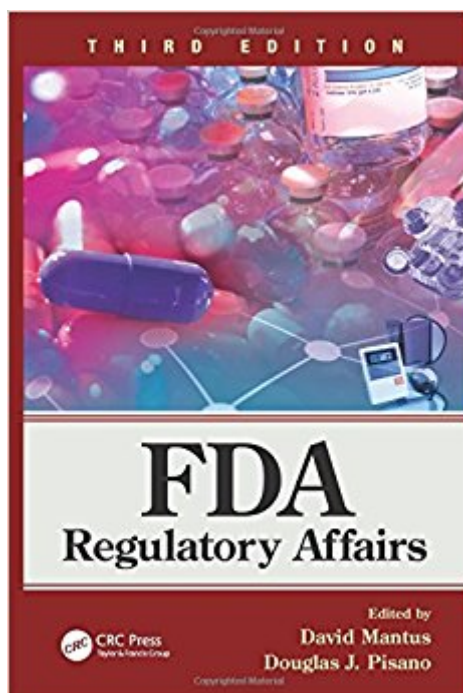


The book was found

# FDA Regulatory Affairs: Third Edition



## Synopsis

FDA Regulatory Affairs is a roadmap to prescription drug, biologics, and medical device development in the United States. Written in plain English, the concise and jargon-free text demystifies the inner workings of the US Food and Drug Administration (FDA) and facilitates an understanding of how the agency operates with respect to compliance and product approval, including clinical trial exemptions, fast track status, advisory committee procedures, and more. The Third Edition of this highly successful publication: Examines the harmonization of the US Federal Food, Drug, and Cosmetic Act with international regulations on human drug, biologics and device development, research, manufacturing, and marketing Includes contributions from experts at organizations such as the FDA, National Institutes of Health (NIH), and PAREXEL Focuses on the new drug application (NDA) process, cGMPs, GCPs, quality system compliance, and corresponding documentation requirements Provides updates to the FDA Safety and Innovation Act (FDASIA), incorporating pediatric guidelines and follow-on biologics regulations from the 2012 Prescription Drug User Fee Act (PDUFA) V Explains current FDA inspection processes, enforcement options, and how to handle FDA meetings and required submissions Co-edited by an industry leader (Mantus) and a respected academic (Pisano), FDA Regulatory Affairs, Third Edition delivers a compilation of the selected US laws and regulations as well as a straightforward commentary on the FDA product approval process thatâ™s broadly useful to both business and academia.

## Book Information

Hardcover: 400 pages

Publisher: CRC Press; 3 edition (January 23, 2014)

Language: English

ISBN-10: 1841849197

ISBN-13: 978-1841849195

Product Dimensions: 1 x 6.2 x 9 inches

Shipping Weight: 1.3 pounds (View shipping rates and policies)

Average Customer Review: 4.0 out of 5 stars 8 customer reviews

Best Sellers Rank: #297,325 in Books (See Top 100 in Books) #21 inÂ Books > Medical Books > Pharmacology > Clinical #67 inÂ Books > Textbooks > Medicine & Health Sciences > Reference > Drug Guides #111 inÂ Books > Medical Books > Medicine > Reference > Drug Guides

## Customer Reviews

"This book covers the myriad of topics related to the development and marketing of all types of

drugs and devices regulated by the U.S. Food and Drug Administration. The book discusses not only the relevant regulations, but also the administrative processes that sponsors must deal with in getting their products approved for marketing. ... This book covers many of the important issues in enough depth to answer basic questions related to the components of an IND (investigational new drug) and an NDA (new drug application), as well as IDE (investigational device exemption) and 510k applications. ... This book fulfills a unique need for those seeking a book for a course in drug and device development or for those in the industry who seek information outside of their area of expertise." •Michael R. Jacobs, BS, PharmD, Temple University School of Pharmacy, in Doody's Book Reviews

David Mantus worked in the biotechnology and pharmaceutical industry for more than 20 years. He not only served as vice president of regulatory affairs at Cubist Pharmaceuticals but also held various regulatory roles at Sention Inc., Shire Biologics, PAREXEL, the Massachusetts Public Health Laboratory, the Massachusetts College of Pharmacy and Health Sciences, and Procter and Gamble Pharmaceuticals. He received his BS in chemistry from the College of William and Mary, his MS and Ph.D in chemistry from Cornell University, and was a post-doctoral research fellow in biomedical engineering at the University of Washington. He is currently associate professor of pharmaceutical sciences and director, regulatory affairs at MCPHS University in Boston. Douglas J. Pisano received his Ph.D in law, policy, and society from Northeastern University. He has participated on the editorial advisory boards of multiple journals and received the Special Service Award for the Enhancement of Regulatory Education from the Regulatory Affairs Professionals Society. A registered pharmacist and active member of several professional organizations, including the American Association of Colleges of Pharmacy and the Regulatory Affairs Professionals Society, he formerly served as dean of the School of Pharmacy at Boston and is currently professor of pharmacy administration, vice-president of academic affairs, and provost at MCPHS University in Boston.

Absolutely loved the book. I looked at it first in a library and then realized that I must own it. Somebody wrote in his/her review that the material seems to be repetitive. Cannot say this not true, but I found it to be a positive feature. Each chapter is written by a different author(s), so some repetition comes from different people trying to cover their topic and share their perspective on things. It was helpful to me personally, as it gives a better picture of the interplay between various topics.

This book has proven, thus far, to be the best reference source I have come across as a student pursuing a Graduate Certificate in Regulatory Affairs. I would highly recommend this book for all students in a Regulatory Affairs program of study. Wish I would have known about this book when I first started the program.

I was not satisfied by this book. First of all, in the chapters there are many repetitions of the same Topics. For instance, the same concept of GMP is repeated across the different chapters, wasting pages and making the Reader bored. This is because each chapter was written by a different Person. So my understanding is that the Editors failed to harmonize the Contents across the book. Moreover, some Topics like a Medical Device PMA are not deeply discussed, and this is by far the most complex aspect of the regulations that me as buyer I was expecting. Finally, the book does not have Workflows to simplify the understanding as well as Examples. Would have been very useful if, for instance, the authors take the example of a recent Approved drug or device and explain more in Details how to do it. PS: In the book you find much more space on the drugs than for device/biologics. So, be careful of what is your interest.

The book was in good standing and it helped me out on my course, Introductory to Regulatory Affairs for Drugs, Biologics and Medical Devices

This is a great book for FDA RA study and would recommend it.

too pricy for the content of information

I was required to purchase this as the assigned textbook for a graduate course. I have not read all of it yet, but have read at least four chapters already. This book contains generally good information; however, it really could have benefited from better editing. It contains numbers of grammatical mistakes and some sections read as if they were first written in another language and then translated into English. The way the chapters have been ordered also strikes me as somewhat illogical. Not the best book I have purchased on this subject.

Great product and seller!

[Download to continue reading...](#)

FDA Regulatory Affairs: Third Edition Regulatory Affairs for Biomaterials and Medical Devices (Woodhead Publishing Series in Biomaterials) The Crisis in Telecommunications Carrier Liability: Historical Regulatory Flaws and Recommended Reform (Topics in Regulatory Economics and Policy) Third Eye: Third Eye Activation Mastery, Easy And Simple Guide To Activating Your Third Eye Within 24 Hours (Third Eye Awakening, Pineal Gland Activation, Opening the Third Eye) ISO 13485:2016, Third Edition: Medical devices - Quality management systems - Requirements for regulatory purposes A Practical Guide to FDA's Food and Drug Law and Regulation, Fifth Edition ServSafe Essentials Korean 5e Update Edition with Answer Sheet, ServSafe Essentials with AnswerSheet Update with 2009 FDA Food Code The FDA and Worldwide Quality System Requirements Guidebook for Medical Devices, Second Edition Passing on Bypass Using External CounterPulsation : An FDA Cleared Alternative to Treat Heart Disease Without Surgery, Drugs or Angioplasty. SECOND EDITION Clinical Trials, Second Edition: Study Design, Endpoints and Biomarkers, Drug Safety, and FDA and ICH Guidelines The AA Effect & Why You've Never Heard of the FDA-Approved Drugs that Treat Alcoholism (Rethinking Drinking Book 1) FDA, You Were Wrong!: Stopping Innovation, Stops America! Protecting America's Health: The FDA, Business, and One Hundred Years of Regulation The FDA and Worldwide Quality System Requirements Guidebook for Medical Devices Product Development Planning for Health Care Products Regulated by the FDA The Circuitous Route by a Group of Novices to a New FDA Approved Cancer Therapy: HOW DID WE DO THIS? New Drugs: An Insider's Guide to the FDA's New Drug Approval Process for Scientists, Investors and Patients Clinical Trials: Study Design, Endpoints and Biomarkers, Drug Safety, and FDA and ICH Guidelines Mastering and Managing the FDA Maze: Medical Device Overview: A Training and Management Desk Reference for Manufacturers Regulated by the Food and Drug Administration Corporate Warriors: The Rise of the Privatized Military Industry, Updated Edition (Cornell Studies in Security Affairs)

[Contact Us](#)

[DMCA](#)

[Privacy](#)

[FAQ & Help](#)