

SEMINAR

Fast imaging - electrons or X-rays ?

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**PLEASE NOTE
CHANGE OF DATE,
TIME AND VENUE**

Wednesday 12 October, 2011

4.00pm – 5.00pm

Science Lecture Theatre S14, Building 29

Abstract

The relative strengths and weaknesses of X-ray and electron beams for fast, high-resolution imaging will be discussed, toward the goal of making a molecular movie. Aspects of the comparison include the following :

- i). The much higher scattering cross section of electrons, and resulting multiple scattering.
- ii) The existence of aberration-corrected lenses for electrons, providing sub-Angstrom probes.
- iii) Field-emission sources are brighter than current-generation synchrotrons. Our measurements of degeneracy for a field-emitter, and our laser-driven photofield fast source will be reviewed.
- iv) A cold field-emitter generates perhaps a hundred electrons per picosecond, insufficient to form an image. Hence the need for "stroboscopic" methods where a triggerable, repetitive process can be found. Multiple delays and projections are needed for a 3D movie. An X-ray free electron laser produces about $1E13$ photons in 20 fs, but practically all either do not interact with the sample, or are annihilated in production of damaging photoelectrons. Hard X-rays minimize multiple scattering. EELS allows parallel detection, impossible with X-rays
- v) The search for triggerable, repeating processes in materials science. First results from our time-resolved pump-probe diffraction at the LCLS free-electron hard X-ray laser will be shown.
- vii) Differences in time resolution between the methods. Current XFELs produce 10 fs pulses at a repetition rate of about 120 Hz, with correspondingly fast detector readout.
- viii) The effects of coulomb interactions on electron beams in producing unwanted energy spread and beam divergence, and the reduction of these effects at high energy due to relativistic effects.
- ix). The discovery that radiation damage can be avoided by using a sufficiently fast X-ray pulse which terminates before damage begins (1). (Can we outrun knock-on damage with electrons ?).
- x). XFEL diffraction from protein nanocrystals provides atomic resolution and femtosecond time resolution, opening up new possibilities for structural biology: (1) For solution of the phase problem (2). Single-shot patterns from many identical viruses in random orientations can be merged and reconstructed in 3D.
- xi) Is the use of scattering from many identical molecules, randomly oriented in solution, a better approach, in the light of the work of Z. Kam (4), which does not require modelling ?

Recent experimental results from the latest fast XFEL imaging of photosynthetic membrane proteins and the cathepsin enzyme will be reviewed (5). High-energy (3 MeV) fast electron diffraction results, with $7E5$ electrons per 100 fs pulse and a 500 micron diameter beam, will also be shown and discussed (6), in addition to electron diffraction movies of the CDW in TaS_2 .

1 H. Chapman et al. "Femtosecond protein nanocrystallography". Nature 470 73 (2011).

2 J. C. H. Spence et al. "Phasing XFEL nanocrystal data". Optics Express. 19, 2866 (2011).

3 M. Seibert et al "Single Mimivirus imaged on-the-fly with X-ray laser. Nature 470, 78 (2011)

4. D. Saldin et al . Phys Rev Letts. 106, 115501 (2011).

5 J.C.H. Spence. X-ray lasers for biology. Rep Prog Phys. (2012). In preparation.

6 Y. Murooka et. al. Appl Phys Letts 98, 251903 (2011)

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Visitors are most welcome: Please note that there is a designated Visitors Car Park (S2) clearly ground-marked by white paint and tickets, at a cost of \$3.50/hour for up to 3 hours, available from a dispensing machine. This high-rise car park is located on the following Clayton Campus Map, Ref. E3.