

Compliant, Effective, Efficient.

Exclusive Video Interview: Designing a Winning CAPA System

CAPA is a critical Quality System subsystem that, when executed correctly, can provide your organization with a clear payback in terms of improved compliance, effectiveness and operational efficiency.

In this exclusive interview and supplemental article, industry expert Larry Mager discusses the importance of CAPA, and how to design and manage a compliant, effective and efficient CAPA System in your organization.

[Download the video](#)

[Read the full article](#)

PathWise and Irish Medicines Board Offer Intensive CAPA Training in Dublin

Last month, representatives from PathWise and the Irish Medicines Board presented a 2-day, intensive CAPA workshop in Dublin. The event, held at the St. Helens hotel on Stillorgan Road, drew 35 participants from life science companies across Europe.

Course participants learned enhanced Corrective and Preventive Action techniques, while representatives from the Irish Medicines Board shared local regulatory material, including Common Struggles Faced by Irish Manufacturers, Best Practices for Quality System Improvements and Future Policies and Standards.

[Download the Irish Medicines Board Presentation](#)

[Read the full story](#)

New LinkedIn Group! Corrective and Preventive Action (CAPA)

PathWise is pleased to offer a new resource platform and discussion board for life science professionals! Areas of focus include Corrective and Preventive Action and related processes, systems and best practices. Membership is free and features valuable networking opportunities, as well as the latest news and updates impacting the life science industry.

[Join the Corrective and Preventive Action \(CAPA\) Group](#)



PathWise, Inc.

www.PathWise.com

866.580.PATH



**PATHWISE
TRAINING AND
DEVELOPMENT**

Facing an uphill battle?

Pathwise, Inc. offers training and development for medical device and pharmaceutical companies in the areas of quality and compliance. We help our clients to develop and implement standards that improve the manufacturing process for products that serve citizens around the world.

CAPA

Risk Management

Documentation

Root Cause Analysis

Auditing

QSR

Upcoming Training Events

CAPA Requirements and Industry Practice

Arlington, VA

June 22-24, 2009

Integrating ICH Q9 and ISO 14971 into the Quality System

Recorded Webinar

Speaker: Ed Bills

Best Practices to Manage an FDA Inspection

Recorded Webinar

Speaker: Larry Mager

Enroll online at www.PathWise.com or call 866.580.PATH for more information.