Poster Presentations

[MS7-P08] NMR crystallography of membrane proteins <u>Anthony Watts</u>

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Solid state NMR methods are very versatile with respect to sample geometry and form1, and both amorphous and $(2D \text{ and } 3D^{1})$ crystalline materials can be studied to give high resolution spectra of use in deriving structural models. Being a short-Trange technique (dipolar interactions exist over Ås only), these crystals can be very small (nanoor amorphous), and certainly much smaller than can be manipulated for conventional single crystal diffraction approaches - powder samples still contain short-Trange order. Indeed, biomolecules that crystallize into subµm crystals but of little use for diffraction, can be ideal for study by solid state NMR.

Here some examples will be given of how we have devised methods and resolved highresolution structural information from both 2D and 3D crystals of membrane proteins^{2,3,4}. Both back-bone², and ligand/prosthetic group details^{5,6} have been obtained, usually ahead of the XRD or EM structures – some comparison will be presented where the crystal structures have been subsequently resolved.

References

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