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The bestselling and most useful aid available for finding all references to FDA and DEA regulations, Interpharm Master Keyword Guide: 21 CFR Regulations of the Food and Drug Administration, is used in hundreds of active pharmaceuticals, pharmaceutical, biotechnology, diagnostic, and device manufacturing companies. And it is in use by every FDA district in the United States to sort their way through their own regulations. Each of the over 20,000 entries is quoted in context to provide instant access to every noun, phrase, and concept used by the DEA and FDA. The KEYWORD and SECTION TITLE are shown in upper case, the Subpart Title and/or Part Title are shown in capitals and lower case.

How to use this guide:

1. Look up the keyword of interest
2. Note the context in which the keyword is mentioned in the section of title and the details of the subpart or part title to determine if it is the reference you need
3. When you find the correct reference, use the section number provided to look up the details of the regulations in the Code of Federal Regulations Title 21

Updated to include the latest changes in 21 CFR, the Interpharm Master Keyword Guide: 21 CFR Regulations of the Food and Drug Administration, 2002-2003 Edition makes it easy to find the exact section you need and apply it correctly.

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