



Good Clinical Practice: Pharmaceutical, Biologics, and Medical Device Regulations and Guidance Documents Concise Reference; Volume 2, Guidance

Mindy J. Allport-Settle

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Selected FDA GCP/Clinical Trial Guidance Documents Grouped by Topic: * FDA Overview and Orientation * Introduction to GCP * Part I: General * Part II: Institutional Review Boards (IRBs) and Informed Consent * Part III: Drugs and Biologics * Part IV: Medical Devices * Part V: Manufacturing Requirements for Investigational Products * Part VI: Electronic Data Reference Tools * Part VII: Combined Glossary and Index for all Quality Guidance Documents

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