Regulatory And Quality Considerations In Molecular Pharmaceutics

8:45 AM - 10:15 AM: Session 1: Regulatory Guidelines for Molecular Pharmaceutics

Discussion on regulatory guidelines.
Protocols for ensuring compliance with FDA, EMA, and other regulatory bodies in the development of molecular pharmaceuticals.

10:15 AM - 10:30 AM: Coffee / Tea / Snacks Break

Networking and refreshments.

10:30 AM - 12:00 PM: Session 2: Quality by Design (QbD) in Molecular Pharmaceutics

Practical session on applying QbD principles.
Protocols for integrating Quality by Design (QbD) approaches to improve drug quality and manufacturing processes.

12:00 PM - 1:00 PM: Lunch Break

Catered lunch and networking opportunity.

1:00 PM - 2:30 PM: Session 3: Stability Testing and Shelf-Life Prediction

Hands-on training on stability testing.
Protocols for predicting shelf-life and ensuring the stability of pharmaceutical compounds over time.

2:30 PM - 2:45 PM: Short Break

Time for a stretch and informal discussions.

2:45 PM - 4:15 PM: Session 4: Risk Management in Drug Development

Practical session on risk management in drug development.
Protocols for identifying and mitigating risks during the molecular drug development process.

4:15 PM - 4:30 PM: Coffee / Tea / Snacks Break

Last networking opportunity with snacks.

4:30 PM - 5:30 PM: Closing Session: Implementing Changes and Technology Adoption

Group discussions on implementing new techniques learned today.
Dialogue on overcoming challenges in adopting new technologies in similar sectors.
Feedback session and closing remarks.
Certificate Issue

5:30 PM: Workshop Concludes