Life & Health

Global Language Courses

34 Languages:

Arabic
Chinese (Simplified)
Chinese (Traditional)
Czech
Danish
Dutch Nederlands
English
Finnish
Flemish
French (Canada)
French (France)
German
Greek
Hebrew
Hindi
Hungarian
Indonesian
Italian
Japanese
Korean
Malaysian (Malay)
Norwegian
Polish
Portuguese (Brazil)
Portuguese (Portugal)
Romanian
Russian
Slovak
Spanish (Spain)
Spanish (Latin America)
Swedish
Thai
Turkish
Vietnamese

Translated Courses

UL EduNeering's learning solutions reflect today's global business environment. To support our global clients, we have translated many of our popular courses into multiple languages as specified in this brochure.

Custom Translations

Any of our 700+ standard or custom courses can be translated into a specific language upon request.

If you have subcontractors, research sites and manufacturing operations around the world, join other global companies that rely on ComplianceWire® as their single enterprise-wide compliance learning platform for users in both Asia and Europe.

Global LMS Interface Options

ComplianceWire, our exclusive online learning management system, provides the option of enabling users to change their user interface into any of the 34 supported languages.

Course Title	CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
DCRI Courses:														
Audits and Inspections: Identifying Fraud and Misconduct	DCRI-04	✓												
China Food and Drug Administration (CFDA): Clinical Trial Regulation in China	DCRI-14	✓												
CREATe Refresher Course	DCRI-13	✓												
Drug Safety and Adverse Event Reporting	DCRI-10	✓												
Electronic Data Capture	DCRI-12	✓												
Evolution of Clinical Research and Drug Safety	DCRI-01	✓												
How is Clinical Research Regulated?	DCRI-03	✓												
Human Research Protection Program	DCRI-07	✓												
Informed Consent – Part 1	DCRI-08	✓												
Informed Consent – Part II	DCRI-09	✓												
Phases of Clinical Research	DCRI-02	✓												
Recruitment, Retention and Lost to Follow-Up	DCRI-11	✓												
Responsibilities of a Clinical Research Coordinator (CRC) in FDA-Regulated Studies	DCRI-05	✓												
Responsibilities of Investigators Conducting FDA-Regulated Studies	DCRI-06	✓												

Course Title	CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
Ethics Courses:														
Detecting and Preventing Fraud	RH0024	✓	✓				✓			✓				✓	
Conflicts of Interest	RH0007	✓	✓				✓			✓				✓	
Foreign Corrupt Practices Act (FCPA)	RH0014	✓	✓			✓	✓	✓					✓	✓	
Global Anti-Bribery	RH0041						✓	✓		✓	✓	✓	✓	✓	✓
Handling Confidential Information	RH0006	✓	✓				✓			✓				✓	
Privacy and Data Protection	RH0027	✓	✓				✓			✓				✓	

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
HR & Environmental Health and Safety Courses:														
Basic Radiation Awareness	EHS05							✓						
Bloodborne Pathogens – Health Care Workers	EHS09							✓						
Confined Space Entry	EHS15													✓
Driver Safety Program (DSP)	EHS110						✓ (EU)							✓
Electrical Safety	EHS23													✓
Ergonomics: Body Mechanics and Fitness	EHS33					✓								
Forklift Safety	EHS34													✓
Hazard Communication	EHS37							✓						✓
Hazardous Waste Determination	EHS38													✓
Heat Stress	EHS44													✓
Hot Work Permits	EHS47													✓
Improving Productivity	EHS49													✓
Introduction to Workplace Safety	EHS53													✓
Lockout/Tagout – Authorized	EHS58													✓
Machine Safeguarding	EHS59													✓
Personal Protective Equipment	EHS67						✓							
Slips, Trips and Falls	EHS98													✓

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
Miscellaneous Courses:														
Welcome to the ComplianceWire® Platform	SYS01					✓								

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
GMP FDA Enforcement Courses:														
FDA Establishment Inspection Report Writing	FDA26	✓												
FDA Good Guidance Practices (GGPs)	FDA21						✓							
MDR Regulation 1: Overview and General Provisions	FDA63		✓											
MDR Regulation 2: Device User Facility, Importer and Reporting Requirements	FDA65		✓											
MDR Regulation 3: Requirements for Individual Adverse Event Reports	FDA66		✓											
Part 11: Electronic Records; Electronic Signatures	FDA31						✓							

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
GMP Medical Device Courses:														
Complaint Management for Medical Device Manufacturers	DEV46	✓	✓				✓	✓					✓	
Design Control Regulations for Medical Device Manufacturers	DEV40	✓	✓				✓	✓					✓	
Effectively Responding to FDA 483s and Warning Letters	PHDV70												✓	
Essentials of an Effective Calibration Program	PHDV75												✓	
Global Regulatory Strategy and Planning Process	DEV54			✓										
Good Documentation Practices for Medical Device Manufacturers	DEV56	✓					✓ (EU)							✓
Handling an FDA Inspection	PHDV74	✓	✓											
Introduction to Quality System Regulation (QSR)	DEV43	✓	✓				✓ (EU)	✓					✓	
Medical Device Packaging, Labeling and Distribution	DEV41						✓	✓						
Meeting GMP Training Requirements	PHDV76						✓							
Orientation to GMP Compliance	PHDV73	✓				✓	✓							
Principles of Good Documentation	PHA74	✓					✓ (CA/EU)						✓	✓
Q9: Quality Risk Management	ISPE09		✓											
Quality Systems Inspection Technique (QSIT)	DEV42	✓					✓							
Understanding GMPs for Facilities and Equipment	PHDV63						✓							

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean	Dutch	French	German	Hebrew	Italian	Polish	Russian	Latin: Portuguese (Brazil)	Latin: Spanish (Europe)	Latin: Latin America
GMP Pharmaceutical Courses:														
Batch Record Reviews	PHA53						✓							
Biotechnology: An Overview of Compliance Considerations	PHDV68												✓	
Change Control	PHA35						✓							
Effectively Responding to FDA 483s and Warning Letters	PHDV70												✓	
Essentials of an Effective Calibration Program	PHDV75												✓	
Failure Investigations for Pharmaceutical Manufacturers	PHA59						✓							
GMP Principles for Batch Records	PHA60					✓								
GxPs	PHDV61						✓							
Handling an FDA Inspection	PHDV74	✓												
Introduction to GMPs	PHA38						✓		✓				✓		✓
Maintenance and Cleaning of Drug Manufacturing Equipment	PHA44						✓							✓	
Meeting GMP Training Requirements	PHDV76						✓							
Orientation to GMP Compliance	PHDV73	✓				✓	✓							
Packaging and Labeling of Finished Pharmaceuticals	PHA39						✓							✓	
Principles of FDA Inspections for Pharmaceutical Manufacturers	PHA61						✓							
Principles of Good Documentation	PHA74	✓					✓ (CA/EU)							✓	✓
Understanding GMPs for Facilities and Equipment	PHDV63						✓							
Writing and Reviewing SOPs	PHA48						✓							

2014 Scheduled Course Translations

All of the courses listed are being translated into Simplified Chinese, Japanese and Korean languages in 2014.

Those marked with a "✓" are complete as of the publication of this brochure.

Course Title	English Language CODE	Asia: Chinese (Mandarin)	Asia: Japanese	Asia: Korean
Global Regulatory:				
Q9: Quality Risk Management – ICH Q9	ISPE09		✓	
A Guide to ISO 13485 – The Quality Management System for Medical Devices	DEV50	✓		
Global Regulatory Strategy and Planning Process	DEV54	✓		✓
EU Medical Device Directive: Part I — Introduction	MDD01		✓	✓
EU Medical Device Directive: Part II – Specific Procedures	MDD02		✓	
The Approval Process for New Medical Devices in the US	DEV47			
Design Control Regulations for Medical Device Manufacturers	DEV40	✓	✓	
ISO 14155: Obligations of Sponsors and Monitors for Medical Device Trials	GCP03			✓
Audits and FDA Inspections:				
Pre- and Post-Approval FDA Inspections	PHDV66			
Principles of Auditing	PHDV69	✓		✓
Handling an FDA Inspection	PHDV74	✓	✓	✓
QSIT 4: The Corrective and Preventive Actions Subsystem	FDA53			
Quality System Inspection Technique	DEV42	✓		✓
Risk Management 1: Key Concepts and Definitions	FDA29			✓
Complaints, Recalls and FDA MDR Regulations:				
Handling a Product Recall	PHDV64	✓	✓	
Failure Investigations for Medical Device Manufacturers	DEV45			
Complaint Management for Medical Device Manufacturers	DEV46	✓	✓	
The MDR Regulation 1: Overview and General Provisions	FDA63	✓	✓	✓
MDR Regulation 2: Device User Facility, Importer and Reporting Requirements	FDA65	✓	✓	✓
MDR Regulation 3: Requirements for Individual Adverse Event Reports	FDA66	✓	✓	✓
FDA Quality System Regulation (QSR):				
Introduction to the Quality System Regulation (QSR)	DEV43			✓
QS Regulation 3: Design Controls	QSR03			
QS Regulation 5: Identification & Traceability; Production and Process Controls	QSR05			
QS Regulation 7: Corrective and Preventive Action	QSR07			✓
QS Regulation 8: Labeling and Package Control; Handling, Storage, Distribution and Installation	QSR08			✓
FDA Import Operations:				
Import Operations 1: Background	FDA37			
Import Operations 2: The Process	FDA42			
Import Operations 3: Other Activities	FDA43			
GMP Key Concepts:				
GMP Principles of SOPs	PHA64			
Key Concepts of Process Validation	PHDV77			
Essentials of an Effective Calibration Program	PHDV75			
FDA Training and Qualification Requirements	PHA67	✓		
Environmental Control and Monitoring	PHDV87			
Risk Management 1: Key Concepts and Definitions	FDA29	✓		

About UL EduNeering

UL EduNeering is a business line within UL Life & Health's Business Unit. UL is a premier global independent safety science company that has championed progress for 120 years. Its more than 10,000 professionals are guided by the UL mission to promote safe working and living environments for all people.

UL EduNeering develops technology-driven solutions to help organizations mitigate risks, improve business performance and establish qualification and training programs through a proprietary, cloud-based platform, ComplianceWire®.

For more than 30 years, UL has served corporate and government customers in the Life Science, Health Care, Energy and Industrial sectors. Our global quality and compliance management approach integrates ComplianceWire, training content and advisory services, enabling clients to align learning strategies with their quality and compliance objectives.

Since 1999, under a unique partnership with the FDA's Office of Regulatory Affairs (ORA), UL has provided the online training, documentation tracking and 21 CFR Part 11-validated platform for ORA-U, the FDA's virtual university. Additionally, UL maintains exclusive partnerships with leading regulatory and industry trade organizations, including AdvaMed, the Drug Information Association, the Personal Care Products Council and the Duke Clinical Research Institute.

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