From Molecule to Market: R&D and Regulatory Considerations

Description

From Molecule to Market: R&D And Regulatory Considerations builds on the business overview of the *Intro to Business* course. This course delivers an outline of clinical, regulatory, and quality considerations and practices. Attendees will be provided foundational information about the Biopharma industry and regulations via instruction, storytelling, and breakout group discussions. The course will explore the decision-making process by touching on relevant topics such as market share and phases of development.

Format

Online: Course will occur over three days, 2.5 hours per day.

Topical Agenda

Session 1:

- The History of Regulations
- Industry Overview
- Research & Development: An Overview of R&D

Session 2:

- Roles in Clinical Trials
- Marketing Considerations

Session 3:

- Compliance via Regulatory & Quality Process