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- Time-resolved research using fs laser as pump and SR as probe
- Source requirements
- Racetrack microtron design study at LBNL



- Non-equilibrium phase transitions: coherent phonons in InSb
- Warm dense matter: Al at high T, solid p
- Photogeneration of halogen radicals observed via time-resolved X-ray absoorption spectroscopy C. Bressler
- Dynamics in complex materials: polarons in  $Nd_{1/2}Sr_{1/2}MnO_3$
- Single x-ray pulse x-ray crystallography of CO-myoglobin, photoactive yellow protein, K. Moffat and M. Wolf (ESRF)
- X-ray diffraction of InSb undergoing a phase transition A. Rousse (ENSTA)
- Research programs at APS, UCSD, proposals at SLS, CLS

## Time-resolved x-ray diffraction of laser-excited InSb





- Atomic disordering on a time scale of a vibrational period ?
- Coherent acoustic phonons generation by ultrafast laser pulse (~100 fs)
- Lindenberg et al., Phys. Rev. Lett., **84**, 111, 2000

### Coherent acoustic phonons



- measured
- - calculated from dynamical diffraction

Fit parameters:

laser attenuation depth: 100 nm elect.-acoustic coupling time : 12 ps thermal strain: 0.17% max instantaneous strain: 0.08% max

![](_page_3_Picture_6.jpeg)

## Non-thermal melting

![](_page_4_Picture_1.jpeg)

![](_page_4_Figure_2.jpeg)

- Just below damage fluence (15 mJ/cm<sup>2</sup>)
- 3 ps decrease faster than
  acousto-electric response time
  (12 ps)
- Bond softening leads to nonthermal melting.

![](_page_5_Picture_1.jpeg)

- Warm dense matter is at high T ( $\sim$ 10,000 K) and solid  $\rho$ , at interface between solid state and plasma physics, these conditions exist in planets
- Measure Near-Edge X-ray Absorption (XANES) as probe of electronic structure
- Comparison and test of calculations being done at LLNL
- First system: aluminum, hydrodynamic properties well studied at high T

![](_page_5_Figure_6.jpeg)

# Structural evolution of simple photochemical reaction: I<sup>-</sup> photodetachment

![](_page_6_Picture_1.jpeg)

- Reaction:  $I^- + hv \rightarrow I + e_{sol} \text{ in } H_2O$ .
- Solvated electron has been extensively studied by femtosecond optical spectroscopy. Evolving structure of solute and solvent cage cannot easily be derived.
- Use x-ray absorption fine structure (XAFS) to determine time-resolved structure of solvent cage.

![](_page_6_Figure_5.jpeg)

![](_page_7_Picture_1.jpeg)

- Photon energy range: 100 eV to 10 keV
  - laser penetration depth  $\sim 0.1 \ \mu m$  while x-ray penetration depth increases with photon energy
- Repetition rate: 10 kHz
  - present state-of-art fs laser with mJ pulse energy
  - sample recovery, for semiconductor  $\sim 100$  ns
- Flux:  $\sim 10^{10}$  1/s 0.1% bw, Brightness:  $\sim 10^{14}$  1/s 0.1% bw
- Pulse duration: 100 fs, synchronized to a laser with same accuracy
  - period of typical molecular vibration
  - pulse duration of commercially available fs laser
- Tunable

# Dedicated facility for an ultra-fast x-ray spectroscopy

![](_page_8_Figure_1.jpeg)

• Racetrack microtron suits above defined goals at a relatively modest cost

• Electron bunch length is compressed from  $\sim 10$  ps to  $\sim 1$  ps before entering the last straight

• X-ray pulse is compressed from  $\sim 1$  ps to  $\sim 0.1$  ps in the beamline

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![](_page_9_Picture_1.jpeg)

X-ray pulse compression is due to a correlation between the longitudinal and transverse positions of electrons inside the electron bunch created by the RF orbit deflection in a cavity in the beginning of the final straight.

![](_page_9_Figure_3.jpeg)

An optical scheme for pulse compression with a collimating mirror and a double asymmetrically cut crystal monochromator

![](_page_9_Figure_5.jpeg)

![](_page_10_Picture_1.jpeg)

#### Undulator:

period = 14 mm, peak magnetic field = 1.5 T gap = 5 mm number of periods = 100

#### X-ray flux and brightness corresponding to the spontaneous undulator radiation at 9-th harmonics.

Beam energy, GeV	2
X-ray wavelength, Å	1.5
Flux, x-rays/sec/0.1% bw	$2 x 10^{10}$
Brightness, (usual units)	$2x10^{14}$
Pulse length (FWHM), fs	100