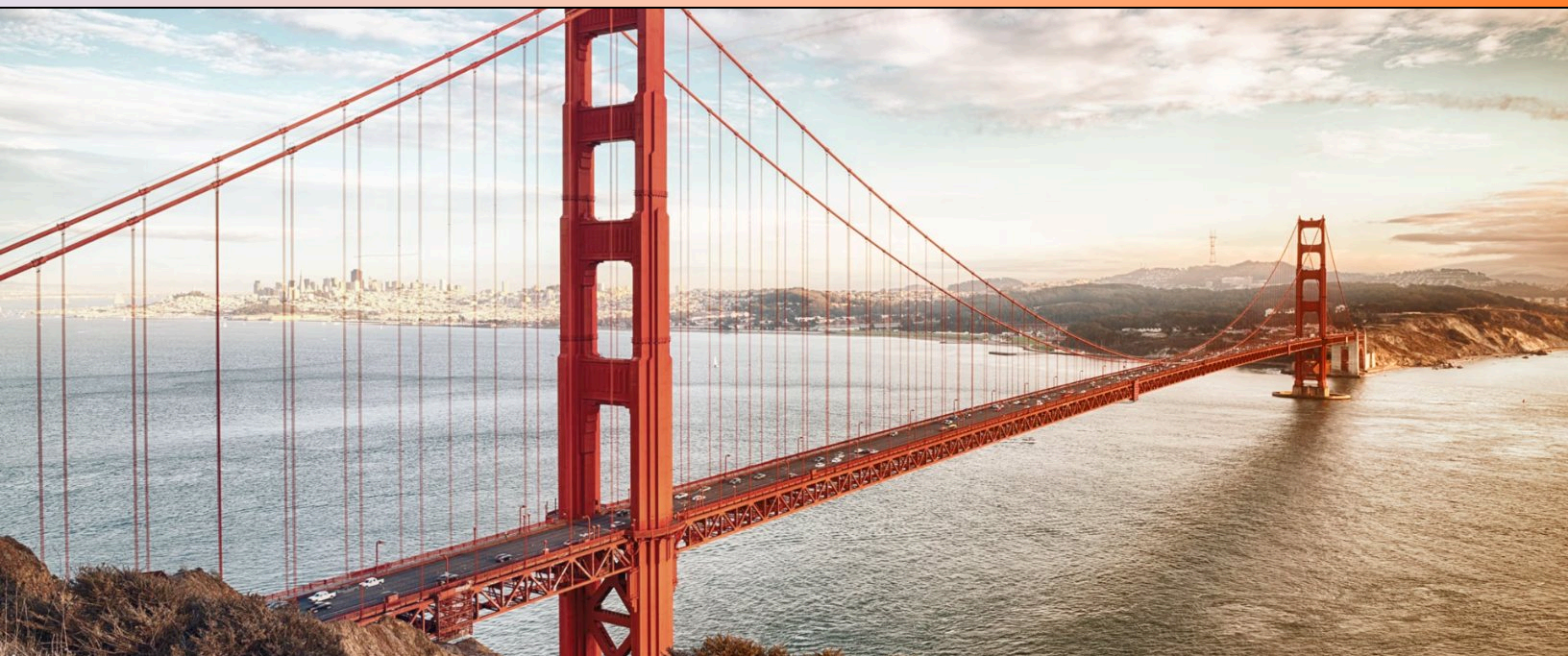


BridgeBio PharmD Fellowship Program

A two-year Regulatory Affairs PharmD fellowship
based in San Francisco, California

2027-2029



OUR MISSION

To find, develop, and deliver breakthrough medicines for genetic diseases to patients as quickly and safely as possible.

ABOUT BRIDGEBIO

Impatient for Patients

BridgeBio Pharma Inc. (BridgeBio) is a biopharmaceutical company with the mission to discover, create, test and deliver transformative medicines to treat patients who suffer from genetic diseases. BridgeBio's pipeline of development programs ranges from early science to advanced clinical trials. BridgeBio was founded in 2015 and its team of experienced drug discoverers, developers, and innovators are committed to applying advances in genetic medicine to help patients as quickly as possible.

BridgeBio exists to bring meaningful medicines to patients as quickly and as safely as possible. We're passionate about fostering open and transparent relationships with patients and their families by staying connected, listening to their experiences, and applying their insights into our work.

[SEE THE FULL BRIDGEBIO PIPELINE →](#)

2000+

patients are enrolled in our clinical trials globally

100%

of our pipeline contains first-in-class therapies



RECOGNITION

#1 Best Place to Work – BioSpace 2026, small biotech companies



THE BRIDGE WAY

Moving at the speed of patient need.

Our decentralized model rethinks what's possible for patients with genetic conditions. We call this The Bridge Way, and it's how we accelerate discovery, reduce the risk of failure, and deliver results.

HUB AND SPOKE Our hub-and-spoke model is designed to unlock speed, precision, and scalability. Teams operate autonomously as individual 'spokes', focused on a specific condition or drug candidate, while our central 'hub' provides clinical, regulatory, and commercial capabilities & expertise.

OUR VALUES

01 Put patients first

We exist to help people living with genetic conditions. To do that, we listen to and work alongside communities to navigate challenges and co-create solutions, whether that means developing new treatments, shaping thoughtful trial designs, or providing other meaningful forms of support. Putting patients first means that compassion and understanding drive every step of our work. Always.

02 Think Independently

Experimentation is expected. We ask "Why?" or "Why not?", challenging convention and approaching every problem with a rigorous, first-principles mindset.

03 Speak up

Great ideas come from a broad range of people (not just leaders) and should be worthy of consideration and testing. We encourage everyone to share their thoughts candidly so we get to the best possible answer.

04 Every Minute Counts

We work with urgency, giving autonomy to focused teams who can make informed decisions faster. Because the quicker we act, the sooner people are connected with meaningful treatments.

05 Let Science Speak

We're driven by data and inspired by discovery. Our model is designed to give every potential medicine a fair chance to prove itself guided by clear criteria, grounded in science, and led by trust in data.

Our growing fellowship program

In partnership with Northeastern University, BridgeBio is offering a two-year Regulatory Affairs PharmD fellowship. The program consists of a 2-year assignment within the Regulatory Affairs team. The PharmD Fellow will work both remotely and on-site at the BridgeBio office location in San Francisco, CA.

LENGTH	Two years, 2027–2029
CORE	Regulatory Affairs
BASED	San Francisco, CA — hybrid



Monica B. Sandberg

Vice President, Regulatory Affairs, Eidos Therapeutics · Fellowship Program Director

We at BridgeBio are excited to be able to collaborate with Northeastern University for our 6th fellowship cohort cycle. Our growing fellowship program seeks to provide Doctor of Pharmacy (PharmD) graduates with hands-on and well-rounded academic and industry experiences that will help prepare them for a successful career in the pharmaceutical industry. Upon completion of this 2-year fellowship, BridgeBio fellows will be equipped with the technical skills necessary to navigate the unique and ever-changing landscape of Regulatory Affairs in a biopharmaceutical company. Additionally, fellows will have the opportunity to work in cross-functional teams and expand their professional network.



Elisa Tsuji, PharmD

Associate Director, Regulatory Affairs, Calcilytix · Fellowship Preceptor

"The BridgeBio Regulatory Affairs Fellowship was a transformative experience that not only deepened my understanding of global regulatory frameworks but also gave me direct involvement in high-impact projects. The hands-on training and close mentorship allowed me to develop critical skills and insights that are essential for navigating the complexities of regulatory affairs."



David Leung, PharmD, MS

Second Year Regulatory Affairs Fellow

"BridgeBio employs a unique operational model that maximizes its potential to transform areas of deep unmet need into approved therapies. The Regulatory Affairs Fellowship has provided me with a platform—supported by strong mentorship and cross-functional collaboration—to holistically develop the technical skills and professional experience needed to support all facets of the regulatory affairs landscape."



Amy Nguyen, PharmD

First Year Regulatory Affairs Fellow

"What sets the BridgeBio Regulatory Affairs fellowship apart is the pairing of dedicated mentorship with genuine autonomy to pursue the areas that interest you most. I was trusted with meaningful projects early on, and that breadth of responsibility across multiple functional areas has built the technical foundation this field demands."

Objectives

Regulatory strategy

- Manage, develop, and implement regulatory strategy in support of development programs.
- Develop proficiency within the Veeva regulatory management system where you will support the planning, preparation and execution of high-quality regulatory submissions (e.g. Clinical Trial Application (CTA)/Investigational New Drug (IND) application and amendments, annual reports, initial license applications (New Drug Application (NDA)/Marketing Authorization Application (MAA), etc.) and supplements/variations).
- Develop and maintain knowledge of International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use (ICH), Good Clinical Practice (GCP), Good Manufacturing Practice (GMP), and other relevant global guidelines to ensure compliance of regulatory strategies and submissions.
- Work in cross-functional teams including Clinical, Statistics, Medical Affairs, Commercial, and Nonclinical groups to help operationalize regulatory strategy.
- Support regulatory interactions with regulatory bodies including preparation of meeting requests and briefing documents.

Global labeling

- Gain in-depth knowledge of labeling regulations regarding Company Core Data Sheet (CCDS), US Package Insert (USPI), Summary of Product Characteristics (SmPC), and Structured Product Labeling (SPL).
- Support development of regulatory labeling strategy and the negotiation and maintenance of competitive labeling with global health authorities.
- Maintain core labeling documents.
- Provide strategic guidance on labeling regulations, competitor labeling, and labeling trends and work with cross-functional team to ensure that the regulatory labeling strategy is aligned with the global regulatory strategy.
- Assist in preparation of responses to labeling-related queries from health authorities.

Advertising and promotion (marketing, legal, regulatory)

- Support regulatory review of promotional and non-promotional materials in accordance with business goals and objectives, health authority regulations, ICH guidelines, Pharmaceutical Research & Manufacturers of Americans (PhRMA) guidelines, company policies and established precedents and recommend revisions/actions that achieve fair balance.
- Support the review and approval of promotional and non-promotional materials in a cross-functional promotional review committee that includes Commercial, Medical Affairs, and Legal.
- Coordinate timely and accurate review of materials for submission to the United States Food and Drug Administration (FDA).



Northeastern University Pharmaceutical Industry Fellowships

ABOUT OUR PROGRAM

OVERVIEW

Northeastern University Pharmaceutical Industry Fellowships Program is a two-year experiential program designed to advance lifelong learning and the education and training of PharmD graduates. Critical to the success of the program is our ability to prepare fellows to meet the ongoing workforce needs in various areas of industry.

Our program provides fellows an opportunity to work with our innovative biopharmaceutical industry partners while collaborating with Northeastern University faculty in the areas of professional and career development, service, scholarship, and teaching.



OUR MISSION

Our mission is to provide the highest quality training for future biopharmaceutical industry professionals by combining industry expertise with Northeastern University's renowned tradition of lifelong and experiential learning.

CORE VALUES

- ☑ Innovation
- ☑ Lifelong Learning
- ☑ Social Impact Through Drug Development
- ☑ Quality Industry Pharmacy Training

FOCUS AREAS

LEARN

Pursue graduate degrees or certificates in Regulatory Affairs, Business, Public Health, and more through tuition reimbursement.

TEACH

Teach pharmacy students in various small and large group classes. Earn a Teaching Certificate of achievement.

RESEARCH

Perform research with faculty and students. Present data at conferences. Publish your findings. Generate literature.

NETWORK

Boston has a lot to offer, both socially and professionally. Our program takes advantage of it all!

Thanks to the collaboration and dedication of our industry partners over the course of nearly 10 years, our program is now the 3rd largest industry fellowship program in the nation.

OUR PROGRAM PILLARS

TEACHING & SCHOLARSHIP

Teaching and Learning Seminar Series provides contextual activities and reflection on adult education and pedagogy outcomes.

PROFESSIONAL & CAREER TRAINING

Professional Development and Career Training Series are customized to engage fellows in appropriate and professional conduct for success.

SERVICE

Through their service on committees, fellows have an active connection within the community and program. This allows an opportunity to demonstrate leadership development and skills.



Northeastern University Pharmaceutical Industry Fellowships Program provides a dynamic academic environment offering fellows the opportunity for a wide breadth of experiences.



PROGRAM OPPORTUNITIES

Develop teaching skills through participation in our Teaching and Learning Seminar Series

Utilize a layered learning model in experiential education by co-precepting students on pharmacy practice experiences including Northeastern's unique co-op program

Facilitate small and large group didactic education in partnership with a faculty mentor

Create, present, and publish scholarly research through collaborative industry and university relationships

Engage with faculty who participate in various interdisciplinary graduate programs including biotechnology, nanomedicine, immunology, health informatics, and drug discovery

Network with local residents and other Northeastern fellows via professional development programs, teaching seminars, and participation on fellowship committees

OUR PARTNER COMPANIES

SINCE 2015

Through these exciting partnerships, Northeastern fellows collaborate and learn from each other, further positioning them to be successful in both academic and industry settings. Fellows are empowered to shape their experience, as well as the future of the program, through leadership on the Professional Development & Networking and Recruitment committees.



26

Industry Partnerships

74

Active Fellows

156

Fellowship Alumni

99%

Employment in Industry within 90 days of Fellowship Completion



OUR TEAM



Andrew Orr-Skirvin

Faculty Director



Amanda Scott

Faculty Manager



Sherisse Mayala-Nelson

Program Manager



Sophia Sawtelle

Senior Program Coordinator



Julia Van

Director of Corporate
and Foundation Relations



Milini Rambukwella

Human Resources
Associate



Dayna D'Angelo

Budget Coordinator



Jason Lancaster

Clinical Professor
Fellowship Faculty Manager



Adam Wooley

Clinical Professor
Fellowship Faculty Manager

**CONNECT
WITH US!**



Instagram

[@nufellowship](https://www.instagram.com/nufellowship)



LinkedIn

[Northeastern Pharmaceutical
Industry Fellowships](#)

BOSTON, MA:

Northeastern University is in the heart of Boston, steps away from the famous Fenway Park and downtown Boston. Surrounded by top tier academic and research institutions, the area is home to pioneers of innovation who are committed to intellectual curiosity and scientific advancement. The greater Boston area is a cultural hub, offering world-class attractions and rich history.

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YEAR 1 FELLOWS



Shyanne Casapao PharmD
Abbott
Global Medical Affairs



Leah Song, PharmD
Alnylam
Global Scientific Communications



Azhar Hussain, PharmD, DHA, MBA
Abbott
Global Medical Affairs



Kezia Eapen, PharmD
Alnylam
Global Medical Information



Tania Nguyen, PharmD
Alnylam
US Medical Affairs



Sarika Satishkumar, PharmD
Alnylam
Global Regulatory Affairs



Shadi Barjasteh Mohebby, PharmD
Alnylam
Global Patient Safety
and Risk Management



Zeenat Entezar, PharmD
Alnylam
Global Medical Affairs



Alec Bayne O'Reilly, PharmD
Alnylam
Early Assets Global Medical
Affairs



Wendy Wang, PharmD
Alnylam
Clinical Science



Kate VanBilliard, PharmD
Alnylam
Global Scientific
Communications



Rohan Prasanna Deshpande, PharmD
Alnylam
U.S. Marketing - Rare Disease TTR

YEAR 1 FELLOWS



Hemali Patel, PharmD
Apellis
Medical Affairs



Aiyat Ali, PharmD
Dompe
Global Regulatory Affairs



Christopher Cha, PharmD
BioNTech
Health Economics and
Outcomes Research



Victoria Walsh, PharmD
Intellia Therapeutics
Global Medical Affairs



Amy Nguyen, PharmD
BridgeBio/ Eidos
Regulatory Affairs



Hania Mohsen, PharmD
Ironwood
Regulatory Affairs



Veerkumar Patel, PharmD
Chiesi
North America Medical Affairs



Apostoli Malouhos, PharmD
Pacira
Global Medical Information and
Communications



Cassie Lynch, PharmD
CSL Seqirus
Medical Affairs



Caitlin Raimo, PharmD
Scholar Rock
Global Scientific Communications



Jan Hernandez-Velazquez, PharmD, MS
Dompe
Global Regulatory Affairs



Llaran Turner, PhD
Takeda
Global Medical Affairs

YEAR 1 FELLOWS



Peeper Walker, PharmD
Takeda
Global Medical Affairs



Brinn Mancuso, PharmD
Vertex
Medical Affairs



Tina Zheng, PharmD
Takeda
Global Regulatory Affairs



Xhulio Kampula, PharmD
Vertex
Medical Affairs



Alanna M. Farrell Colón, PhD
Takeda
Global Medical Affairs



Abena Agyei, PharmD, MS
Vertex
Regulatory Affairs



Kathryn LeMoine, PharmD, MA
Vertex
Clinical Science



Alexander McHugh, PharmD
Zevra
Global Medical Affairs



Sophie Friedman, PharmD
Vertex
US Market Access



Ellie King, PharmD
Vertex
Medical Affairs

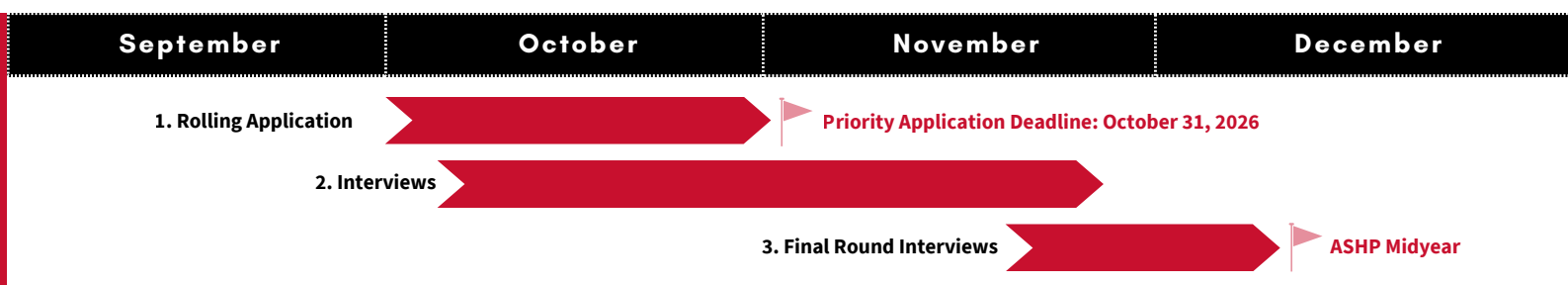


Northeastern University

Pharmaceutical Industry Fellowships

APPLICATION REQUIREMENTS

Fellows are selected on a nationally competitive basis. Unless otherwise noted in the position description, candidates must have a Doctor of Pharmacy degree from an ACPE-accredited college of pharmacy by June 30, 2027. Candidates must apply through [Northeastern's career portal](#) and are encouraged to do so by the priority application deadline of October 31, 2026.



Step 1: Application	Step 2: Interviews	Step 3: Final Round Interviews & Offers
- Fellowship position postings go live on October 1, 2026. We will cross-post positions on ASHP's PPS website. - Complete the application by October 31, 2026, for priority consideration for first-round interviews.	- Phone screenings and initial interviews will be conducted with eligible candidates starting in early October into November. - After the priority deadline, interviews will be conducted with both the sponsor company and Northeastern University.	- Final round interviews will take place virtually or in person during ASHP Midyear in Orlando, FL. Visit midyear.ashp.org for event details. - Northeastern University, in conjunction with the Alliance of Industry Fellowship Associates (AIFA), has agreed to extend offers for fellowships no earlier than December 11th, 2026.

Applications are reviewed on a rolling basis – apply early!

Application Materials:

- Curriculum Vitae (CV)
- Unofficial PharmD Transcript
- Cover Letter

3 Letters of Recommendation:

- Highly encouraged to submit by **October 31, 2026**
- Official Deadline: **November 23, 2026**
- Recommenders must submit LORs using the [link here](#) or follow the instructions on the job posting
- Letter writers should submit **one letter per candidate and follow the instructions on the form carefully**

ADDRESS YOUR COVER LETTER AND 3 LETTERS OF RECOMMENDATION TO:

J. Andrew Orr-Skirvin, PharmD, BCOP
 Clinical Professor, School of Pharmacy
 Chair, Department of Pharmacy & Health System Sciences
 Director of Pharmaceutical Industry Fellowship Program
 360 Huntington Ave, 140TF R218
 Boston, MA 02115

For more information and fellowship resources:

Visit: a27p.com
 Visit: bouve.northeastern.edu/pharmacy/fellowships
 Email: PharmDfellowships@northeastern.edu

Please note: Northeastern University will not provide H-1B, TN, O-1, E-3, or any other type of employment visa sponsorship for the successful applicant to this position, now or in the future. STEM OPT is not compatible for this position. The successful applicant must be able to maintain valid work authorization in the United States throughout the entire appointment without Northeastern University's sponsorship for a visa.