



Code of Federal Regulations Title 21, Volume 4, April 1, 2015 (Paperback)

By -

Regulations Press, 2016. Paperback. Condition: New. Language: English . Brand New Book ***** Print on Demand *****.21 CFR Parts 200-299 covers prescription drug products for human use, manufacturing, labeling, processing, and packing practices, official drug names, and more. Code of Federal Regulations Title 21, Volume 4, April 1, 2015 Containing parts Parts 200 to 299 Part 200; GENERAL Part 201; LABELING Part 202; PRESCRIPTION DRUG ADVERTISING Part 203; PRESCRIPTION DRUG MARKETING Part 205; GUIDELINES FOR STATE LICENSING OF WHOLESALE PRESCRIPTION DRUG DISTRIBUTORS Part 206; IMPRINTING OF SOLID ORAL DOSAGE FORM DRUG PRODUCTS FOR HUMAN USE Part 207; REGISTRATION OF PRODUCERS OF DRUGS AND LISTING OF DRUGS IN COMMERCIAL DISTRIBUTION Part 208; MEDICATION GUIDES FOR PRESCRIPTION DRUG PRODUCTS Part 209; REQUIREMENT FOR AUTHORIZED DISPENSERS AND PHARMACIES TO DISTRIBUTE A SIDE EFFECTS STATEMENT Part 210; CURRENT GOOD MANUFACTURING PRACTICE IN MANUFACTURING, PROCESSING, PACKING, OR HOLDING OF DRUGS; GENERAL Part 211; CURRENT GOOD MANUFACTURING PRACTICE FOR FINISHED PHARMACEUTICALS Part 212; CURRENT GOOD MANUFACTURING PRACTICE FOR POSITRON EMISSION TOMOGRAPHY DRUGS Part 216; PHARMACY COMPOUNDING Part 225; CURRENT GOOD MANUFACTURING PRACTICE FOR MEDICATED FEEDS Part 226; CURRENT GOOD MANUFACTURING PRACTICE FOR TYPE A MEDICATED ARTICLES Part 250; SPECIAL REQUIREMENTS FOR SPECIFIC HUMAN DRUGS Part 290;...



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