**General Position Description:**
The Regulatory Pharmaceutical Fellowship is a two-year, learning based program designed to train pharmacists for careers in government, academia, or the pharmaceutical industry. Each fellow will spend a 9-month rotation at the Food and Drug Administration’s (FDA) Center for Drug Evaluation and Research (CDER) (Silver Spring, MD); a 6-month rotation with Purdue University College of Pharmacy (Indianapolis, IN); and a 9-month rotation at Eli Lilly and Company (Indianapolis, IN) or Johnson & Johnson PRD L.L.C. (Titusville, NJ). The 2011-2013 cycle offers two tracks: Drug Information, and Drug Advertising and Promotion. Deliverables may include research projects, publications, poster presentations, summaries of experiences gained, a teaching portfolio, continuing education presentations, and regulatory projects. Chosen candidates will have various educational experiences in cross-functional departments and have opportunities to publish articles and/or give presentations. Qualified candidates must be a graduate from an ACPE-accredited college of pharmacy, or otherwise eligible for licensure, prior to the start of the fellowship term.

The Fellowship begins in July 2011 and continues until June 2013, with a competitive annual stipend and benefits.

**Program Participants**
**Drug Information:** FDA's CDER Division of Drug Information, Purdue University, and Eli Lilly and Company

**Advertising and Promotion:** FDA's CDER Division of Drug Marketing, Advertising, and Communications, Purdue University, and Johnson & Johnson PRD L.L.C.

For more information about the program, please visit: [http://www.fda.gov/RegPharmFellowship](http://www.fda.gov/RegPharmFellowship)

We encourage interested candidates to forward their CVs to our current fellows for on-site interview consideration during the upcoming ASHP Midyear Clinical Meeting:
Drug Information Fellow, Kimberly Wu at DrugInformationFellowship@gmail.com
Drug Advertising and Promotion Fellow, Nital Patel at DrugMarketingFellowship@gmail.com.
Thank you for your time and consideration.

Best regards,
Sadhna Khatri, MS, PharmD On behalf of:
Catherine Y. Chew, PharmD
CDR, U.S. Public Health Service
Team Leader, Division of Drug Information
Food and Drug Administration Center for Drug Evaluation and Research
Director, Fellowship Program