

Figure S1. Choice of cutoffs for disconnected sub-graphs. Percentage of the total number of nodes in the graph (y-axis) as a function of correlation coefficient (x-axis) for the Δ F508 (top) and Δ I507 (bottom) networks. The correlation coefficient at which at least half the total number of nodes in the graph (half of the protein residues) reside in the largest connected component is chosen as a cutoff in generating the disconnected sub-graph. The dashed line represents half of the maximum number of nodes in the graph.

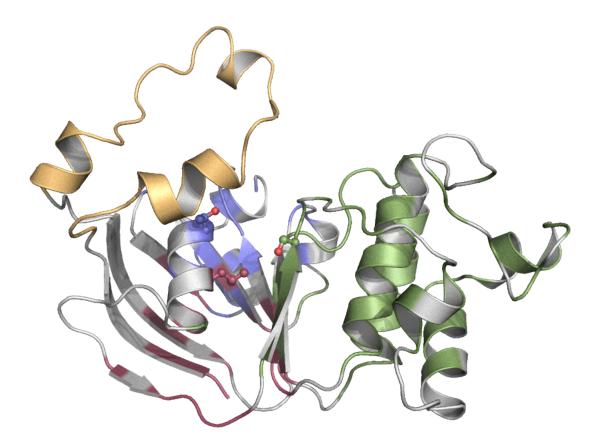


Figure S2. Regions in NBD1 affected by critical nodes. Three critical nodes – S492, T465, and L468 – are shown in ball-and-stick representation. S492 influences residues shown in green, T465 influences those shown in blue and red, and L468 influences those in red. Other critical nodes affect subsets of the residues colored red, green, and blue.

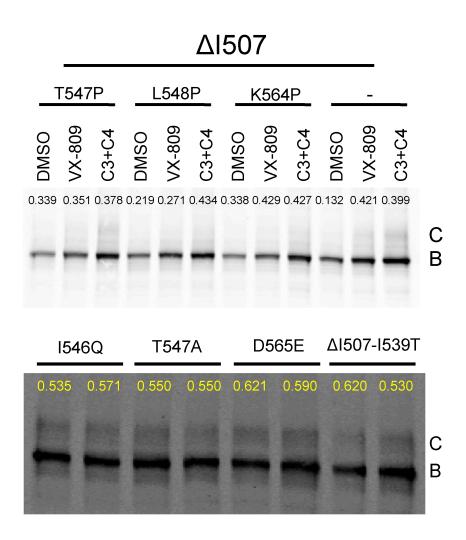


Figure S3. Mutations to critical nodes affect critical maturation pathways in Δ I507-CFTR. HEK cells transiently transfected with Δ I507-CFTR with or without additional mutations. We treat cells with indicated correctors 48 hours post-transfection, and grow at 27°C for an additional 24 hours before harvesting for western blot analysis (20 µg proteins/lane). Ratio of C-band to B-band is displayed below each lane.

Q07DV2 Aotus nancymaae 487-RISFCSQFSWI-497 Q00PJ2 Atelerix albiventris 487-RISFCSQFSWI-497 Q09YK5 Ateles geoffroyi 487-RISFCSOFSWI-497 P35071 Bos taurus 486-RISFCSQYSWI-496 Q2QLB4 Callicebus moloch 487-RISFCSQFSWI-497 Q2QLF9 Callithrix jacchus 487-RISFCSQFSWI-497 Q5U820 Canis fimiliaris 487-RISFCSQFSWI-497 Q2QLC5 Carollia perspicillata 487-RISFCSQFSWI-497 Q07E05 Cavia porcellus 487-RVSFCSQFSWI-497 Q2IBA1 Cercopithecus aethiops 487-RISFCSOFSWI-497 Q07DY5 Colobus quereza 487-RISFC**S**QFSWI-497 A0M8W1 Danio rerio 486-RISYSSOTAWI-496 Q07E42 Dasypus novemcinctus 487-RISFCSQFSWI-497 A4ZE37 Dicentrarchus labrax 486-RISFSPQTSWI-496 Q2QL74 Didelphis marsupialis 487-RISFCSQFSWI-497 Q2QLA3 Equus caballus 486-RISFCSQFSWI-496 A0M8T4 Felis catus 487-RISFCSQFSWI-497 A0M8U4 Gallus gallus 488-RISFSPQVSWI-498 Q2IBF6 Gorilla gorilla 487-RISFCSQFSWI-497 P13569 Homo sapiens 487-RISFC<mark>S</mark>QFSWI-497 Q07DX5 Hylobates leucogenys 487-RISFCSOFSWI-497 Q108U0 Loxodonta africana 487-RISFCSQFSWI-497 07JII8 Macaca fascicularis 487-RISFCSQFSWI-497 Q7JII7 Macaca fuscata 487-RISFCSQFSWI-497 09TUO2 Macaca memestrina 487-RISFCSOFSWI-497 000553 Macaca mulatta 487-RISFCSOFSWI-497 A4D7T4 Macropus eugenii 487-RISFCSQFSWI-497 Q2QL83 Microcebus murinus 487-RISFCSQFSWI-497 Q09YJ4 Muntiacus muntjak 486-RISFCSQYSWI-496 Q07DW5 Muntiacus reevesi 486-RISFCSOYSWI-496 P26361 Mus musculus 487-RVSFC**S**QFSWI-497 O07E16 Mustela putorius 487-RISFCSOFSWI-497 Q07DZ6 Ornithorhynchus anatinus 487-RISFSPQFSWI-497 Q00554 Oryctolagus cuniculus 487-RISFCSQFSWI-497 Q2QLH0 Otolemus garnettii 487-RISFCSQFSWI-497 000555 Ovis aries 486-RISFCSOYSWI-496 Q2QLE5 Pan troglodytes 487-RISFCSOFSWI-497 Q9TSP5 Papio anubis 487-RISFCSQFSWI-497 Q2IBE4 Pongo abelii 487-RISFCSQFSWI-497 P34158 Rattus norvegicus 487-RVSFS**S**QISWI-497 Q2IBB3 Rhinolophus ferrumequinum 487-RISFCSQFSWI-497 Q09YH0 Saimiri boliviensis 487-RISFCSQFSWI-497 098TY5 Salmo salar 486-RISFSPOTSWI-496 P26362 Squalus acanthias 488-RISYS**P**QVPWI-498 Q6PQZ2 Sus scrofa 487-RISFCSQFSWI-497 Q5D1Z7 Trichosurus culpecula 487-RISFCSQFSWI-497 P70034 Xenopus laevis 488-RISFSPOVSWI-498

Figure S4. Sequence conservation in CFTR from different species. Amino acid sequences of CFTR from different species from the Uniprot database. For clarity, an 11 residue window with S492 (or equivalent) as the center of the window is shown. The serine at position 492 (or equivalent) is conserved in other species (red) with a substitution only to proline (green).

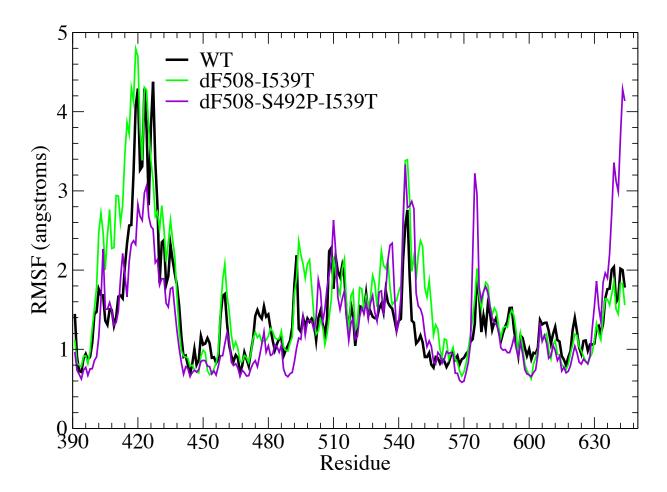


Figure S5. I539T mutation does not rescue NBD1 dynamics in Δ **F508-CFTR.** The I539T mutation, commonly used in the experimental setting to affect maturation of Δ F508-CFTR, does not return the profile of root mean square fluctuations (RMSF) (green) to wild type levels (black), especially in the dynamic regulatory insertion (RI) region, residues 404-435, and the critical central structure of the domain, residues 490-570. When combined with the S492P substitution (purple), these fluctuations are rescued.

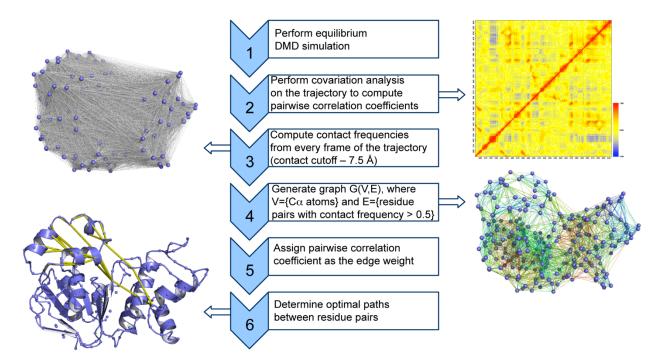


Figure S6. Computational methodology for determination of allosteric networks. The flow chart outlines the steps involved in determination of optimal paths between chosen sites in a given protein.

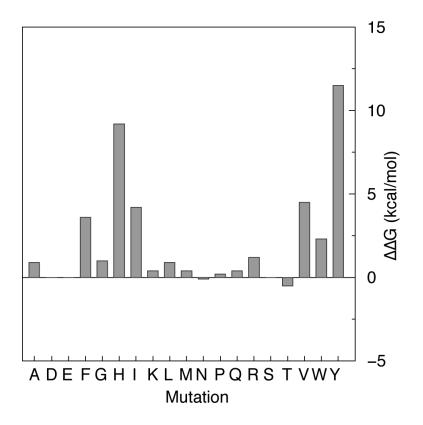


Figure S7. Exhaustive mutagenesis of S492 does not improve stability. Change in ΔG upon mutation of S492 to various amino acids. Negative values denote stabilizing mutations, while positive values denote destabilizing mutations. Bars represent $\Delta\Delta G$ values calculated from 500 independent iterations for every substitution.

-	Sheep			Horse			Shark			Frog		
	37°C		27°C	37°C		<u>27°C</u>	37°C		27°C	37°C		27°C
	WT	ΔF	ΔF	WT	ΔF	ΔF	WT	ΔF	ΔF	WT	ΔF	ΔF
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Figure S8. **Effect of F508 deletion on CFTR maturation in various species.** Western blots indicate the extent of maturation of WT and Δ F508-CFTRs from the species indicated in HEK-293 cells at 37°C or 27°C. The more rapidly migrating band (arrow) contains the core-glycosylated, endoplasmic reticulum localized form, and the more slowly migrating band contains CFTR with more complex oligosaccharides. The latter form of the shark protein (asterisk) features greater mobility than that of the other species due to the addition of one rather than two N-linked oligosaccharide chains. Like human Δ F508-CFTR, the two mammalian proteins mature at the lower but not at the higher temperature, whereas the non-mammalian orthologs mature at both temperatures.

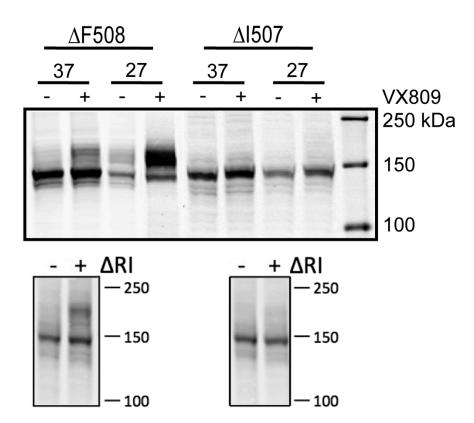


Figure S9. Δ I507-CFTR is not rescued by low temperature or VX809 corrector. While Δ F508-CFTR can be rescued by lowering temperature to 27 °C, treatment with VX809 corrector, or both, Δ I507-CFTR is not rescued by either of these treatments, alone or in combination.