

# KEY COMPONENTS OF GOOD LABORATORY PRACTICES (GLP)

Organization and Personnel



Building and Facilities



Equipment and Machinery



Standard Operating Procedures (SOPs)



Sample Handling



Method Validation



**Quality Control** 



Data Management



Compliance & Ethics



Continuous Improvement



## **Guide To Good Laboratory Practices**

**LP Steffe** 

#### **Guide To Good Laboratory Practices:**

**GLP Essentials** Milton A. Anderson.2002 **Reg-words** U.S. Food and Drug Administration, 1978 Good Laboratory Practice (GLP) eRegs & Guides - For Your Reference Book 1 eRegs & Guides, 2014-10-25 Good Laboratory Practice GLP 21 CFR Title 58 Good Laboratory Practice for Non Clinical Laboratory Studies 21 CFR Title 9 Animals and Animal Products PART 1 Definition of Terms 21 CFR Title 9 Animals and Animal Products Part 2 Regulations 21 CFR Title 9 Animals and Animal Products Part 3 Standards 21 CFR Title 29 Part 1910 1450 Occupational exposure to hazardous chemicals in laboratories 21 CRF Title 29 Labor 1910 1 1910 9 21 CFR Title PART 11 Electronic Records Electronic Good Laboratory Practice Regulations Sandy Weinberg, 2007-01-19 Recent changes in the interpretation Signatures and enforcement of 21 CFR Part 11 have shifted the focus of Good Laboratory Practice GLP regulations to concentrate on the acceptance of electronic signatures the archiving of data the security of electronic documents and the automation of laboratory procedures This all encompassing Fourth Edition addresse 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 Good Laboratory Practice OECD Principles and Guidance for Compliance Monitoring OECD, 2005-12-16 This publication unites all of the OECD documents related to Good Laboratory Practice and compliance monitoring and in the Annex reproduces the three OECD Council Decisions related to the Mutual Acceptance of Data in the Assessment of Chemicals The Indispensable Guide to Good Laboratory Practice (GLP) Mark Gregory Slomiany Ph. D., Mark Gregory Slomiany, 2009-10 Familiarity with Good Laboratory Practice GLP is a necessity for those working in or aspiring to work in biotech pharma and even academic research where partnerships with industry require ever greater awareness of drug development regulations. Thus this guidebook serves to not only summarize GLP regulations but explain the interaction of these various components in the development of a GLP compliant study Far from being long winded special attention has been paid to the brevity of each chapter If more detail is needed the reader can easily refer to the official FDA verbiage provided below each chapter summary Inspired by the lack of affordable succinct and easily accessible training options available to students scientists and entrepreneurs Dr Slomiany set out to create a guidebook which conveys the underlying principles of GLP in an easily digestible format while providing resources for more in depth study A Laboratory Quality Handbook of Best Practices Donald C. Singer, 2001-09-25 Based on the work of a collection of experts from the laboratory science and quality assurance fields A Laboratory Quality

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