

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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