ePath

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ePath is knowledge based web training. It focuses on terms, definitions, regulations and expectations.

PathWise

WE BUILD

KNOWLEDGE

ePath is the PathWise online learning managment system. Modules can be purchased for individuals, small, or large groups. ePath is the PathWise solution for your on-demand learning needs!

BENEFITS

There are many benefits for your organization from chosing epath web based training.

Fulfill New Hire and FDA Requirements

Ensure employees are receiving consistent training across the board, whether it is for new employee onboarding, or to mainting annual training records.

Be Compiant to Regulations

ePath training covers many important regulations and best practices. Ensure employees have the knowledge for a compliant, effective and efficient quality system.

Flexible Mobile Learning

Use a desktop, laptop, tablet or phone. Our web based training can be accessed anywhere online!

Optimize Training Time

Complete training when and where it works for your employees, and your organization.

COMPLIANCE

Learn about the laws and regulations that govern the regulated life science industry.

PathWise Compliance Web Based Training (WBT) modules teach the laws and regulations that govern the regulated life science industry. This group of knowledge-based online training focuses on terms, definitions, regulations and expectations.

ePath

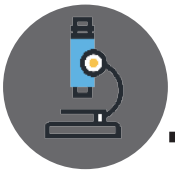
COMPLIANCE TRAINING MODULES

Quality Systems Regulation (QSR)

This course discusses why the QSR is important, and how it supports your business and compliance drivers. The course provides a breakdown of the QSR requirements and includes subpart detail, and highlights the differences between FDA and ISO standards. **AVAILABLE IN SPANISH.**

Current Good Maufacturing Practices (cGMP)

This course focuses on 21 CFR Part 210 & 211 cGMP in Manufacturing, Processing, Packing, or Holding of Drugs and Finished Pharmaceutical. It discusses the role of the FDA in the cGMP process, the requirments from the FDA, and subpart detail.

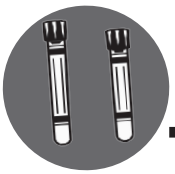


Biologics 21 CFR 600, 601, 610

This course provides attendees with a high level understandign of the FDA requirements for biologics in 21 CFR parts 600, 601, and 610. Major topic areas covered are general requirements, licensing, and biologics standards.

Good Documentation Practices (GDP)

This course helps your organization be consistent in documenting activities to exceed regulatory expectations. The course provides a breakdown of the requirements, and shares best practices and industry standards for GDP. **AVAILABLE IN SPANISH.**



Combination Products

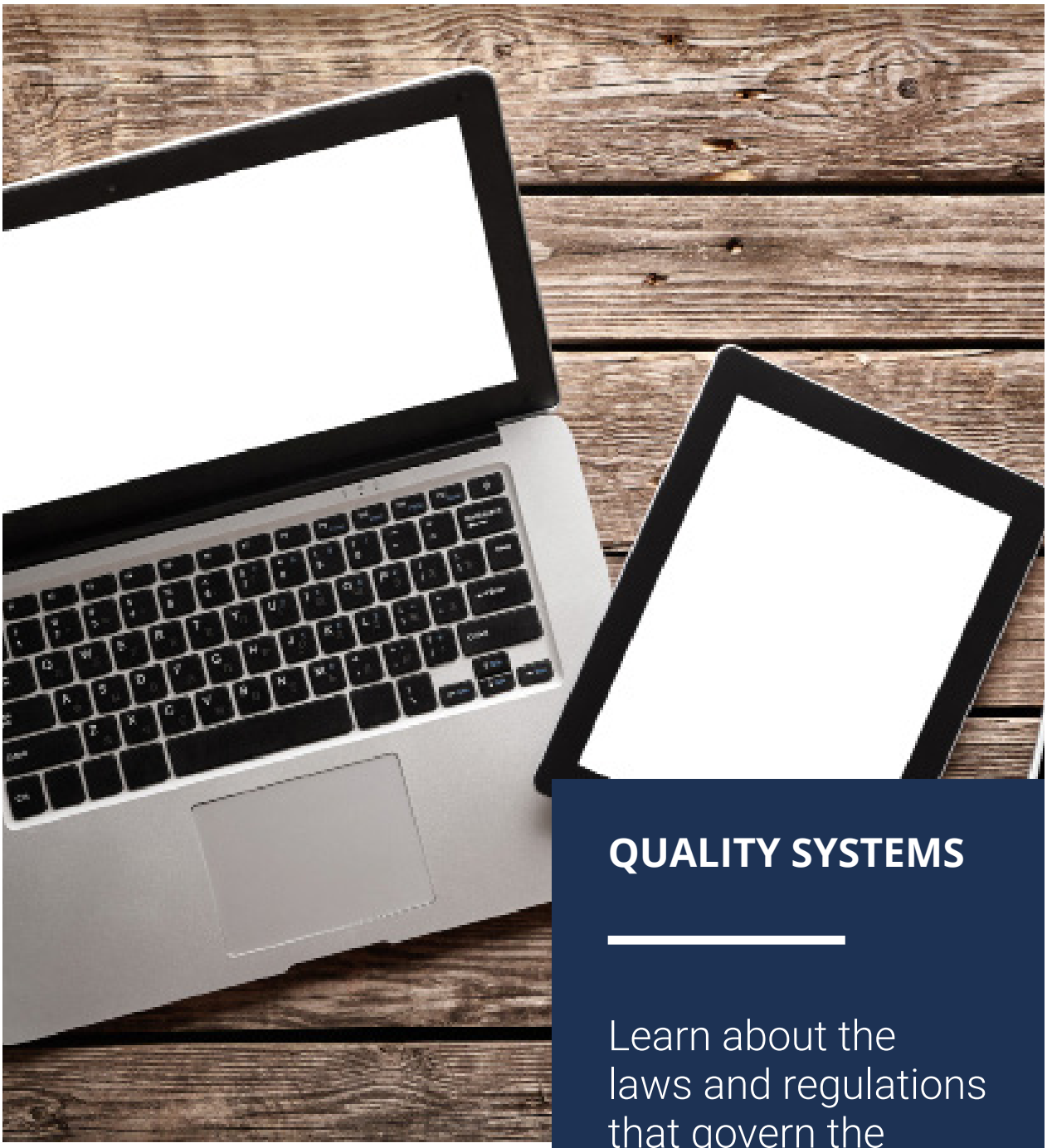
This course provides a clear regulatory understanding of the obligations manufacturers of Combination Products have. It focuses on the Guidance Document issued by the FDA in January 2017 for cGMP requirements for combination products.

Electronic Records and Signatures

This web based training modules gives an overview of electronic records and signatures. It goes through requirements for 21 CFR Part 11, Public Docket 92S-0251, open and closed systems and electronic signatures. The course also discusses the importance in using good documentation practices.

Good Clinical Practices (GCP)

This one hour training will help you understand GCP, including the recent updates to the regulations in E6(R2), ICH E6(R1), how the regulation came to be, and why they are essential for your organization.



QUALITY SYSTEMS

Learn about the laws and regulations that govern the regulated life science industry.

With the PathWise Quality Systems Web Based Training (WBT) Modules you and your team can learn about different pieces of your quality management system, to ensure your product is up to regulations, and meets your customer's needs.

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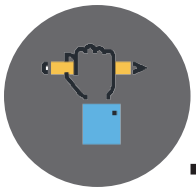
QUALITY SYSTEM TRAINING MODULES

INTRO TO INVESTIGATIONS AND CAPA

The Investigations and CAPA training module discusses a high-level understanding of why the Investigations and CAPA is important, and how it supports your business. It also highlights important terminology , regulations, and requirements.
AVAILABLE IN SPANISH.



INSPECTION READINESS

This three-module course teaches how to prepare, what to expect, the roles and responsibilities, and how to manage an inspection. The modules can be assigned independently to different personnel in your organization based on their involvement with inspections.

INTRO TO QUALITY CHANGE CONTROL

This training module focuses on best practice approaches to support the quality change control system. It outlines the change control process, the related regulations, and provides background for you employees and partners to understand the system's importance.

INTRO TO TECHNICAL WRITING

This course focuses on a high-level understanding of technical writing and why it is important to your business and compliance drivers. Learners will learn the expectations for technical writing, and discover key steps in planning and preparing documents.

INTRO TO HUMAN FACTORS AND PERFORMANCE

This training discusses a basic understanding of Human Performance, introducing Human Factors and Performance into your organization. It provides an overview of the influences in human factors and performance to systems, organizations, and people.

INTRO TO QUALITY RISK MANAGEMENT

This course covers covers Quatliy Risk Management key terms, and what the FDA requires in defining the Quality Risk Management process. It describes the Risk Models used in both Medical Device and Pharmaceutical industries, including available tools best suited for each situation.

PRODUCTIVITY

Learn about how to improve productivity in your organization.

With the PathWise Productivity Web Based Training (WBT) Modules you and your team can learn about how to improve productivity in your organization. Learn how best to utilize your resources, and utilize your staff to their full potential.

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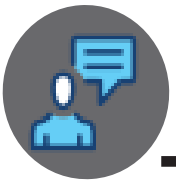
PRODUCTIVITY TRAINING MODULES



Effective Meeting Management

This two part web based training module focuses on planning and executing effective meetings in your organization. Learn why meetings are important, what roles are involved, how to best plan for meetings, and how to overcome challenges that may arise.