This is your new

Medical Devices Law and Regulation Answer Book 2015

Edited by Suzan Onel and Karen M. Becker

Medical Devices Law and Regulation Answer Book 2015 walks you through the current regulatory requirements governing medical devices and describes every aspect from premarket requirements for specific types of devices to postmarket regulation and ongoing government enforcement and investigation. This comprehensive work distills the essential elements of a complex regulatory environment and provides in a single resource a practical guide to the complexities of FDA regulation of medical devices, as well as important related topics not commonly included in surveys of the field.

With over thirty contributors from a variety of law firms and consulting firms specializing in medical device work, this book provides practical guidance on how to handle everyday questions on a wide variety of topics as well as what issues are likely to arise and how to avoid them.

Summary of Contents

1 Overview of the Legal Framework for Medical Device Regulation in the United States
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(continued on reverse)
PLI is proud to publish this practical, concise, and easy-to-use guide to Medical Devices Law and Regulation. If you have any questions or comments, please contact us (see “Questions About This Book?” on the page following the title page).