Preambles to GCP Regulations

Preambles to Good Clinical Practice Regulations for Drug Studies

Contains preambles to the following sections of Title 21 (Food & Drugs) of the Code of Federal Regulations:

Part 50:	Protection of human subjects
Part 54:	Financial disclosure by clinical investigators
Part 56:	Institutional Review Boards
Part 312:	Investigational new drug application
Part 314:	Applications for FDA approval to market a new drug

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CLINICAL RESEARCH RESOURCES