

## Beth Rhoades

Cornell NanoScale Science & Technology Facility  
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### Interests

The application of nanotechnologies to biology-based questions; nanomedicine, cytometry and microfluidics, self-assembling molecules and microbial structures.

### Education

BA in Biology, University of Colorado

Ph.D. in Microbiology, Colorado State University

Thesis: The Immunopathogenesis of Murine Pulmonary Tuberculosis.

### Professional

**2008 to present Biology liaison, Cornell NanoScale Science & Technology Facility.** Promote biology-driven nanoscale projects in the CNF and NNIN. Specifically provide support for biology-based users, promote educational programs in nanobiotechnology, support microfluidics, microscopy & printing processes, recruit new users from the medical community.

**2006 to 2008 Sr. Research Associate, Cornell** in the Dept. of Microbiology and Immunology conducting research on the inflammatory activity of mycobacterial cell walls. Experience in microbiology, immune cell biology, lipid biochemistry and flow cytometry.

**2005 Associate Scientist, Lovelace Respiratory Research Inst. Albuquerque, NM.**

Research in mycobacterial disease and adjuvant activity. Experience in establishing new biosafety level-2 and level-3 laboratories.

**2000 to 2004 Post-doc, Cornell University** in the lab of David Russell establishing murine models of inflammation. Experience in animal models and immunology protocols.

**1997 to 2000 Post-doc, Washington University, St. Louis, MO** in the lab of David Russell studying the release of bioactive glycolipids from intracellular mycobacteria. Experience in molecular cell biology.

### Recent Publications

Wang Z, Schwab U, Rhoades E, Chess PR, Russell DG, Notter RH. 2007. Peripheral cell wall lipids of *Mycobacterium tuberculosis* are inhibitory to surfactant function. *Tuberculosis* Dec 20 Epub.

Mwandumba HC, Squire SB, White SA, Nyirenda MH, Zijlstra EE, Molyneux ME, Russell DG, Rhoades ER. 2007. Alveolar macrophages from HIV-infected patients with pulmonary tuberculosis retain the capacity to respond to stimulation by LPS. *Microbes Infect.* 9:1053-60.

Hsu FF, Turk J, Owens RM, Rhoades ER, Russell DG. 2007. Structural characterization of phosphatidyl-myo-inositol mannosides from *Mycobacterium bovis* Bacillus Calmette Gérin by multiple-stage quadrupole ion-trap mass spectrometry with electrospray ionization. II. Monoacyl- and diacyl-PIMs. *J Am Soc Mass Spectrom.* 18:479-92.

Howard ST, Rhoades E, Recht J, Pang X, Alsup A, Kolter R, Lyons CR, Byrd TF. 2006. Spontaneous reversion of *Mycobacterium abscessus* from a smooth to a rough morphotype is associated with reduced expression of glycopeptidolipid and reacquisition of an invasive phenotype. *Microbiology* 152:1581-90.

Geisel RE, Sakamoto K, Russell DG, Rhoades ER. 2005. In vivo activity of released cell wall lipids of *Mycobacterium bovis* bacillus Calmette-Guérin is due principally to trehalose mycolates. *J Immunol* 174:5007-15.