ANNOUNCING THE GATEWAYFDA BLOG

September 14, 2011

Entering the U.S. market can be difficult—and when your product is regulated by the U.S. Food and Drug Administration, it can be downright intimidating.

Hodgson Russ is proud to announce the launch of GatewayFDA, a blog that provides analysis, commentary, and resources to help foreign pharmaceutical and medical device companies make sense of ever-evolving FDA regulations and understand the complete range of U.S. legal issues that may complicate bringing a drug, medical device, or other FDA-regulated product or service to market in the United States.

GatewayFDA is written by a team of attorneys experienced in successfully guiding FDA-regulated companies through regulatory, corporate, intellectual property, licensing, product liability, and other critical U.S. legal challenges.

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