

U.S. FOOD & DRUG ADMINISTRATION

TITLE LINE 1 TITLE LINE 2 TITLE LINE 3

Document Version: \$VERSION Document Version Date: \$VERSIONDATE

Version History

Version Number	Implemented By	Revision Date	Approved By	Description and Reason for Change
1.0	Kyle Thomsen	09 Dec 2019		First version of FDA document template.
1.1	Kyle Thomsen	10 Dec 2019		Modularized template using the subfiles package.
1.2	Kyle Thomsen	11 Dec 2019		 Created the versionRecord command in order to facilitate the creation of changelog entries, such as this one. Created the keyTermRecord command in order to facilitate the creation of tables containing key terms and definitions. Created the referenceRecord command in order to facilitate the creation of reference tables. However, I am contemplating substituting this with one of the usual LATEX citation or reference environments.
\$VERSION	Aurotech Corp.	\$VERSIONDATE		\$CHANGELOG

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1 Subfile Section One

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Appendix A: Key Terms

The following table provides definitions and explanations for terms and acronyms relevant to the content presented within this document.

Term	Definition
ADSL	Subject-Level Analysis Data
ASCII	American Standard Code for Information Interchange
CBER	Center for Biologics Evaluation and Research
CDASH	Clinical Data Acquisition Standards Harmonization
CDER	Center for Drug Evaluation and Research
CDISC	Clinical Data Interchange Standard Consortium
CSR	Clinical Study Report
eCTD	Electronic Common Technical Document
FDA	Food and Drug Administration
GLP	Good Laboratory Practice
ICH	International Council for Harmonisation
IND	Investigational New Drug
NDA	New Drug Application
OIMT	Office of Information Management and Technology
OND	Office of New Drugs
PDF	Portable Document Format
RG	Reviewer Guides
SAS	Statistical Analysis System
XML	eXtensible Markup Language
XPORT	SAS Transport Version 5
a term	a definition
another term	another definition

Appendix B: References

The following table summarizes the documents referenced in this document.

Document Name	Description	Location
a document	it's a document	document.org - accessed 01 Jan 20XX
CDER Manual of Policies & Procedures	CDER's Manual of Policies and Procedures (MAPPs) are federal directives and documentation of internal policies and procedures. MAPPs are required by law and made available to the public to make CDER a more transparent organization.	https://www.fda. gov/about-fda/ center-drug-evaluation-and-research-c cder-manual-policies-procedures-mapp (accessed 22 Nov 2019)



Appendix C: System Design Document Approval

The undersigned acknowledge that they have reviewed the **DIMES DevSecOps and Data Sharing Data Sharing Document** and agree with the information presented within this document. Changes to this **Data Sharing Document** will be coordinated with, and approved by, the undersigned, or their designated representatives.

Signature:	Date:
Print Name:	
Title:	
Role:	Project Manager