

Crystal Growth & Orientation in Nanoporous Polymer Monoliths

Benjamin Hamilton¹, Mike Ward², Marc Hillmyer (PI)³

¹University of Minnesota – Chemical Engineering & Materials Science, ²NYU - Chemistry, ³University of Minnesota – Chemistry

NNIN Facilities utilized: Characterization Facility & Nanofabrication Center

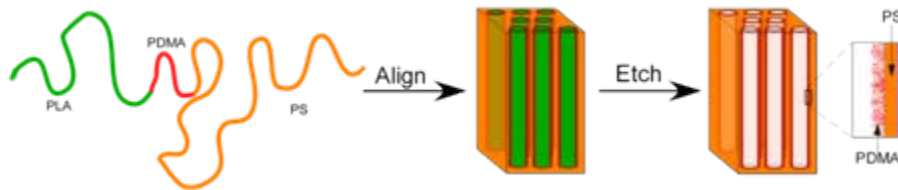
Research Interests

- ◆ polymorphism of glycine crystals confined to nanometer length scales
- ◆ porous polymer monoliths with well-ordered cylindrical pores as crystallization reactors
- ◆ glycine orientation within the well-ordered pores

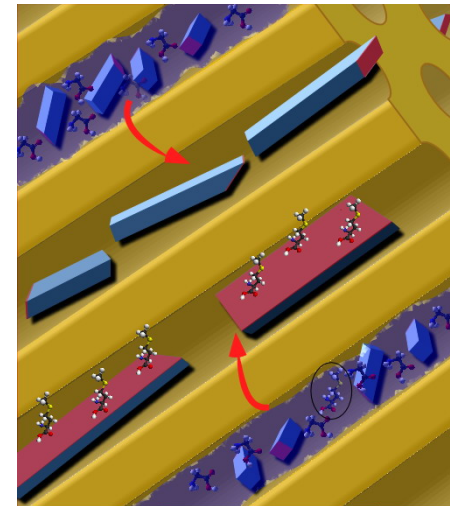
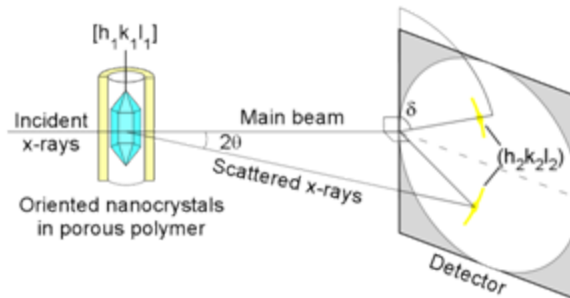
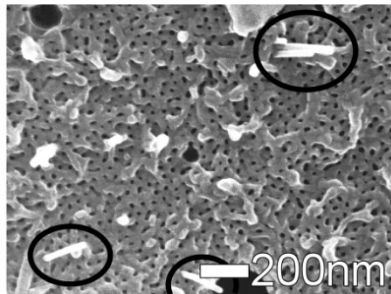
Results

- ◆ Macroscopically unfavored β -glycine crystals form preferentially within pores (vs. α in the bulk)
- ◆ Crystals adopt preferred orientation consistent with fastest growing direction parallel to pore direction
- ◆ Orientation can be altered with racemic auxiliaries chosen to inhibit the fastest-growth direction.

Porous Monoliths from PS-PDMA-PLA triblock terpolymers:



SEM, SAXS, and 2D- μ XRD performed at CharFac:



Publications

- ◆ Hamilton et. Al. *Cryst. Growth Des.* **2008**, 8, 3368-3375.
- ◆ Hamilton et. Al. *J. Am. Chem. Soc.* **2009**, 131, 2588-2596.