

Abstract Submitted  
for the MAR13 Meeting of  
The American Physical Society

**THz Microscopy of Anisotropy and Correlated Motions in Protein Crystals**<sup>1</sup> KATHERINE NIESSEN, GHEORGHE ACBAS, Department of Physics, SUNY at Buffalo, Buffalo, New York, EDWARD SNELL, Hauptman-Woodward Medical Research Institute and SUNY at Buffalo, Buffalo, New York, ANDREA MARKELZ, Department of Physics, SUNY at Buffalo, Buffalo, New York — We introduce a new technique, Crystal Anisotropy Terahertz Microscopy (CATM) which can directly measure correlated intra-molecular protein vibrations. The terahertz (THz) frequency range (5-100  $\text{cm}^{-1}$ ) corresponds to global correlated protein motions, proposed to be essential to protein function [1, 2]. CATM accesses these motions by removal of the relaxational background of the solvent and residue side chain librational motions. We demonstrate narrowband features in the anisotropic absorbance for hen egg-white lysozyme (HEWL) single crystals as well as HEWL with triacetylglucosamine (HEWL-3NAG) inhibitor single crystals. The most prominent features for the HEWL crystals appear at 45  $\text{cm}^{-1}$ , 69  $\text{cm}^{-1}$ , and 78  $\text{cm}^{-1}$  and the strength of the absorption varies with crystal orientation relative to the THz polarization. Calculations show similar anisotropic features, suggesting specific correlated mode identification is possible. 1. Hammes-Schiffer, S. and S.J. Benkovic, Relating Protein Motion to Catalysis. *Annu. Rev. Biochem.*, 2006. 75: p. 519-41. 2. Henzler-Wildman, K.A., et al., Intrinsic motions along an enzymatic reaction trajectory. *Nature*, 2007. 450(7171): p. 838-U13.

<sup>1</sup>This work supported by NSF MRI2 grant DBI295998.

Katherine Niessen  
Department of Physics, SUNY at Buffalo, Buffalo, New York

Date submitted: 09 Nov 2012

Electronic form version 1.4