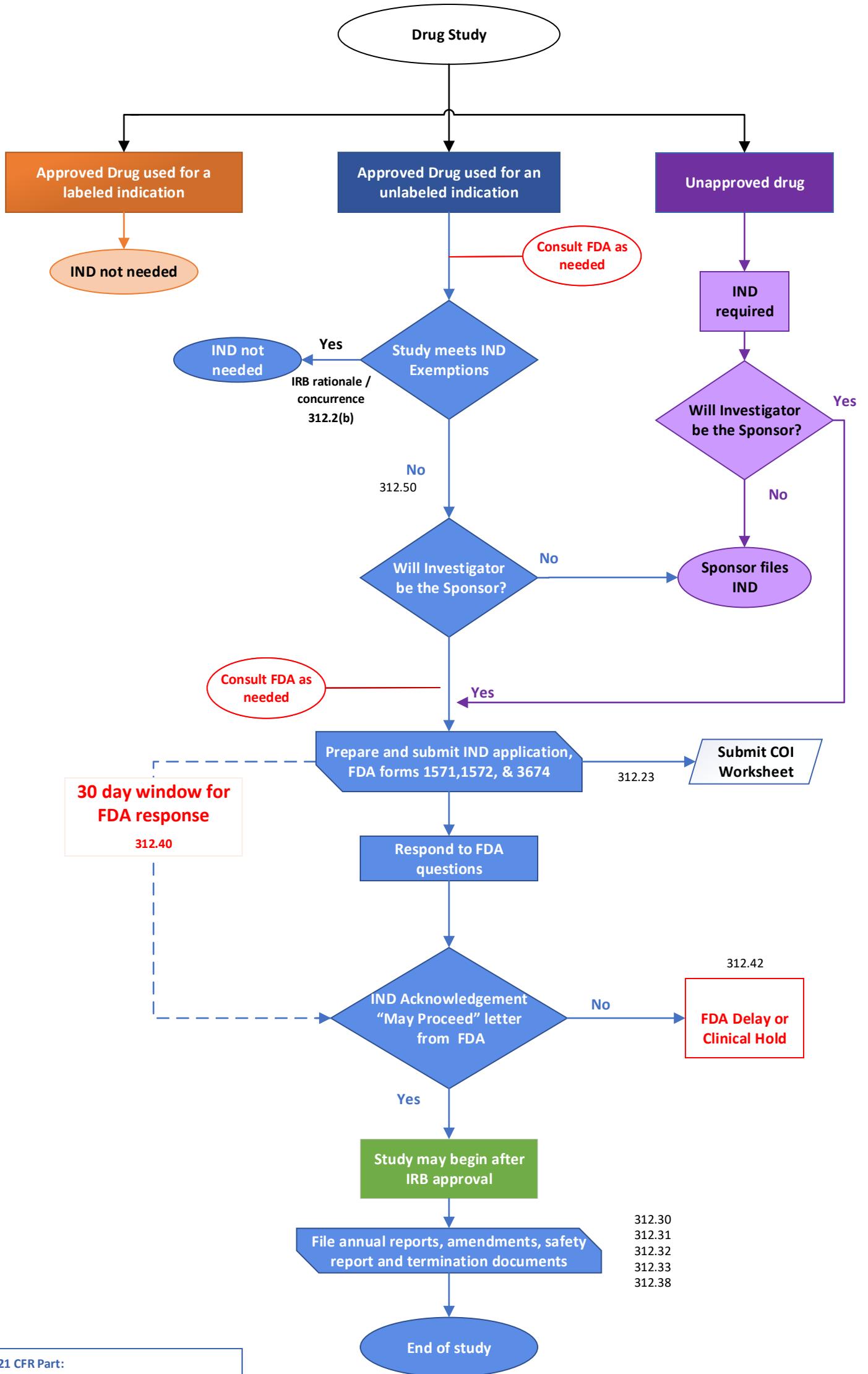


# Sponsor / Investigator IND Flowchart for Clinical Drug Study



- 21 CFR Part:**
- 312.2(b) Exemptions
  - 312.50 Investigator Responsibilities
  - 312.23 IND Content and Format
  - 312.40 Administrative Actions
  - 312.42 Clinical Hold
  - 312.30 Amendments
  - 312.31 Information Amendments
  - 312.32 Safety Reports
  - 312.33 Annual Reports
  - 312.38 Withdrawal IND

- 312.30
- 312.31
- 312.32
- 312.33
- 312.38