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Code Of Federal Regulations, Title 21, Food And Drugs, Pt. 200-299, Revised As Of April 1, 2016



Synopsis

The Code of Federal Regulations is a codification of the general and permanent rules published in the Federal Register by the Executive departments and agencies of the United States Federal Government. CFR Title 21, Parts 200-299 include labelling, prescription drug advertising, prescription drug marketing, registration of producers of drugs and listing of drugs in commercial distribution, medication guides to prescription drugs, pharmacy compounding, controlled drugs, drugs: official names and established names, and more. Other related products: Drug Master File (Blue Polyethylene Folder) is available

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Book Information

Paperback: 247 pages

Publisher: Office of the Federal Register; Revised edition (July 21, 2016)

Language: English

ISBN-10: 0160932726

ISBN-13: 978-0160932724

Product Dimensions: 5.8 x 0.5 x 9.1 inches

Shipping Weight: 1.1 pounds

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