# REGULATORY PHARMACEUTICAL FELLOWSHIPS

Government
Industry
Academia

2024-2026

Jointly sponsored by:

FDA

janssen
PHARMACEUTICAL COMPANIES OF Johnson&Johnson

abbvie

Johnson&Johnson

Lilly

PURDUE UNIVERSITY

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# FELLOWSHIP OVERVIEW

## PROGRAM DESCRIPTION

The purpose of the Regulatory Pharmaceutical Fellowship program is to train selected candidates in one of three tracks focused on drug information, medication safety, or regulatory advertising & promotion. The program provides participants with the unique opportunity to learn from mentors in their chosen specialty track across three diverse settings in government, academia, and industry. Graduates of the fellowship program are qualified to pursue careers in any of the three practice settings.

## FELLOWSHIP BENEFITS

- Competitive stipend
- Reimbursement for relocation during fellowship and professional travel expenses
- Enrollment in the Indiana Pharmacy Teaching Certificate (IPTeC) Program
- Vacation and University holidays
- Optional Purdue University benefits package (health, Rx, vision, and dental)

## 2024-2026 RECRUITMENT

**Drug Information**
2 Positions Available

**Medication Safety**
1 Position Available

## ELIGIBILITY REQUIREMENTS

The fellow must be a graduate from an ACPE-accredited college of pharmacy, or otherwise eligible for licensure as a pharmacist, prior to the start of the fellowship term. All applicants must be U.S. citizens.

# APPLICATION PROCESS

**All interested applicants must submit the following via the online portal:**

1. Letter of intent, specifying track and sponsor(s) of interest
2. Contact information for three references, including name, email, and phone
3. Curriculum Vitae
4. Official transcripts

Please address all fellowship application materials to the Regulatory Pharmaceutical Fellowship Program Director, Dr. Amy Sheehan.

### DEADLINE

All application materials should be submitted to the online portal below no later than 11:59 pm EST on November 13th, 2023

### FINAL INTERVIEWS

Final interviews will be conducted virtually starting the week of November 27th

### PRELIMINARY INTERVIEWS

Preliminary interviews will be conducted virtually

Starting the week of October 30th for the Med Safety Track

Starting the week of November 14th for the Drug Info Track

Recognizing that the choice of a Post-Doctoral Industry Fellowship is an important decision, Purdue University College of Pharmacy, in conjunction with the Academic Industry Fellowship Alliance (AIFA), has agreed to extend offers for Fellowships no earlier than December 13th, 2023.

APPLICATION PORTAL: https://hiring.science.purdue.edu/pharm

PORTAL OPENS **OCTOBER 1ST, 2023** FOR THE 2024-2026 RECRUITMENT CYCLE

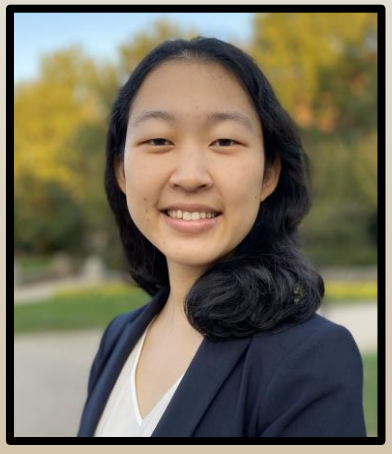
Additional fellowship opportunities are available through Purdue University. Visit our website at: https://www.phpr.purdue.edu/fellowships/pharmaceutical-industry-affiliated-fellowships

# CURRENT FELLOWS

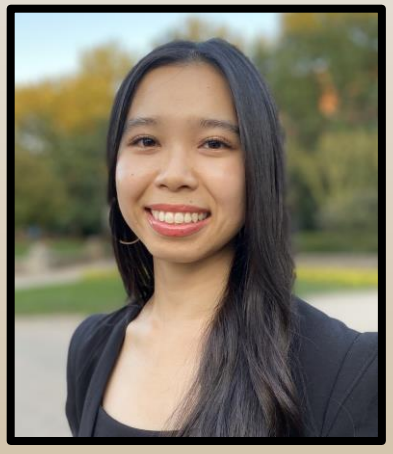
KORI ADAIR
Drug Information

KYOUNGEUN KIM
Drug Information

JULIANNA HICKEY
Drug Information



ANLY LI
Drug Information



MICHELLE LE
Medication Safety

NIKHILA VISWANATHAN
Medication Safety

SARAH TANIOS
Medication Safety

RICHARD LEE
Advertising & Promotion

# DRUG INFORMATION

## PROGRAM OVERVIEW

**ACADEMIA**

The Purdue University experience exposes the fellow to academia and an institutional-based drug information center at Indiana University Health. Fellows will conduct a research project for presentation at a national meeting and publication in a peer-reviewed pharmacy journal. Significant teaching experience in and out of the classroom is provided.

**West Lafayette & Indianapolis, IN**

**FDA**

The FDA experience provides an opportunity for fellows to refine their drug information skills in a regulatory setting. Fellows respond to drug information inquiries from patients, health care professionals, and regulated industry; create and disseminate content; and assist with a number of high-profile initiatives.

**Silver Spring, MD**

**INDUSTRY**

This experience provides the opportunity for the fellow to gain training as a member of a medical information team in the pharmaceutical industry. The fellow will develop skills related to development and delivery of drug information to healthcare professionals, payors, consumers, and business partners through a variety of methods.

**See locations on Page 7**

## CURRENT FELLOWS

**Kyoungeun Kim, PharmD**
Industry Sponsor:
Eli Lilly and Company
Second Year Fellow: 2022-2024 Cycle
Graduated from University of Illinois Chicago

**Anly Li, PharmD**
Industry Sponsor:
Eli Lilly and Company
First Year Fellow: 2023-2025 Cycle
Graduated from the University of Michigan College of Pharmacy

**Kori Adair, PharmD**
Industry Sponsor:
Janssen Scientific Affairs, LLC
Second Year Fellow: 2022-2024 Cycle
Graduated from Texas Tech University

**Julianna Hickey, PharmD, MPH**
Industry Sponsor:
Janssen Scientific Affairs, LLC
First Year Fellow: 2023-2025 Cycle
Graduated from Campbell University College of Pharmacy & Health Sciences

# DRUG INFORMATION

## FELLOWSHIP SCHEDULE: 2 POSITIONS, 1 FELLOW EACH

POSITION 1

| PURDUE UNIVERSITY | FDA | ELI LILLY AND COMPANY |
|---|---|---|
| College of Pharmacy | Division of Drug Information | Global Medical Information Indianapolis, IN |
| **6 months: 07/2024 – 12/2024** | **6 months: 01/2025 – 06/2025** | **12 months: 07/2025 – 06/2026** |

POSITION 2

| PURDUE UNIVERSITY | JANSSEN PHARMACEUTICALS | FDA |
|---|---|---|
| College of Pharmacy | Medical Information & Knowledge Integration Horsham, PA | Division of Drug Information |
| **6 months: 07/2024 – 12/2024** | **12 months: 01/2025 – 12/2025** | **6 months: 01/2026 – 06/2026** |

**Anly Li**

First Year Drug Information Fellow

"Through this fellowship, I have had multiple opportunities to advance my drug information and writing skills through experiences answering drug information questions, teaching and precepting pharmacy students, and leading a research project. I am grateful for the support of my preceptors and co-fellows, and I am excited to continue learning more about the role of drug information in academia, regulatory, and industry settings."

**Julianna Hickey**

First Year Drug Information Fellow

"I chose this fellowship with Purdue University, Janssen Scientific Affairs LLC, and the FDA because of the unparalleled and diverse drug information experiences it would provide. So far, I have enjoyed growing professionally in an academic setting and helping pharmacy students learn about the importance of drug information."

# DRUG INFORMATION

## PROGRAM PRECEPTORS

**Amy H. Sheehan, PharmD | Purdue University**
hecka@purdue.edu

For over twenty years, Amy has worked with Purdue University College of Pharmacy and the Indiana University (IU) Health Center for Medication Management, where she contributes to the provision of comprehensive drug information services for all IU Health-affiliated hospitals. She has published over 40 peer reviewed articles in the pharmacy literature and authored book chapters for therapeutics and drug information textbooks. Amy serves on the Indiana Pharmacy Teaching Certificate (IPTeC) Program Executive Committee and is currently a member of the Editorial Board for the Annals of Pharmacotherapy. She serves as course coordinator for a drug information and advanced literature evaluation class.

**Megan (Cuomo) Walter, PharmD | FDA**
Past Fellow 2018-2020 Cycle
Megan.Walter@fda.hhs.gov

Megan Walter received her BS and PharmD degrees from the University of North Carolina Eshelman School of Pharmacy. Upon graduation, she completed the Regulatory Pharmaceutical Fellowship in Drug Information with Purdue University, Janssen Scientific Affairs, LLC, and FDA. After completing the fellowship, Megan accepted a position as a pharmacist in the Division of Drug Information at FDA. Megan answers drug information questions from consumers, healthcare professionals, and the pharmaceutical industry.

**Sara Roach, PharmD | FDA**
Sara.Roach@fda.hhs.gov

Sara Roach is a pharmacist in the FDA's Division of Drug Information where she provides timely, accurate, and useful information to consumers, health care professionals, and industry. Sara is part of the REMS@FDA database team that updates and manages the FDA's public REMS database. Also, as a member of DDI's Podcast team, Sara assists in producing various podcast series for the Center for Drug Evaluation and Research. Sara enjoys precepting fellows and pharmacy students, as well as mentoring new staff. She holds a Doctor of Pharmacy degree from Wingate University School of Pharmacy.

**Kathy Mybeck, PharmD | Eli Lilly and Company**
Mybeck_Kathy@lilly.com

Throughout her 20+ year career at Eli Lilly and Company, Kathy has held various roles within Medical Information and Regulatory Affairs. Kathy has provided global medical information support and led development of launch portfolio medical information responses and strategies for endocrine and oncology products. Kathy also served as an Implementation Lead for the Regulatory Transformation initiative by partnering with the Labeling department on process and system updates. Kathy continues to coach her team members on medical information-related activities and precept PharmD students from a variety of schools.

**Samina Ali, PharmD | Janssen Scientific Affairs, LLC**
sali2@its.jnj.com

Samina Ali earned her BS and PharmD degrees from Rutgers University College of Pharmacy and completed a Pharmacy Practice Residency at Mount Sinai Medical Center in New York City. Samina has led and supported Medical Information activities for multiple Janssen products in oncology, diabetes, virology, GI, women's health and urology. She is currently responsible for strategy and review of scientific content, promotional and sales training materials for prostate cancer products.

# MEDICATION SAFETY

## PROGRAM OVERVIEW

**ACADEMIA**

The fellow will have unique experiences in academia, such as publishing original research, delivering presentations, and teaching. The fellow will actively participate in practice-based research to foster the discovery and delivery of information and practices to enhance medication safety.

**West Lafayette & Indianapolis, IN**

**INDUSTRY**

The fellow at AbbVie will gain experience in safety signal detection and evaluation, as well as the development of regulatory documents. The fellow will actively contribute to team decisions on the collection, detection, assessment, monitoring, and prevention of adverse events for pre- and post-marketed compounds.

**North Chicago, IL**

**FDA**

The fellow will participate in intra- and inter-center projects in pre- and post-market arenas. The fellow will focus on research opportunities to contribute to the review, development, modification, and assessment of a Risk Evaluation and Mitigation Strategy (REMS) for prescription drugs and biologics in accordance with the current FDA standards.

**Silver Spring, MD**

## FELLOWSHIP SCHEDULE: 1 POSITION, 1 FELLOW

| PURDUE UNIVERSITY | ABBVIE | FDA |
|---|---|---|
| College of Pharmacy | Pharmacovigilance and Patient Safety<br>North Chicago, IL | Division of Risk Management (DRM)<br>Silver Spring, MD |
| **4 months:<br>07/2024 – 10/2024** | **12 months:<br>11/2024 – 10/2025** | **8 months:<br>11/2025 – 06/2026** |

**Sarah Tanios**

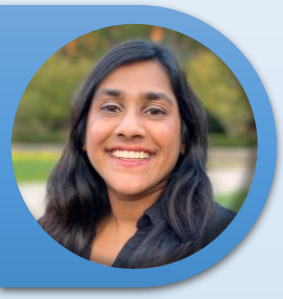
First Year Medication Safety Fellow

"I chose this particular fellowship because of its perfect duality as a well-rounded experience that is also incredibly immersive. It is rare to come across a program like this one that provides deep engagement in not only industry, but also FDA/regulation, and academia. From my fellowship experiences so far and those upcoming, it is clear that this program provides excellent structure to becoming a versatile pharmacovigilance and medication safety professional."

# CURRENT FELLOWS

**Michelle T. Le, PharmD**
Industry Sponsor:
AbbVie
Second Year Fellow: 2022-2024 Cycle
Graduated from Chapman University

**Sarah Tanios, PharmD**
Industry Sponsor:
AbbVie
First Year Fellow: 2023-2025 Cycle
Graduated from Midwestern University
Chicago College of Pharmacy



**Nikhila Viswanathan, PharmD, MS**
Industry Sponsor:
Novartis
Second Year Fellow: 2022-2024 Cycle
Graduated from Temple University

# PROGRAM PRECEPTORS

**Kyle Hultgren, PharmD | Purdue University**
khultgre@purdue.edu

Kyle Hultgren is the director of the Purdue University College of Pharmacy's Center for Medication Safety Advancement. Dr. Hultgren's current research includes extensive work on dashboards and measurement systems for the evaluation and improvement of medication use systems as well as large adverse event data set analysis. He holds multiple copyrights on mobile computer simulation technology and two patents on medical devices designed to improve patient safety. Dr. Hultgren lectures nationally and internationally on safe medication use practices and teaches regularly in the Doctor of Pharmacy curriculum where he was the 2016 Dr. Aziz Teaching Award recipient.

**Dan Degnan, PharmD, MS, CPPS, FASHP | Purdue University**
ddegnan@purdue.edu

Dan Degnan currently serves as the Associate Director for the Professional Skills Laboratory and is a Clinical Assistant Professor of Pharmacy Practice (Courtesy) at the Purdue University College of Pharmacy. He has an appointment at the Regenstrief Center for Healthcare Engineering at Purdue as a Clinical Research Associate with expertise and research interests in the area of medication safety technology, advanced pharmacy automation, pharmacy operations and high reliability in healthcare. Prior to his role at Purdue, Dr. Degnan served as the Medication Safety Officer at Community Health Network in Indianapolis for almost 10 years. He has held leadership positions in national organizations for both pharmacy and healthcare quality. He is the 2022 recipient of the Dr. Aziz Outstanding Teacher of the Year at the College of Pharmacy.

**Ephrem Abebe, BSPharm, MPharm, MS, PhD | Purdue University**
eabebe@purdue.edu

Ephrem Abebe is an Assistant Professor in the Department of Pharmacy Practice, College of Pharmacy, Purdue University. He teaches in and co-coordinates the Patient Safety and Informatics course for third year pharmacy students. He also mentors graduate students pursuing MS and PhD degrees in the Health Services, Outcomes, and Policy (HSOP) graduate program in the Department of Pharmacy Practice. Dr. Abebe leads a highly interdisciplinary, federally-funded, research program that leverages approaches from health services research, implementation science, human factors engineering, and participatory design. A core focus of Dr. Abebe's research program is related to understanding and supporting the longitudinal medication use experience of patients and their family caregivers as they journey through the healthcare system. Dr. Abebe was born and raised in Ethiopia. He is passionate about mentorship and professional development of trainees and strives to create a safe and welcoming research and learning environment for all individuals.

# MEDICATION SAFETY

## PROGRAM PRECEPTORS

### Adrienne M. Rothstein, PharmD | AbbVie

Adrienne.Rothstein@abbvie.com

Adrienne Rothstein is a Scientific Director at AbbVie, focused on late stage oncology products. Adrienne received her BS in Pharmacy from St. John's University and her PharmD from the University of Cincinnati. After graduation, she pursued a pharmacy practice residency at Stanford University Hospital and then worked in drug information and pharmacovigilance at Elan Pharmaceuticals. Adrienne later worked at the FDA for 10 years in the Office of Surveillance and Epidemiology, Division of Pharmacovigilance and the Office of New Drugs, Division of Reproductive, Urology and Bone Products. After joining AbbVie in 2014, Adrienne led a team of safety scientists responsible for safety surveillance for oncology and immunology products (development and post-approval), before her current role as a Scientific Director to monitor product safety, deliver presentations, set strategy and author/edit a variety of regulatory submissions for AbbVie's oncology products.

### Alexandra Terry, PharmD, BCPS | AbbVie

Alexandra.Terry@abbvie.com

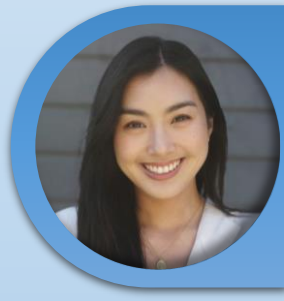
Alexandra Terry is an Associate Director in Safety Data Sciences at AbbVie. Her current role within the Pharmacovigilance and Patient Safety organization involves developing and authoring surveillance strategies, signal evaluations, aggregate safety reports, and global submission activities. She supports a variety of investigational and post-marketed product teams in the oncology and neuroscience therapeutic areas. Alexandra received her Doctor of Pharmacy degree from University of Illinois at Chicago. Prior to joining AbbVie, she completed a Visiting Scientist Fellowship at Eli Lilly and a PGY-1 pharmacy practice residency at OSF Saint Francis Medical Center.

### Charlotte Moureaud, PharmD | AbbVie

Past Fellow 2019 - 2021 Cycle

Charlotte.Moureaud@abbvie.com

Charlotte Moureaud is an Associate Director in Safety Data Sciences at AbbVie. She received her Doctor of Pharmacy degree from Rutgers University, Ernest Mario School of Pharmacy. Following graduation, she completed the Regulatory Pharmaceutical Fellowship in Medication Safety with Purdue University, AbbVie, and FDA. During that time, she obtained her MBA. After completing the fellowship, Dr. Moureaud rejoined the team at AbbVie. She supports oncology products in both late development and post-approval and contributes to signal strategy and assessments. She also strategizes and authors global safety documents submitted to regulatory authorities and supports submission activities.



### Elisa Basa, PharmD | AbbVie

Elisa.Basa@abbvie.com

Elisa Basa is an Associate Director in Safety Data Sciences (SDS) at AbbVie. She received her Doctor of Pharmacy degree from the University of the Pacific Thomas J. Long School of Pharmacy, through the Pre-Pharmacy Advantage program. Following graduation, Elisa worked in retail pharmacy prior to joining AbbVie, where she supported both early and late development oncology products for 4 years before transitioning to support the immunology therapeutic area. As part of her current role within Pharmacovigilance & Patient Safety, Epidemiology, R&D Quality Assurance (PSEQ), Elisa conducts standard safety surveillance activities and collaborates cross-functionally to strategize, review, author, and support activities such as ad hoc safety requests, regulatory submissions, and aggregate safety reports. Elisa has also been a preceptor for Advanced Pharmacy Practice Experiences (APPE) students and Summer SDS Interns at AbbVie.

### Cynthia L. LaCivita, PharmD | FDA

Cynthia.LaCivita@fda.hhs.gov

Cynthia LaCivita is the Director of Risk Management (DRM) in Office of Medication Error Prevention and Risk Management in the Center for Drug Evaluation and Research at the Food and Drug Administration (FDA). DRM provides risk management expertise on design, development, and implementation of programs and initiatives to support the Center's policies related to Risk Evaluation and Mitigation Strategies (REMS) authorities under the Food and Drug Administration Amendments Act (FDAAA) of 2007. Prior to joining the FDA she was an assistant professor in the Department of Pharmacy Practice and Science at the University of Maryland School of Pharmacy and assistant professor of Oncology at University of Maryland Cancer Center, Director of Clinical Standards and Quality for the American Society of Health-System Pharmacists (ASHP) and Director of Education and Special Programs for the ASHP Research and Education Foundation.

NOT RECRUITING

# ADVERTISING AND PROMOTION

## PROGRAM OVERVIEW

**INDUSTRY**

The fellow will develop an understanding of FDA regulations and guidance, industry codes, and more, as related to prescription drug advertising and promotional activities. The fellow will work directly with internal business partners such as marketing, legal, medical, and others to ensure that a broad range of promotional materials are in compliance with regulations and internal policies.

**Titusville, NJ**

**FDA**

The fellow will gain experience in the FDA promotional review process and provide overviews of relevant laws and FDA guidance documents. The fellow will assist in reviewing promotional materials, evaluating draft product labeling, researching and evaluating industry complaints, and working with other functions in the Office of Prescription Drug Promotion (OPDP).

**Silver Spring, MD**

**ACADEMIA**

The fellow will gain exposure to upper-level academia and the different responsibilities of academic administrators. The fellow will gain significant teaching experience through provision of didactic education including coordination of a core PharmD management and marketing course and the experiential training of pharmacy students.

**West Lafayette & Indianapolis, IN**

## FELLOWSHIP SCHEDULE: 1 POSITION, 1 FELLOW

| Johnson & Johnson | FDA | Purdue University |
|---|---|---|
| **9 months** | **9 months** | **6 months** |
| Regulatory Advertising & Promotion | Office of Prescription Drug Promotion | College of Pharmacy |

**Richard Lee**

First Year Advertising and Promotion Fellow

"This fellowship provides an extremely unique opportunity to gain experience in regulatory advertising and promotion in both the pharmaceutical industry and the FDA's Office of Prescription Drug Promotion (OPDP). These differing perspectives allow me to develop the strong foundation necessary to contribute to the development of compliant and effective promotional materials that are accurate, fair-balanced, and truthful/non-misleading. Additionally, teaching experiences at Purdue University's College of Pharmacy strengthen my abilities to communicate, educate and lead - key attributes needed to succeed in a variety of settings."

# CURRENT FELLOW

**Richard Lee, PharmD, MS**
Industry Sponsor:
Johnson & Johnson
First Year Fellow: 2023-2025 Cycle
University of Maryland School of Pharmacy

# PROGRAM PRECEPTORS

**Sheetal Patel, PharmD | Johnson & Johnson**
Past Fellow 2007 - 2009 Cycle
spate120@its.jnj.com

Sheetal Patel is the Head, Compliance & Regulatory Advertising & Promotion, within Pharmaceutical Group Health Care Compliance organization at Johnson & Johnson. Previously, she held the position of Lieutenant Commander, Senior Regulatory Review Officer, at the U.S. Food and Drug Administration, Office of Prescription Drug Promotion.

**Sheela Trivedi, PharmD, CDCES | Johnson & Johnson**
strived6@its.jnj.com

Sheela Trivedi is an Associate Director in the Regulatory Advertising & Promotion (RAP) for the Pharmaceutical sector within the Johnson & Johnson Healthcare Compliance organization. She earned her PharmD from the Philadelphia College of Pharmacy and completed an Ambulatory Care Clinical Pharmacy Practice Residency at the Philadelphia Veterans Affairs Medical Center. Prior to joining RAP, she worked as a Clinical Pharmacy Specialist within the U.S. Department of Veterans Affairs and part of the Medical Information & Knowledge Integration team within Janssen Scientific Affairs. She is also a Certified Diabetes Care and Education Specialist.

**Sam Skariah, PharmD, RAC | CDR, USPHS | FDA**
Past Fellow, 2005 - 2007 Cycle
Sam.Skariah@fda.hhs.gov

Sam Skariah graduated from the University of Illinois at Chicago College of Pharmacy where he received his PharmD. He then went on to complete this same track of the regulatory fellowship that was in conjunction with Purdue University, Eli Lilly and Company, and the FDA. He is a Commander in the United States Public Health Service and has served as Team Leader within FDA's Office of Prescription Drug Promotion (OPDP) since 2013. Prior to that role, he served as an FDA-OPDP reviewer for various therapeutic areas.

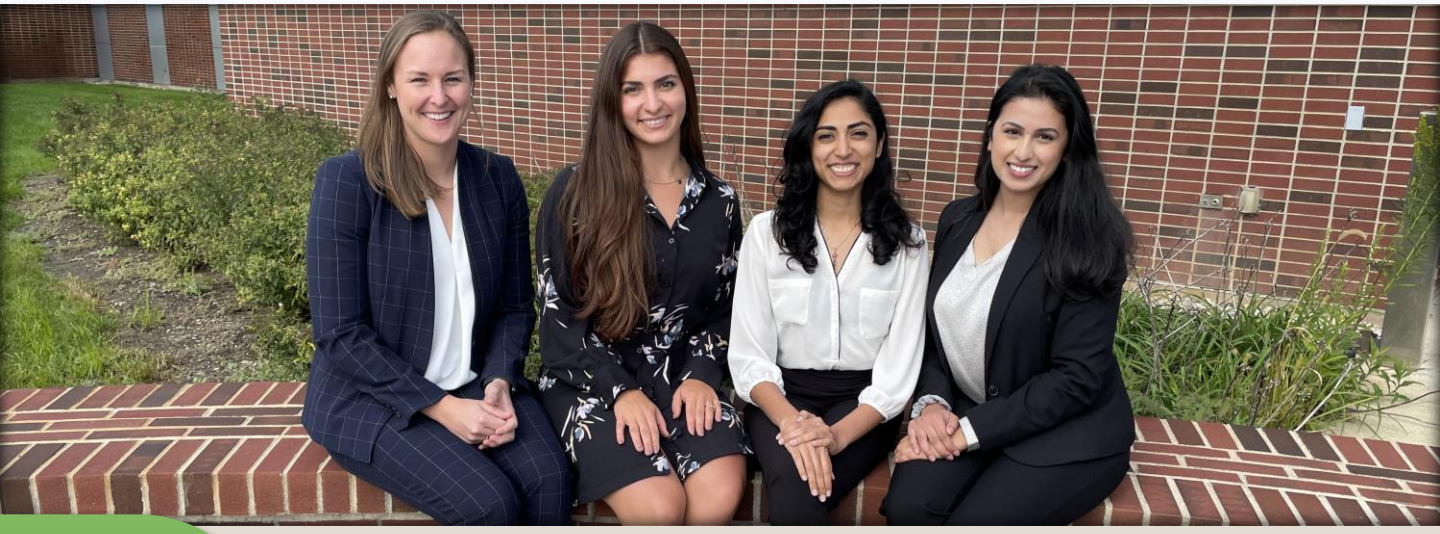
**Jennifer Chen, PharmD, MBA | FDA**
Past Fellow, 2019 - 2021 Cycle
Jennifer.Chen@fda.hhs.gov

Jennifer Chen earned her PharmD from Purdue University College of Pharmacy. She earned an MBA while completing the two-year Regulatory Pharmaceutical Fellowship in the Advertising and Promotion track with rotations at Eli Lilly, FDA, and Purdue University. Following the fellowship, she accepted a position with the Office of Prescription Drug Promotion (OPDP) at FDA. In her current role, Jennifer serves as a reviewer of promotional materials for hematology and oncology products. She also enjoys precepting pharmacy students and serving as a member of a Diversity, Equity, and Inclusion Committee at FDA.

**Kyle Hultgren, PharmD | Purdue University**
khultgre@purdue.edu

Kyle Hultgren is the director of the Purdue University College of Pharmacy's Center for Medication Safety Advancement. Dr. Hultgren's current research includes extensive work on dashboards and measurement systems for the evaluation and improvement of medication use systems as well as large adverse event data set analysis. He holds multiple copyrights on mobile computer simulation technology and two patents on medical devices designed to improve patient safety. Dr. Hultgren lectures nationally and internationally on safe medication use practices and teaches regularly in the Doctor of Pharmacy curriculum where he was the 2016 Dr. Aziz Teaching Award recipient.

# FELLOWSHIP ALUMNI SPOTLIGHT



**Delaney Strong, PharmD**
Medical Information Manager, Vaccines
GSK
**2021-2023 Drug Information Fellow**

"The Purdue University Industry-Affiliated Drug Information Fellowship provided a learning environment where I was able to develop relevant skills while making valuable contributions on each rotation. The experiences, resources, and supportive preceptors in this program successfully prepared me for a career as a drug information pharmacist in an academic, regulatory, or pharmaceutical industry setting."

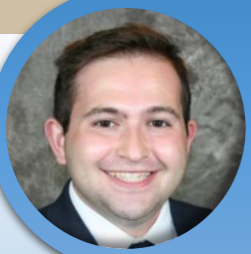
**Divya Desai, PharmD**
Global Medical Information Manager, Diabetes
Eli Lilly and Company
**2021-2023 Drug Information Fellow**

"Purdue's Drug Information Fellowship program was a great fit for me! The program allowed me to pursue my passions for literature evaluation and education, and simultaneously provided me with professional guidance and opportunities to grow the skills I needed for a fulfilling career in medical information."

# FELLOWSHIP ALUMNI SPOTLIGHT


**Amy Bao, PharmD, MPH**
Safety Evaluator, Division of Medication Error Prevention and Analysis
U.S. Food and Drug Administration
**2021-2023 Medication Safety Fellow**

"The Purdue fellowship program in medication safety offers a unique opportunity to learn about applications of medication safety outside of the standard practices of adverse event monitoring and signal detection. During the fellowship, I was able to further cultivate and utilize my strengths to improve patient safety on a global scale through academic, industrial, and regulatory avenues."



**Peter Kokkinias, PharmD, MBA**
Safety Deliverable Manager
AbbVie
**2021-2023 Medication Safety Fellow**

"This program offered me amazing opportunities to network with senior management and hone skills to be a well-rounded pharmacovigilance specialist."

# DRUG INFORMATION

**2021-2023: Divya Desai, PharmD**
Global Medical Information Manager, Diabetes
Eli Lilly and Company

**2021-2023: Delaney Strong, PharmD**
Medical Information Manager, Vaccines
GSK

**2020-2022: Anes Karic, PharmD, MBA, MS**
Medical Science Liaison, Lung Oncology
Janssen Biotech, Inc.

**2020-2022: Shannon Alexander, PharmD**
Manager, Medical Information
Sage Therapeutics

**2020-2022: Katie Comanici, PharmD, MPH**
Associate Director, US Value, Evidence, Outcomes (VEO) Capabilities
Eli Lilly and Company

**2019-2021: Minh Tran, PharmD, MBA**
Medical Science Liaison
AbbVie

**2019-2021: Dylan Vo, PharmD, MBA**
Manager, Global Medical Information
Eli Lilly and Company

**2018-2020: Megan (Cuomo) Walter, PharmD**
Pharmacist, Division of Drug Information
U.S. Food and Drug Administration

**2018-2020: Kaitlin Montagano, PharmD**
Senior Manager, Medical Affairs, U.S.
Astellas Pharma Global Development, Inc.

**2017-2019: Kiersten Rybakov, PharmD**
Manager, Global Medical Information
Eli Lilly and Company

**2017-2019 Jacqueline Wasynczuk, PharmD**
Assistant Professor, Department of Pharmacy Practice
Thomas Jefferson University

**2016-2018: Sandra Bai, PharmD, BCPS**
Labeling Reviewer, Office of Generic Drugs/Office of Regulatory Operations/Division of Labeling Review
U.S. Food and Drug Administration

**2015-2016: Megan N. Freeland, PharmD**
Founder & Health Content Strategist, StockRose Creative, LLC
Founder, Health Professionals to Health Writers Accelerator
Director of Health Communications, Planned Parenthood Federation of America

**2014-2016: Jay R. Fajiculay, PharmD**
Senior Regulatory Health Project Manager, Gastroenterology
Division of Regulatory Operations for Immunology and Inflammation
U.S. Food and Drug Administration

**2013-2015: Bhavani Parikh, PharmD**
US Medical Lead
AstraZeneca

**2012-2014: Andrea M. TenBarge, PharmD**
Sr. Director, Global Medical Affairs Strategy and Transformation
Global Medical Affairs Office
Eli Lilly and Company

**2011-2013: Genevieve Lynn (Ness) Engle, PharmD, BCMAS**
Director, Christy Houston Foundation Drug Information Center
Assistant Professor of Pharmaceutical, Social and Administrative Sciences, Belmont University College of Pharmacy and Health Sciences

**2010-2012: Kimberly (Wu) Chiu, PharmD**
Project Manager
Division of Applied Regulatory Science
Office of Clinical Pharmacology
Office of Translational Sciences
U.S. Food and Drug Administration

**2009-2011: Lindsay E. Wagner, PharmD, MA, BCPS**
Commander, U.S. Public Health Service
Branch Chief, Education & Outreach Branch
Division of Drug Information
U.S. Food and Drug Administration

**2007-2009: Jean Cunningham, PharmD**
Sr. Director, Medical Information
TruLite Health

**2005-2007: Sanjeev K. Bhanot, PharmD**
Sr. Director, Medical Affairs
Merz Aesthetics Inc.

**2003-2005: Tanya Nelson, PharmD**
Principal Scientific Account Lead, East US
Janssen Scientific Affairs, LLC

**2001-2003: John Ng, PharmD**
Consumer Safety Officer, Division of Clinical Compliance Evaluation, Office of Scientific Investigations
U.S. Food and Drug Administration

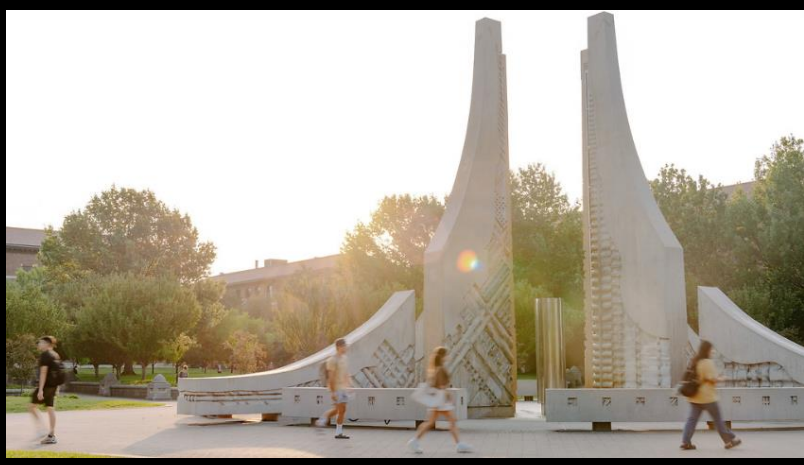
# MEDICATION SAFETY

**2021-2023: Peter Kokkinias, PharmD, MBA**
Safety Deliverable Manager
AbbVie
**2021-2023: Amy Bao, PharmD, MPH**
Safety Evaluator, Division of Medication Error Prevention and Analysis
U.S. Food and Drug Administration
**2020-2022: Morgan Nichols, PharmD**
Manager, Clinical Trial Project Management
Eli Lilly and Company
**2020-2022: Irene Lin, PharmD, MPH**
Senior Safety Data Scientist
AbbVie
**2019-2021: Charlotte Moureaud, PharmD**
Associate Director, Safety Data Sciences
AbbVie
**2019-2021: Jonell Nwabueze, PharmD, MBA**
Leadership Development Program
Eli Lilly and Company
**2018-2020: Danya Faruqi, PharmD**
Associate Director, Safety Data Sciences
AbbVie
**2017-2019: Kathryn Marwitz, PharmD, MPH**
Assistant Professor of Pharmaceutical Sciences
(Social and Administrative Sciences)
Manchester University College of Pharmacy, Natural & Health Sciences
**2016-2018: Kaitlyn Dana Clari, PharmD, PMP**
Project Planner, Associate Director
Pfizer, Groton, CT
**2015-2016: Janelle R. Lenz, PharmD**
Field Scientific Strategy Senior Director - Oncology
Loxo@Lilly
**2014-2015: Trang Truong**
Director, Safety Science Strategic Planning & Operations
BeiGene
**2013-2015: Katelyn Brown, PharmD**
Senior Director, Pharmaceutical Project Manager
Eli Lilly and Company
**2012-2013: Jaclyn A. Jeffries, PharmD, CPh, CPPS**
Executive Director, Safety
AdventHealth

# ADVERTISING AND PROMOTION

**2021-2023: Alison Kwak, PharmD**
Manager, Regulatory Affairs Advertising & Promotion
Sanofi
**2019-2021: Jennifer Chen, PharmD, MBA**
Regulatory Review Officer, Office of Prescription Drug Promotion
U.S. Food and Drug Administration
**2017-2019: Nikki Pedersen, PharmD**
Senior Manager, Regulatory Affairs Advertising & Promotion
Emergent Biosolutions
**2015-2017: John Riehl, PharmD**
Director, Allergan Public Relations and Influencer Consultant
Allergan Aesthetics, an AbbVie Company
**2013-2015: Sam Davis, PharmD**
Principal Consultant, Advertising & Promotion
Opus Regulatory
**2011-2013: Ankur Kalola, PharmD**
Regulatory Review Officer
Office of Prescription Drug Promotion
U.S. Food and Drug Administration
**2009-2011: Nital Patel, PharmD, MBA**
Retiree
**2007-2009: Sheetal Patel, PharmD**
Head, Compliance & Regulatory Advertising and Promotion
Johnson & Johnson Health Care Compliance
**2005-2007: Samuel Skariah, PharmD, RAC**
Commander, U.S. Public Health Service
Team Leader, Office of Prescription Drug Promotion
U.S. Food and Drug Administration
**2003-2005: Amit Patel, PharmD**
Executive Director, Regulatory Affairs Advertising, Promotion & Labeling
Acadia Pharmaceuticals Inc.


PURDUE
UNIVERSITY
College of Pharmacy
Interview
Techniques
Sarah Tanios, Pharm.D.
Michelle T. Le, Pharm.D.
2nd Year Regulatory
Addiction Safety Fellow
AbbVie/FDA



# REGULATORY PHARMACEUTICAL FELLOWSHIPS

Government
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2024-2026

To reach our current fellows and ask questions, email:
Drug Information Track:
DrugInfoFellowship@purdue.edu
Medication Safety Track:
MedSafetyFellowship@purdue.edu
Regulatory Advertising and Promotion Track:
AdPromoFellowship@purdue.edu

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