Due Diligence & Regulatory Information Management Workshop

25 November 2019 @ Glenhove Events Hub

Introduction

This 1-day workshop aims to provide those in Regulatory Affairs and Compliance Departments an overview of the data and GxP requirements related to Health Products.

Who should attend?

- ☑ Responsible Pharmacists
- ☑ Regulatory Information Managers
- ☑ Regulatory Pharmacists
- ☑ QA Managers
- ☑ Product Development Managers
- ☑ Compliance Managers
- ☑ Regulatory Operations Managers

Course Outcomes

At the end of the course, the delegate will have insight and knowledge of the following:

- Legislation related to Health Products
- Classification methodology for Health Products
- Data requirements for submission purposes
- Cohesive compliance information management – Business, RA, QA & PV
- Regulatory Information Management requirements over Products' Lifecycles

Presenter

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