

Pan-Canadian Research Connect Presents:
Navigating FDA Regulations for Canadian
Clinical Trials

LEARNING OBJECTIVE:

- Understanding the FDA Regulatory Framework
- Differentiating FDA Requirements from Canadian Regulations
- Navigating Cross-Border Regulatory Compliance
- Data Quality and Integrity
- Managing Regulatory Inspections and Audits
- Engaging with Regulatory Agencies
- Identifying Resources and Support Networks



Eric S. Pittman

Program Division Director, Bioresearch Monitoring West (BIMO-W), Office of Regulatory Affairs' (ORA), Office of Bioresearch Monitoring Operations, Food and Drug Administration (FDA).









