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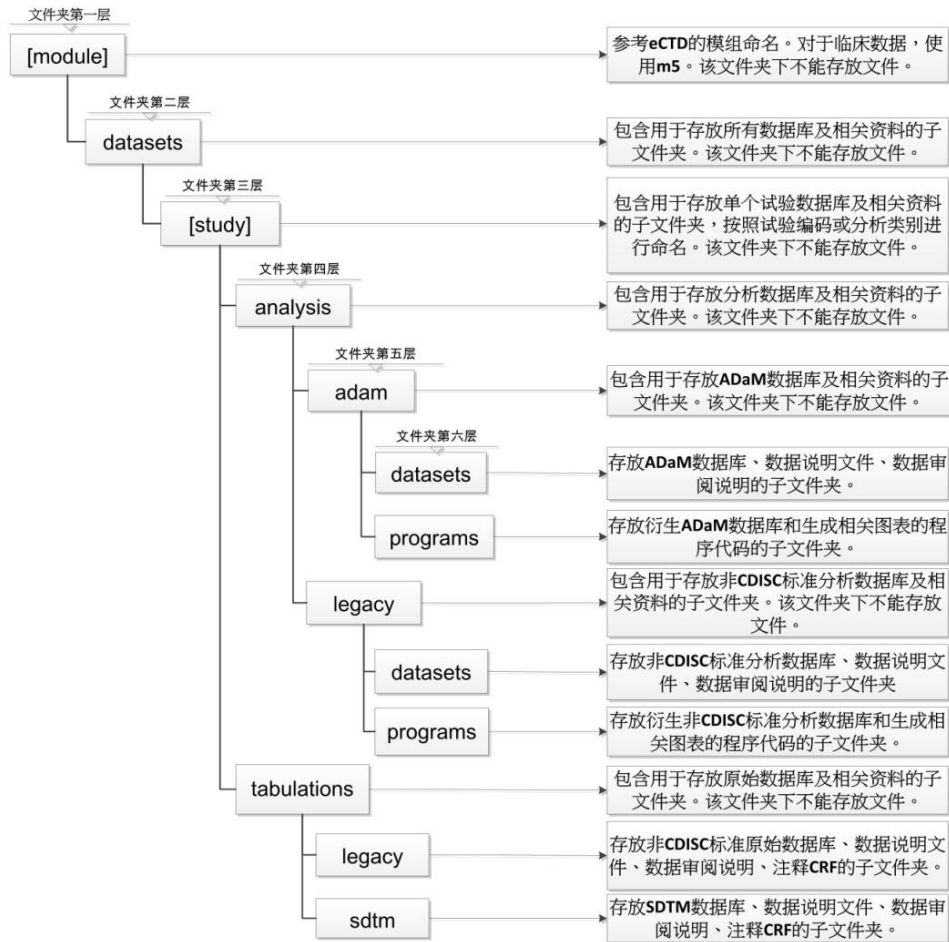
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□□□□□	PDF (Portable Document Format)
□□□□	CRF (Case Report Form)
□□□□□□	eCTD (electronic Common Technical Document)
□□□□□□	ADaM (Analysis Data Model)
□□□□□□□□□□	ICH (International Council for Harmonization of Technical Requirements for Pharmaceuticals for Human Use)
□□□□□□	W3C (World Wide Web Consortium)
□□□□□□	MedDRA (Medical Dictionary for Regulatory Activities)
□□□□□□	XML (Extensible Mark-up Language)
□□□□□□	XSL (Extensible Stylesheet Language)
□□□□□□□□	CDISC (Clinical Data Interchange Standards Consortium)
□□□□□	CSR (Clinical Study Report)
□□□□□□□□	ADSL (Subject Level Analysis Dataset)
□□□□	NDA (New Drug Application)
□□□□□	STF (Study Tagging File)
□□□□□□	SDTM (Study Data Tabulation Model)
□□□□□□	aCRF (annotated Case Report Form)

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[1]. CFDA. □□□□□□□□□□ . 2016 □ 7 □ .

[2]. FDA. Study Data Technical Conformance Guide. Mar 2020.

[3]. PMDA. Revision of Technical Conformance Guide on Electronic Study Data Submissions. Jan 2019.

[4]. CDISC. Study Data Tabulation Model Implementation Guide. Nov 2018.

[5]. CDISC. Analysis Data Model Implementation Guide. Oct 2019.

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