New Opportunities for Structural Biology Research at LCLS and SSRL Aina E. Cohen, representing the entire SSRL-SMB group

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Abstract.

Femtosecond crystallography (FX) is an emerging method that expands the structural information accessible from very small or very radiation sensitive macromolecular crystals. Utilizing extremely bright, short-time-scale X-ray pulses produced by an X-ray free electron laser (XFEL), this method exploits a 'diffraction before destruction' phenomenon where a still diffraction image is produced by a single X-ray pulse before significant radiation induced electronic and atomic rearrangements occur within the crystal. A goniometer-based experimental setup for FX experimentation is available to general users at the new Macromolecular Femtosecond Crystallography (MFX) instrument of the LCLS XFEL. LCLS-MFX began experiments on July 1, 2016. This instrument is based on developments at SSRL and LCLS-XPP to provide an efficient framework for goniometer-based FX experiments using automated strategies tailored to handle a variety of sample requirements, crystal sizes and experimental goals. The instrument has recently been upgraded to include support for fully automated sample exchange and data collection at room temperature and controlled humidity. These developments coupled with improvements in data processing algorithms make it possible to derive high resolution crystal structures using only 100 to 1000 still diffraction images. The MFX project and recent results using radiation sensitive crystals in limited supply and a variety of crystal delivery methods for serial diffraction data collection will be described.

New methods originally developed for serial diffraction experiments at XFELs, are proving valuable at synchrotron sources to study protein dynamics. Recent results using the MESH injector at the SSRL undulator micro-focus station BL12-2 will be described. Building on experiences at BL12-2, a next-generation undulator microfocus beam line, BL12-1, which will provide a preeminent capability for serial diffraction in the US, is under development. BL12-1 will be outfitted with a broad bandpass capability which will provide exceptional brightness, smaller microbeams and a high number of reflections when rastering crystals on the fly or using crystal injectors. It will be equipped with a high speed EIGER PAD detector and a high speed multi-axis goniometer, enabling new approaches for data collection and phasing. Similarities in instrumentation, existing and new sample delivery systems, and software environments will form the foundation of a synergistic relationship between the SSRL BL12-1 and a new Macromolecular Femtosecond crystallography instrument (MFX) at LCLS, through a Gateway approach.