Butler University

PHARMACEUTICAL FELLOWSHIP PROGRAM

 $Fellowship \\ Application$



One-Year Fellowship Tracks

ACADEMIC: Butler University

Longitudinal Experience

INDUSTRY: Recordati Rare Diseases Inc. July 2025–June 2026

Not recruiting for the 2025-2026 cycle

Two-Year Fellowship Tracks

ACADEMIC: Butler University

July 2025-October 2025

INDUSTRY: Partner (Lilly or

Regeneron)

 $November\,2025-October\,2026$

REGULATORY: FDA

November 2026-June 2027

Fellowship Director

John B. Hertig, PharmD, MS, CPPS, FASHP, FFIP

Associate Professor
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"What we love most about our fellowships are the multitude of experiences in academia and the pharmaceutical industry that will provide a comprehensive outlook on medical innovation and outcomes. After completion of our fellowships, we will be equipped with all of the experiences necessary to provide optimal advances in healthcare with a patient-centric approach." - Butler second-year fellows Rachel Matthews and Hannabeth Owens, and first-year fellow Angie Neddeau

The Butler University Pharmacy Fellowships Program has three positions available with unique focus areas. All fellows will learn best practices in teaching and education from award-winning faculty at Butler University in addition to training with leading experts in the pharmaceutical industry. Within our two-year track, fellows will also gain valuable experience at the USFDA. Fellows in this program receive a truly unique experience, preparing them to follow their passions while becoming leaders in their respective fields.



GLOBAL PATIENT SAFETY and PHARMACOVIGILANCE

As a fellow at Eli Lilly & Company, you will have opportunities to work crossfunctionally on pre-and post-marketed compounds to assess and evaluate adverse events in the context of biological plausibility, indication for use, concomitant medications, and comorbid conditions to determine if the adverse events are drug-related. You will actively participate in various aspects of pharmaceutical



innovation, use optimization, and global safety.

ELI LILLY and COMPANY FELLOW Jermiya Jackson

REGENERON

MEDICATION ERROR PHARMACOVIGILANCE and RISK MANAGEMENT

At Regeneron, you will be a part of a leading biotech company in evaluating safety and mitigating risk. The opportunity to work on numerous medications in the pipeline ranging from eye disease to metabolic disease will provide you exposure to multiple therapeutic areas. The range of experiences will also allow you to work



cross-functionally as a team to promote medication safety.

REGENERON FELLOW Sarah Leonard



MEDICAL INFORMATION and MEDICAL AFFAIRS

During your fellowship at Recordati
Rare Diseases Inc., you will collaborate
cross-functionally to create assets
aligned with our medical strategy and
work on initiatives related to Medical
Affairs operational excellence across
therapeutic areas. You will develop
a strong foundation in Medical
Information and broad awareness of
Medical Affairs operations while serving



the underserved rare disease communities of North America.

RECORDATI FELLOW Angie Neddeau

Not actively recruiting for the 2025-2026 cycle

The Office of Medication Error Prevention and Risk Management's (OMEPRM) mission is to increase the safe use of drug products and improve public health by minimizing use error related to the naming, labeling, packaging, or design of drug products; and developing effective and efficient Risk Evaluation and Mitigation Strategies (REMS) for certain drug products that ensure the benefits outweigh its risks. You will actively participate in a range of experiences within OMEPRM to promote medication safety.