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Free Webinar! Fitting Human Factors in the Product Development Process

The FDA has recently stated that more than one third of medical device incidents involve use error, and more than half of device recalls for design problems involve the user interface. These statistics have prompted the agency to strengthen its initiative requiring medical manufacturers to conduct appropriate human factors studies, analyses, and tests.

In this complementary webinar, hear industry expert [Pat Patterson](#) discuss the analysis of use errors and how they can be controlled throughout the product development process.

View the webinar [here](#).

View the article [here](#).

PathWise, Inc.

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How to Prepare for an FDA Investigation

Audits by third parties don't have to be stressful events. The level of preparation within your organization can make a significant difference in your level of success. In this article, we offer some helpful hints and techniques that will enable your organization to perform smoothly during an FDA inspection.

View the full article [here](#).

Will the FDA's Risk Management Model for Drugs Migrate to Devices?

FDA's device center may be moving toward imposing more risk management measures on moderate-risk devices, on the model of recent statutory risk management authorities for drugs, industry lawyers observe.

Read the full article [here](#).

PATHWISE TRAINING AND DEVELOPMENT

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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.

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