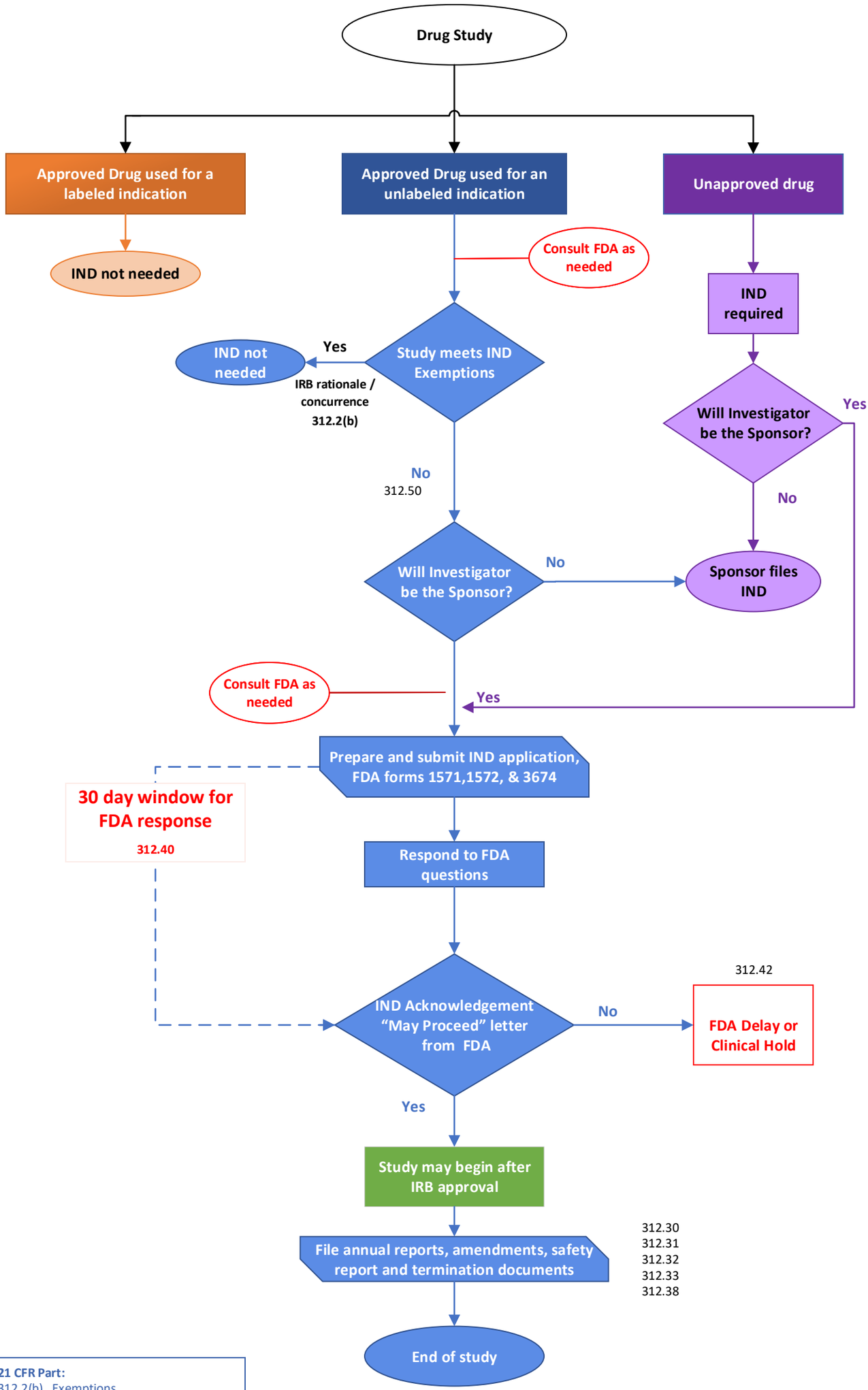
Sponsor / Investigator IND Flowchart for Clinical Drug Study



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