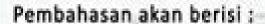


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21CFR58 regulatory requirements from leading industry experts with a vast knowledge and expertise in their fields The approach used takes the regulations provides interpretations and references to examples and regulatory actions Data integrity and the use of electronic systems in compliance with 21CFR11 Electronic Records Electronic Signatures are also discussed Unique volume covering FDA inspections of GLP facilities Provides a detailed interpretation of GLP Regulations Presents the latest on electronic data management in GLP Describes GLP and computer systems validation Can be referenced repeatedly in supporting daily hands on implementation of the CFR requirements *Good Laboratory Practice Regulations, Third Edition, Revised and Expanded* Sandy Weinberg,2002-11-06 Fully updated and revised to include the latest information since publication of the first edition in 1989 the Second Edition of this highly praised reference covers all aspects of the Food and Drug Administration s FDA Good Laboratory Practice GLP regulations and techniques for implementation The book details specific standards and general guidelines for the management of efficient and effective research environment A guide to the current standards and requirements of good laboratory management the book examines essential theoretical principles for anticipating new and emerging interpretations of GLP in a variety of laboratory settings

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