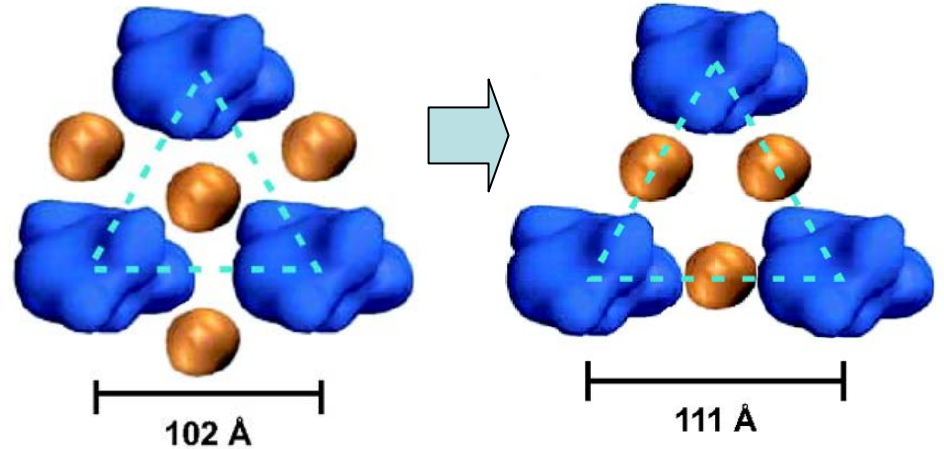
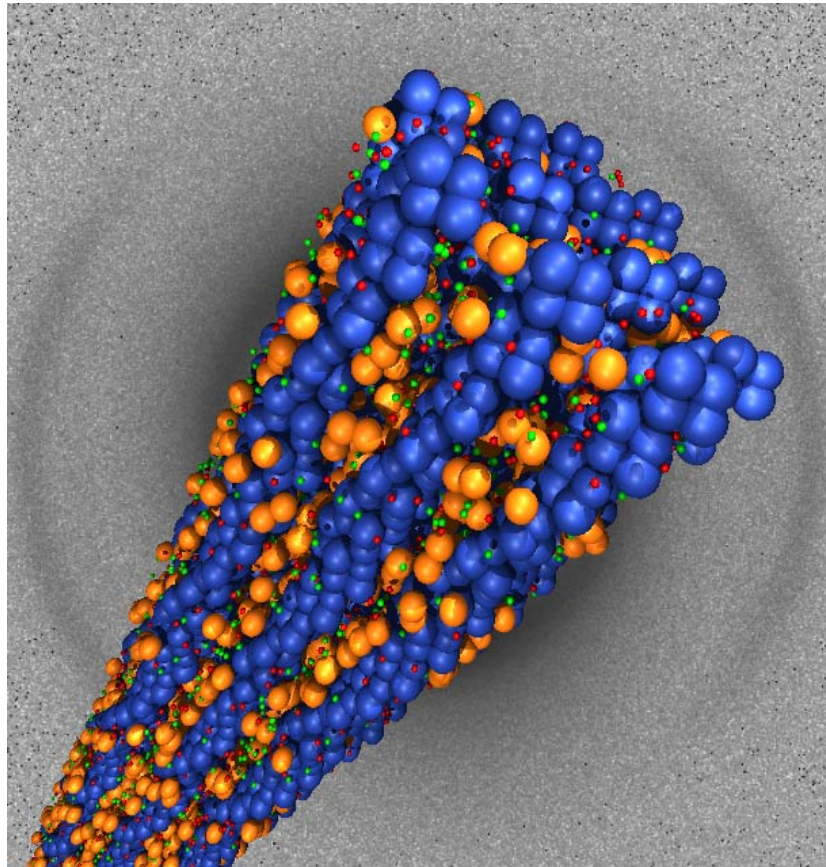


Actin-lysozyme complex in cystic fibrosis sputum



WT lysozyme (orange) and F-actin (blue) in end view. Stable structure, unaltered by mucin.

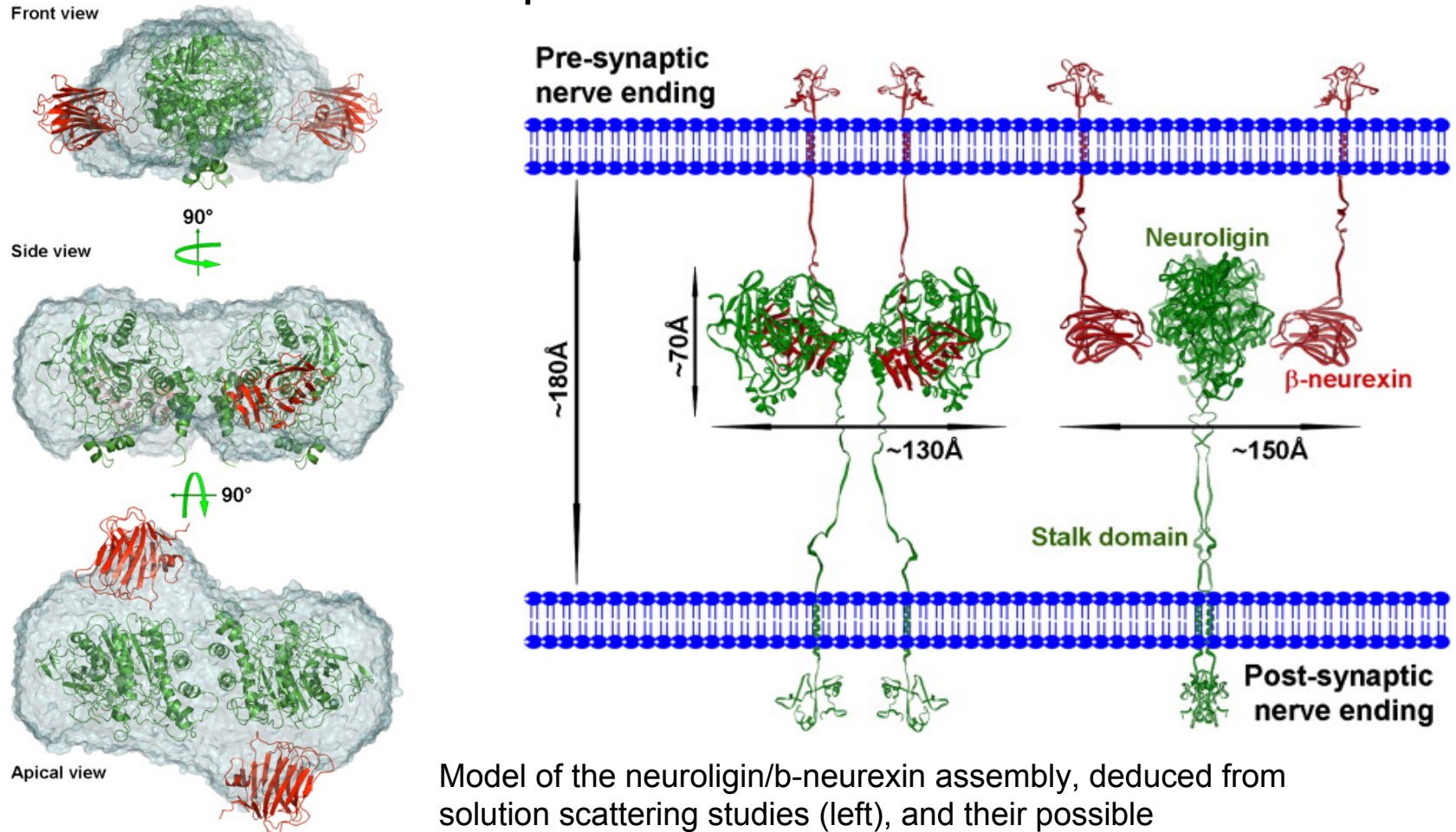
Charge-reduced lysozyme mutant (orange) and F-actin (blue) in end view. This structure is unstable.

This research confirms the existence of the lysozyme-actin complex in cystic fibrosis sputum and demonstrates that engineered lysozyme might prevent infected mucus molecules from forming ordered bundles that trap antimicrobials.

Sanders, L.K., *et al.*, PNAS 104, 15994-9 (2007)

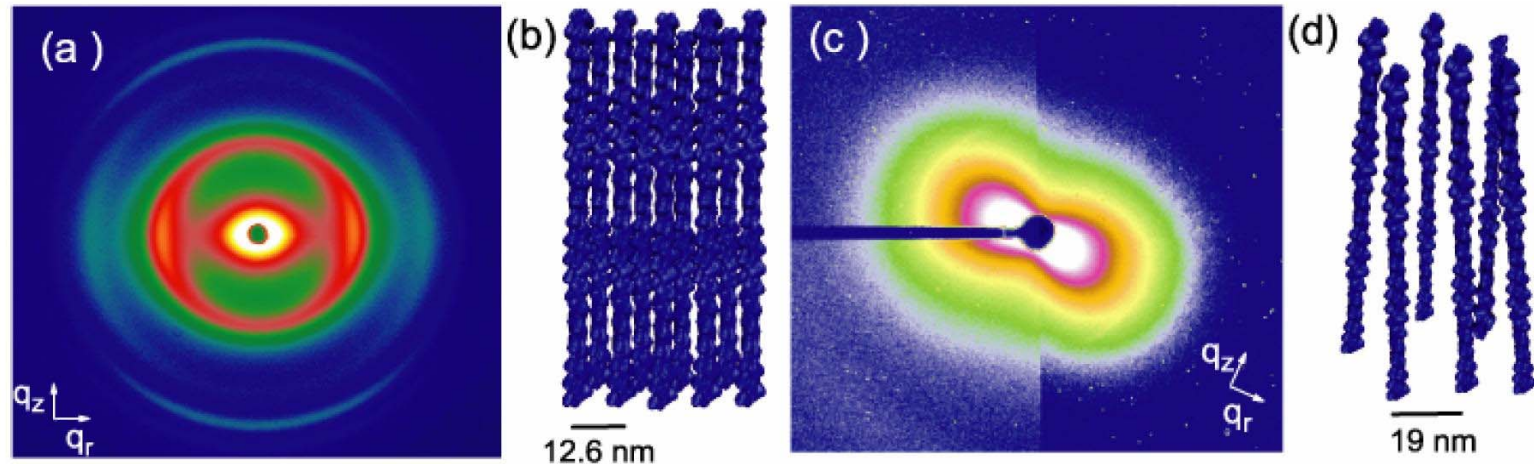
“Control of electrostatic interactions between F-actin and genetically modified lysozyme in aqueous media”

Molecular organization of neuroligin/neurexin complex in synapses. Implication in neurodevelopment disorders, including autism spectrum disorder.



Model of the neuroligin/ β -neurexin assembly, deduced from solution scattering studies (left), and their possible configuration in the synaptic space (right).

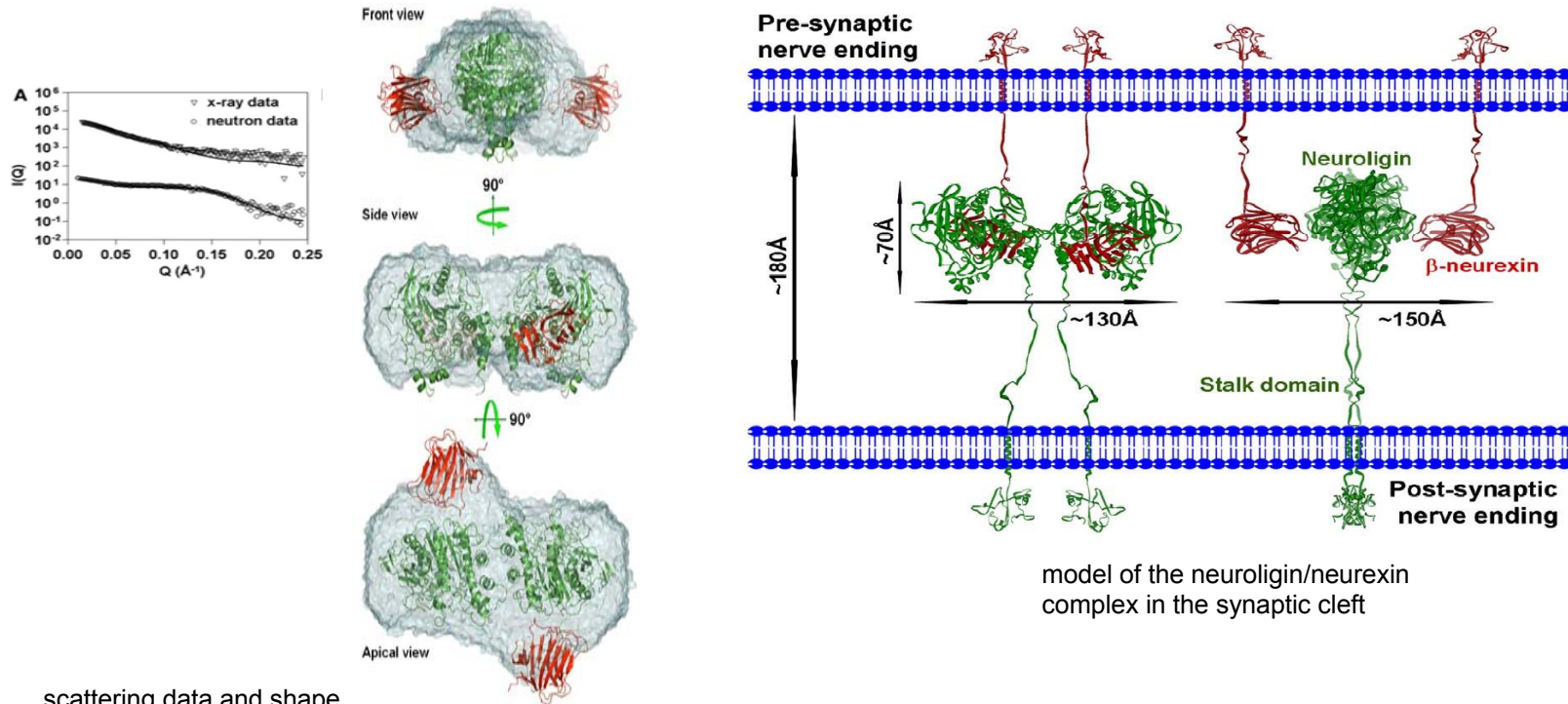
Structural Polymorphism of the Actin-Espín System: A Prototypical System of Filaments and Linkers in Stereocilia



Espíns are a type of cross-linking protein that are responsible for the formation of large parallel f-actin bundles. Scattering experiments have shown that normal espíns cause the actin to arrange in highly ordered hexagonally packed rigid bundles, whereas a deafness causing mutation of the espín is able to induce only a weak nematic order, resulting in bundles of much lower stiffness.

(a) hexagonally coordinated peaks are shown by packed F-actin-espín bundles (b);
(c) the scattering from the bundles with mutant espíns only show nematic order (d)

Synaptic Arrangement of the Neuroligin/b-Neurexin Complex Revealed by X-Ray and Neutron Scattering



scattering data and shape reconstruction of the neuroligin/neurexin complex

model of the neuroligin/neurexin complex in the synaptic cleft

The three dimensional solution structure of the neuroligin/neurexin complex, determined by SAXS and SANS methods, allowed for the development of a structural model of their spatial arrangement in the synaptic cleft. As mutations in the neuroligin and neurexin gene are linked to autism, this provides a structural framework for understanding altered recognition by these proteins in neurodevelopmental disorders.