PVU

PharmaVigilant University

Course Catalog

PharmaVigilant
adminedu@pharmavigilant.com

Course Catalog: PharmaVigilant Introduction and Overview

Course Name	Description	Audience	Duration
PharmaVigilant Introduction and Overview for Clients	Provides background on the Company, roles, overview of services we will provide, and roles in the clinical trial process	All new clients	2 hours

Course Catalog: InSpire

Course Name	Description	Audience	Duration
InSpire for CRCs	Use the InSpire system to enter data and respond to queries during a clinical trial.	Clinical Trial coordinators	8 hours
InSpire for Sponsors	Use the InSpire system to audit the progress of a trial.	Clinical Trial Sponsors who audit the study data.	8 hours
InSpire for Investigators	Use the InSpire system to enter data, respond to queries, and enter electronic signatures.	Clinical Trial Investigators	1 hour
Inspire for rSDV Specialists	Use the InSpire system to examine and verify data, respond to discrepancies, and approve queries.	Remote Source Document Verification Specialists	4 hours
InSpire for Data Managers	Use the InSpire system to analyze, verify, clean and manage clinical data.	Data Manager	5 hours

Course Catalog: I-Vault

Course Name	Description	Audience	Duration
I-Vault for CRCs	Use the I-Vault system to scan source worksheets during a clinical trial.	Clinical Trial coordinators at trial sites	8 hours
I-Vault for Sponsors	Use the I-Vault system to audit the progress of the trial.	Clinical Trial Sponsors who audit and query trial data	2 hours
I-Vault for Investigators	Use the I-Vault system to scan source worksheets and enter electronic signatures.	Clinical Trial Investigators	1 hour
I-Vault for Data Managers	Use the I-Vault system to audit and verify data.	Clinical Trial Data Managers	7 hours
I-Vault for rSDV Specialists	Use the I-Vault system to examine and verify data.	Remote Source Document Verification Specialists	4 hours

Course Catalog: rSDV

Course Name	Description	Audience	Duration
rSDV for Sponsors	Understand the rSDV process and steps during a clinical trial.	Clinical Trial sponsors	1 hour
rSDV for Coordinators and Investigators	Understand your responsibilities and the rSDV process from the rSDV specialist's role.	Clinical Trial Data Coordinators and Investigators	1 hours
rSDV for Data Managers (Demonstration)	Understand the rSDV process to audit, verify, and approve data and report data and adverse events.	Clinical Trial Data Managers	1 hours
rSDV for rSDV Monitors (Hands on)	Comprehensive course on how the rSDV process works to examine, verify, query, and approve data.	Remote Source Document Verification Monitors	2 hours