

CERSI IMMERSION PROGRAM

APRIL 2-20, 2018

STANFORD UNIVERSITY

By enrolling in this program, participants will gain a comprehensive and rigorous overview of regulatory and scientific considerations for the medical product development. Intensive one-day courses are taught by industry subject matter experts in the Spring quarter.

A Statement of Completion is offered to participants who complete all coursework. Limited enrollment.

Intended Audience:

Graduate and professional students, postdocs, academic researchers, life science and early career regulatory professionals

Format:

Face-to-face lectures, discussions and group work

Roadmap:

- April 2 Introduction to U.S. Food and Drug Administration
- April 4 Medical Device Development and Regulation
- April 6 New Drug Development and Regulation
- April 9 Large Molecule Development
- April 11 Non-clinical Safety Assessment
- April 13 Early Clinical Drug Development
- April 16 510(k) Regulatory Pathways for Medical Devices
- April 17 Challenges in the regulation of high cost treatments -
An overview from Latin America
- April 18 Biomarkers in Drug Development and Approval
- April 20 Post Marketing Considerations for Drugs

For more information, visit: cersi.stanford.edu/cersi-immersion-program

