ULTRAVIOLET RAMAN SPECTROSCOPY FOR CHARACTERIZING AMYLOID FIBRIL STRUCTURE AND DYNAMICS. <u>Igor K. Lednev</u>, Dmitry Kurouski and Ludmila Popova, Department of Chemistry, University at Albany, SUNY, 1400 Washington Ave., Albany, NY 12222.

Understanding fibrillogenesis at a molecular level requires detailed structural characterization of amyloid fibrils. We utilized the combination of deep UV resonance Raman (DUVRR) spectroscopy and postmortem hydrogen-deuterium exchange (HX) for probing parallel and anti-parallel β -sheet in fibrils prepared from full-length A β (1-40) and A β (34-42) peptides, respectively. The application of DUVRR spectroscopy allow us to discover a new protein folding/aggregation phenomenon, spontaneous refolding of amyloid fibrils. Mature fibrils prepared from apo- α -lactalbumin spontaneously refold from one polymorph to another as a result of a mild alteration in solution temperature and salinity. This discovery changes the very concept of the extraordinary stability of amyloid fibrils and presages a new approach for potentially regulating the biological activity of fibrils and their associated toxicity.