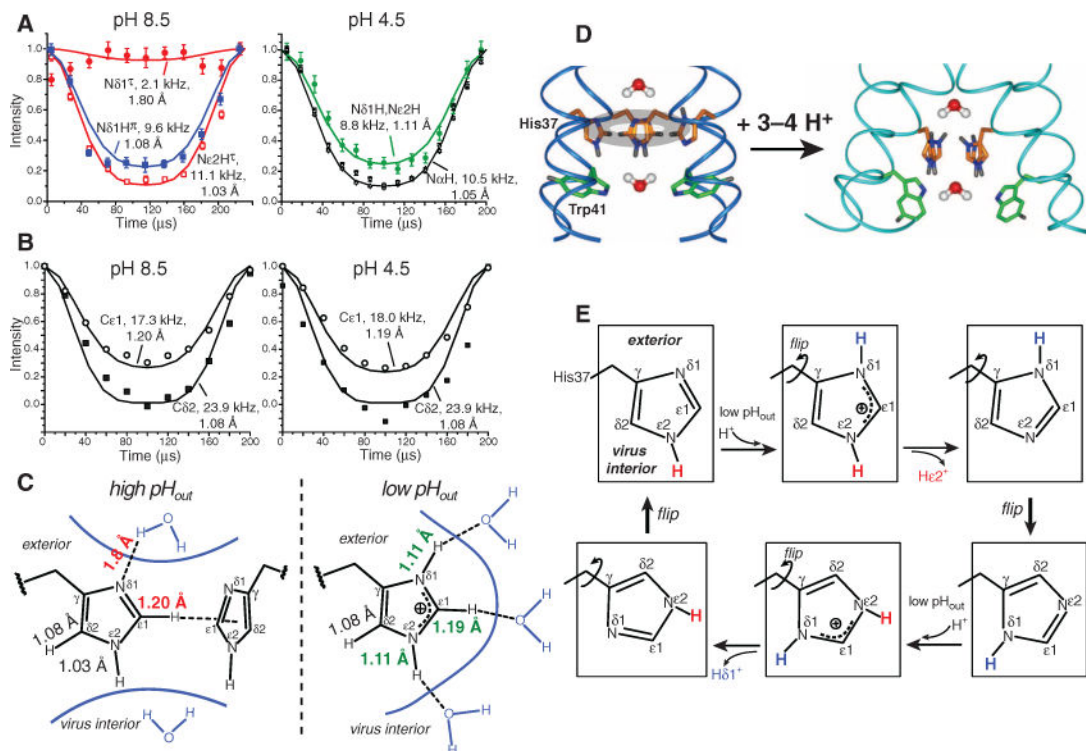


Fig. 3. His³⁷ sidechains reorient at low pH but remain static at high pH at physiological temperature. **(A)** 303 K ¹³C-¹⁵N dipolar couplings. At pH 8.5, a 1:1 combination of C γ -N δ 1 and C ϵ 1-N δ 1 couplings reaches the rigid limit. At pH 4.5, the dominant C γ -N δ 1 coupling is motionally averaged. **(B)** C δ 2-H δ 2 coupling at 308 K is motionally averaged at pH 4.5 but in the rigid limit at pH 8.5. **(C)** Measured order parameters at pH 4.5. **(D)** Two-site jump of imidazolium at low pH. A 45° reorientation around the C β -C γ bond fits the observed order parameters.

**Fig. 4.**

Charged His³⁷ H-bonds with water and undergoes ring flips to relay protons. **(A)** N-H dipolar couplings at 243 K. At pH 8.5, N ϵ 2(τ) is not H-bonded, whereas N δ 1(π) is. The unprotonated N δ 1(τ) shows a weak H-bond. At pH 4.5, both nitrogens show weak couplings and bond stretching. **(B)** C-H dipolar couplings of C δ 2 and C ϵ 1 at 243 K. The C ϵ 1-H ϵ 1 bond is stretched, whereas C δ 2-H δ 2 is not. **(C)** Imidazole bond lengths and H-bond networks at high and low pH. **(D)** His³⁷ structure and dynamics and proposed water orientations across the tetrad. Trp41 may interact with His³⁷ at high pH_{out}. **(E)** Proposed imidazole structural changes in a cycle in which multiple ring reorientations mediate the transfer of two protons.