

GOOD LABORATORY PRACTICES

GLP Principles



Good Laboratory Practice Regulations

USA. Food and Drug Administration



Good Laboratory Practice Regulations:

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market The GLP regulations have developed from rules on how to exactly document the planning conduct and reporting of toxicity studies to a quality system for the management of a multitude of study types from the simple determination of a physical chemical parameter to the most complex field studies or ecotoxicology studies At the same time the term Good Laboratory Practice has become somewhat of a slogan with the aim to characterise any reliably conducted laboratory work

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Orientation Overview of GCP and Introduction to GLP Part I Federal Regulations Relating to Good Laboratory Practice o Parts 58 Good Laboratory Practice for Nonclinical Laboratory Studies o 1987 Final Rule Good Laboratory Practice Regulations Part II Guidance Documents o Bioresearch Monitoring Good Laboratory Practice o Good Laboratory Practices Questions and Answers Part III Redbook 2000 o IV B 1 General Guidelines for Designing and Conducting Toxicity Studies o IV B 2 Guidelines for Reporting the Results of Toxicity Studies Reference Tools Part IV Combined Glossary and Index

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