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# **PRESENTER**

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CDER | US FDA

Lina Cong has a medical and computer background with over ten years of experience on study data standards and study data submissions. She also has ten years of experience on clinical trial data analysis and clinical data management within the pharmaceutical industry.



# Technical Rejection Criteria for Study Data (TRC) and Beyond

PharmaSUG SDE

October 21, 2022

Presenter: Lina Cong

Co-host: Jiang Xu



# Agenda



- Technical Rejection Criteria for Study Data (TRC)
- Tools to Help Industry Pass TRC Validation
- Study Data Validation Overview
- Addressing Common TRC Errors:
  - Error 1734
  - Error 1735
  - Error 1736
- Summary
- Frequently Asked Questions





# Technical Rejection Criteria for Study Data (TRC)



# Technical Rejection Criteria for Study Data (TRC) – What's New



- ❖ CBER Non-clinical study requirements will start after March 15, 2023
- Rule 1734 will no longer check for study ID matching

- Changes to other study data validations:
  - > New Validation Rule 1738 for study ID matching
  - ➤ Change of scope for Validation Rule 1737



# eCTD Validation Updates



Study Data Validation Effective Date for CBER Module 4 Sections: 3/16/2023

Error	Severity	Description
1734	High	A dataset named ts.xpt with information on study start date must be present for each study in required sections*
1735	High	Correct STF file-tags must be used for all standardized datasets and corresponding define.xml files in required sections*
1736	High	For SEND data, DM dataset and define.xml must be submitted in Module 4 required sections*  For SDTM data, DM dataset and define.xml must be submitted in Module 5 required sections*  For ADaM data, ADSL dataset and define.xml must be submitted in Module 5 required sections*

\* Module 4 sections: 4.2.3.1, 4.2.3.2, 4.2.3.4 Module 5 sections: 5.3.1.1, 5.3.1.2, 5.3.3.1, 5.3.3.2, 5.3.3.3, 5.3.3.4, 5.3.4, 5.3.5.1, 5.3.5.2

Please review eCTD Validation Specification all details are not included in this presentation



# eCTD Validation Updates



- Effective date for changes to Rule 1737 and new Rule 1738: 3/16/2023
- Applies to clinical and non-clinical study data submitted to CDER or CBER as part of an NDA, BLA, IND, ANDA, or master file submission

Error	Severity	Description
1737	Medium	For each study referenced by an STF file in Module 4 and Module 5, no more than one dataset of the same file name and leaf title should be submitted using the lifecycle operator 'new'*
1738	Medium	Study ID for STF should match STUDYID listed in the referenced Trial Summary (ts.xpt) file*

Please review eCTD Validation Specification all details are not included in this presentation

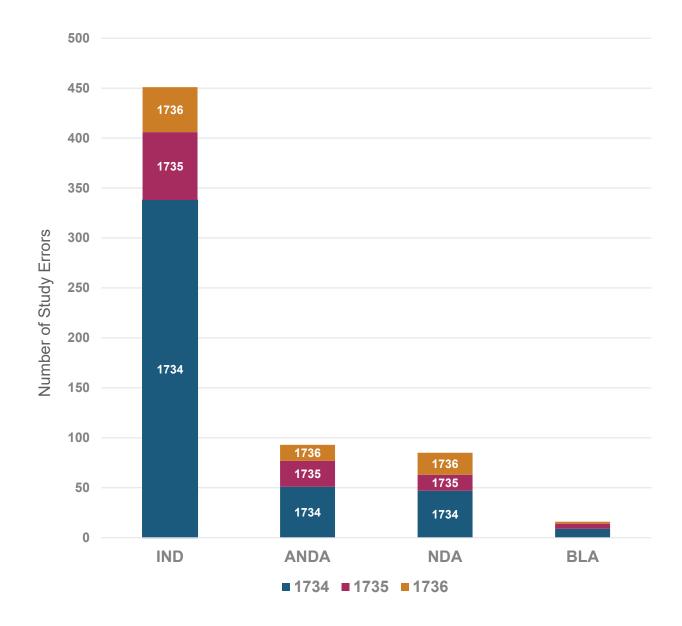


<sup>\*</sup> Applies to all sections except 4.3, 5.2, 5.4, & 5.3.6 Postmarketing reports

## Rejected Submissions

FDA

- 1734 is the most common error and rejection reason for a missing ts.xpt
- Commercial IND submissions have the highest number of errors and rejections overall



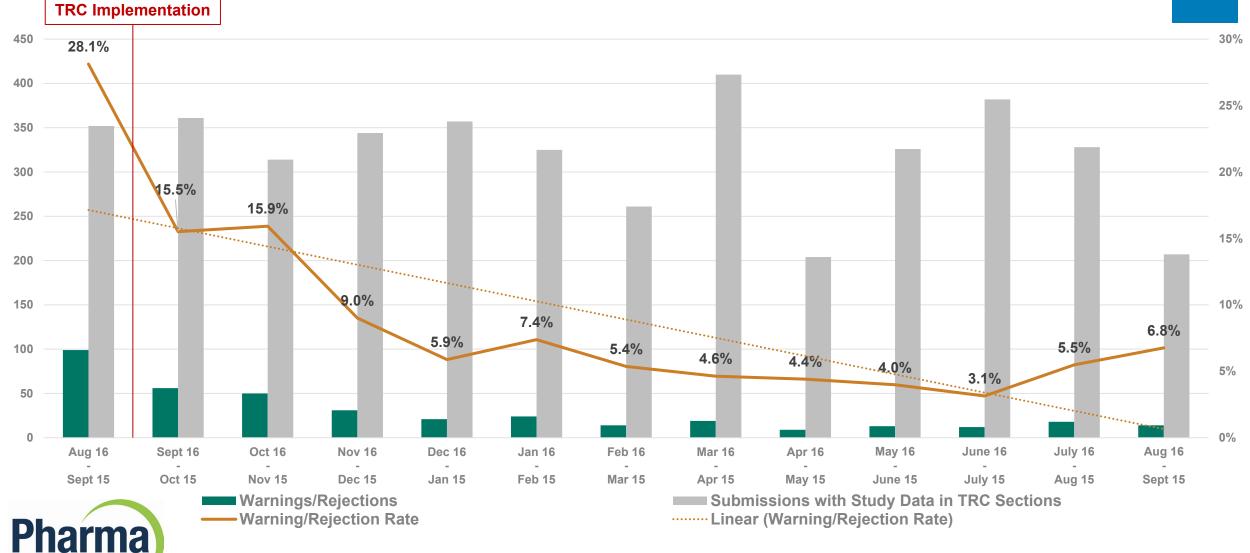


**Notes:** Metrics generated from data between September 15, 2021 - September 15, 2022 CDER began rejecting submissions with TRC errors on September 15, 2021

### **CDER Rejection Rate Trend**

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Notes: Metrics generated from data between August 16, 2021 - September 15, 2022

CDER began rejecting submissions with TRC errors on September 15, 2021

The Aug 16 – Sept 15, 2021 time period includes warnings for TRC errors, not rejections

Warnings/Rejections include submissions rejected due to TRC errors 1734, 1735, and 1736



# Tools to Help Industry Pass TRC Validation



#### The Self-Check Worksheet

FDA

- Designed to walk sponsors through each step of the TRC validation process
- Dynamically guides sponsors through study data requirements based on study information entered
- Helps sponsors prepare study data to submit to the FDA for the first time

**Demonstration Videos & Other Supporting Material** 

<u>Technical Rejection Criteria Self-Check Worksheet</u>



**Self-Check Worksheet Instructions** 

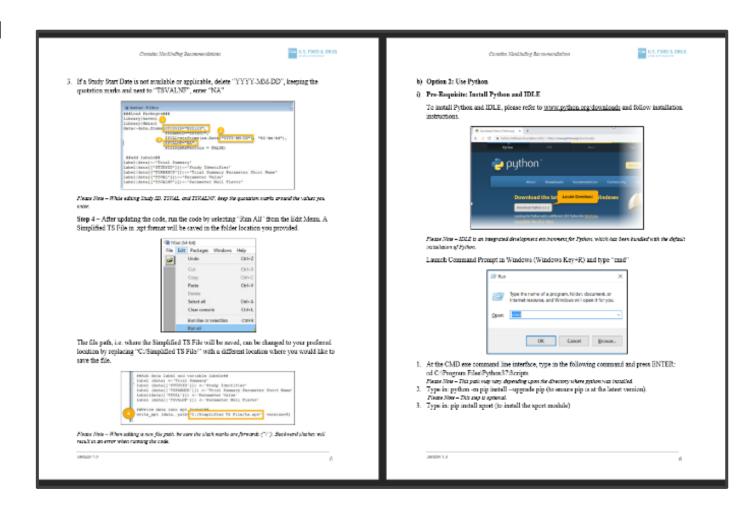
DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration  SELF-CHECK WORKSHEET FOR STUDY DATA PREPARATION						
	ksheet is not required for submissions of st					
prepare newly submitted s	study data to FDA, i.e. studies for which no	files have been previously submitted.				
*Required Field						
Section 1: Application &	Submission Information					
1a. FDA Center* 1b.	Application Type*	1c. Application Number*				
CDER CBER	NDA BLA ANDA Commercial	IND				
1d. eCTD Sequence Number	1e. eCTD Submission Type	1f. eCTD Submission Sub Type				
Note: Repeat Sections 2	through 5 for each study included in the s	submission.				
Section 2: Study Informa	tion					
2a. Study ID*						
za. Glady ib						
	er across application documents. Therefore, the stud study, i.e. STF File, ts.xpt, dm.xpt, etc.)	ly ID must be consistent across all the files				
Yes No	Data is Being Submitted for This Study as Part of					
If you answered "No" in Field 2	2b, do not proceed. This self-check worksheet is des	signed for newly submitted study data.				
2c. Title of the Study						
2d. Study Section - eCTD Heading (Example: m4-2-1-1) <sup>x</sup>						
2e. Module*						
Nonclinical (m4) Clinical (m5)						
2f. Study Dataset Type(s)*						
Tabulation Analysis Other						
If you are submitting tabulation data select "Tabulation." If you are submitting analysis data, select "Analysis." For other types of data, such as Listings datasets, when tabulation or analysis data is not being submitted, select "Other." Additional details and examples are included in the Study Data Self-Check Worksheet Instructions.						
FORM FDA 4061 (11/19)	Page 1 of 3	PSC Publishing Services (301) 445-6740 El				

# The Simplified ts.xpt Creation Guide



- Helps industry create simplified TS files using free and open-source software, R and Python
- Provides step by step instructions to install the necessary software
- Users can copy and paste code samples from the guide into R or Python
- Available on FDA's web page, <u>Study Data</u> for Submission to CDER and CBER
- Demonstration video also available at <u>Study Data for Submission to CDER and CBER</u>
- Additionally, a publicly available tool was developed by PHUSE:

Simplified ts.xpt File Generator (https://geotiger.shinyapps.io/07\_genTS/)





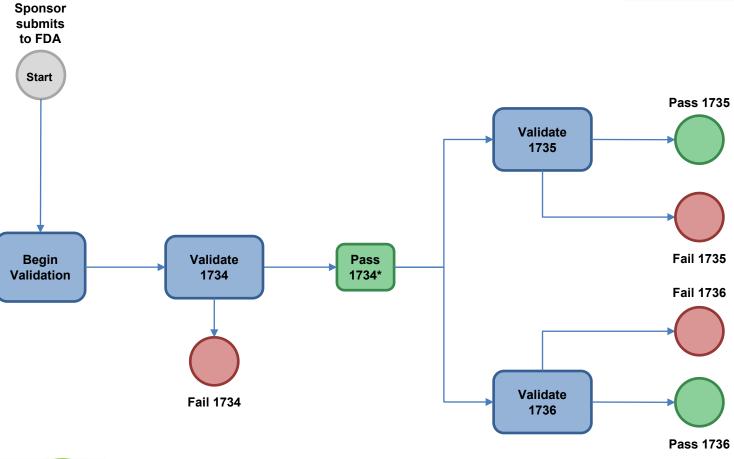
# Study Data Validation Overview



#### TRC Validation Rule Flow









<sup>\*</sup> Not every submission will proceed beyond 1734 to 1735 and 1736, depending on the Study Start Date (SSD)

# Validation Rule Flow: 1737, 1738, & 1789

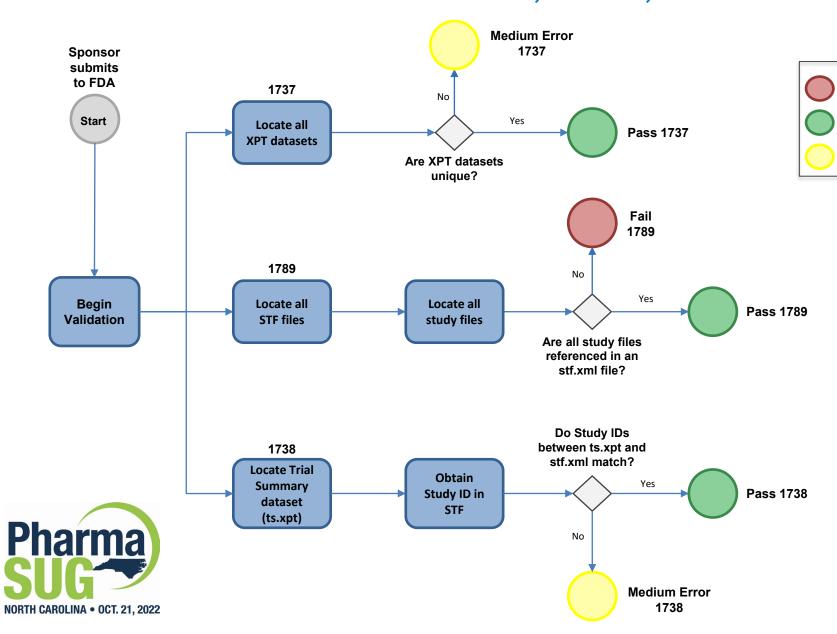


Legend

High validation errors lead to rejection of the submission

Medium severity errors do not lead to a rejection

Validation passes





# Addressing Common TRC Errors Error 1734



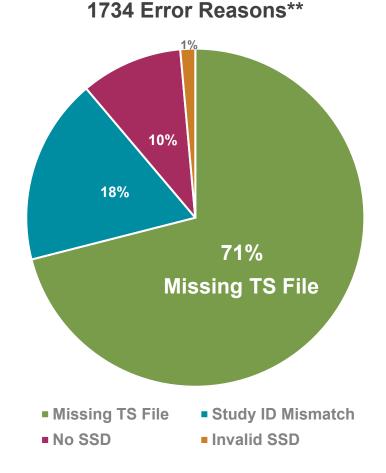
#### Validation Rule 1734

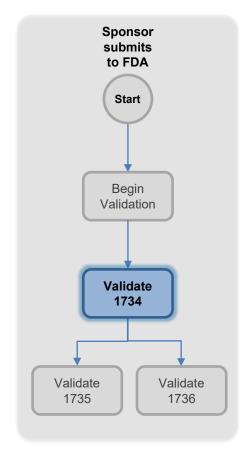


A dataset named ts.xpt with information on study start date must be present for each study in required sections\*

- ✓ Trial Summary Dataset (ts.xpt) is present
- ✓ Study ID (or SPREFID) matches STF Study ID<sup>^</sup>
- ✓ Study start date is provided (or TSVALNF = NA)
- ✓ Study start date is in a valid format

^Moving to Rule 1738, effective 3/16/2023











Providing a TS File for non-clinical studies which require a Simplified TS will address the biggest root cause of Error 1734.

Over 70% of Missing TS File Errors are for non-clinical studies with study reports and no .xpt datasets\*

	M4	M5
Studies with only study reports	73%	N/A
Studies with only study data	0%	8%
Studies with study data and reports**	17%	NA



#### **Option 1**

Use the Simplified ts.xpt Creation Guide to generate a simplified ts.xpt in R or Python:

#### Option 2

Use publicly available tool developed by PHUSE to generate simplified ts.xpt files:

Example of a Simplified TS file for a non-clinical study:						
•	STUDYID	TSPARMCD	TSVAL	TSVALNF		
1	S107	STSTDTC	2014-10-26			



# Addressing Common TRC Errors Error 1735

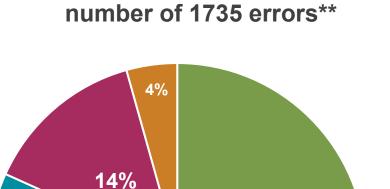


#### Validation Rule 1735

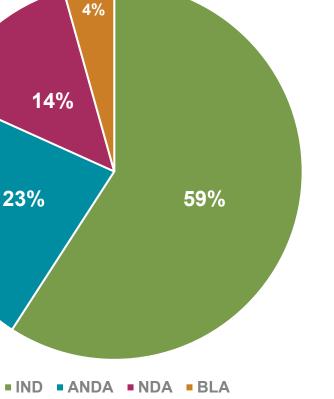
The correct STF file-tags must be used for all standardized datasets and corresponding define.xml files in required sections\*

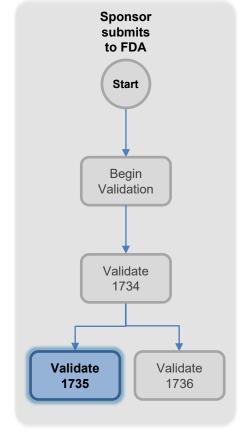
- ✓ Valid file tags for XPT files (e.g., adsl.xpt, dm.xpt) are tagged as:
  - "data-tabulation-dataset-sdtm" for SDTM
  - "analysis-dataset-adam" for ADaM
  - "data-tabulation-dataset-send" for SEND
- ✓ Valid file tags for corresponding define.xml files:
  - "data-tabulation-data-definition" for SDTM & SEND
  - "analysis-data-definition" for ADaM

#### IND submissions have the highest number of 1735 errors\*\*



23%







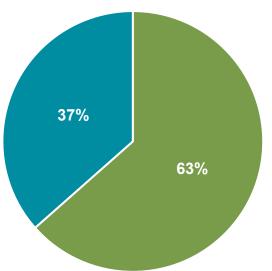
## Addressing the Most Common 1735 Error



**Tags** 

- ❖ The most common cause of 1735 errors is incorrectly tagged define.xml files
- **❖** When preparing STF files, ensure files are tagged properly

#### 1735 Error Reasons\*



- Invalid File Tag for define.xml
- Invalid File Tag for dm.xpt or adsl.xpt

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#### **Example Study Tagging File (STF) for SDTM:**

Sponsors commonly apply the same file tags as datasets or other files for all files submitted, including define.xml files

```
<doc-content
                xlink:href="../../../../../../../../../o001/index.xml#adf5f97a527ef1507a6db8fca2f8b4867" xlink:type="simple">
                <file-tag info-type="us" name="data-tabulation-dataset-sdtm"/>
  dataset domains
Standardized
    (e.g. dm.xpt)
            </doc-content>
            <doc-content
                xlink:href="../../../../,../0001/index.xml#ab54f98276616b94d1d30fa071ffffc36" xlink:type="simple">
                <file-tag info-type="us" name="data-tabulation-dataset-sdtm"/>
            </doc-content>
            <doc-content
                xlink:href="../../../../../../../../0001/index.xml#a7794eaba0442a7c66cbf122fb66ff932" xlink:type="simple">
                <file-tag info-type="us" name="data-tabulation-dataset-sdtm"/>
            Define.xml
            <doc-content
                xlink:href="../../../<del>c./0001/index.xml#a57bb2ed13e2d2feb76</del>06e65d59586355" xlink:type="simple">
                <file-tag info-type="us" name="data-tabulation-dataset-sdtm/>
                                                                                                             Correct
                                                                                                                File
```

File tag for define.xml needs to be corrected to:

"data-tabulation-data-definition"



# Addressing Common TRC Errors Error 1736



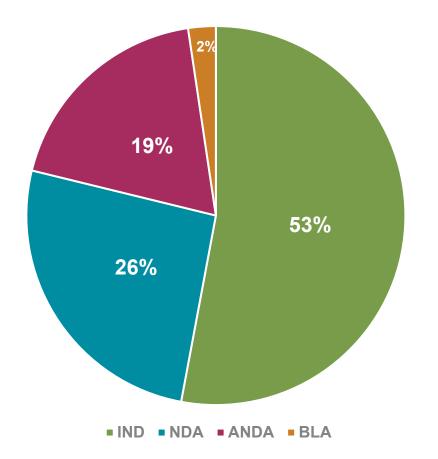
#### Validation Rule 1736

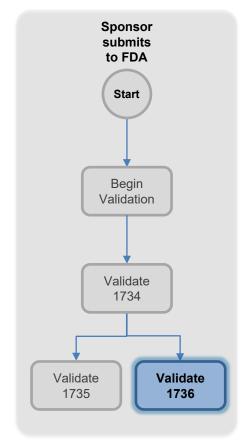


DM datasets and define.xml files are submitted with SDTM and SEND (tabulations) data and an ADSL dataset and define.xml file are submitted with ADaM (analysis) data.

- ✓ For SEND, a DM dataset and define.xml must be submitted in module 4\*
- ✓ For SDTM data, a DM dataset and define.xml must be submitted in module 5\*
- ✓ For ADaM data, an ADSL dataset and define.xml must be submitted in module 5\*

# IND submissions have the highest number of 1736 errors\*\*





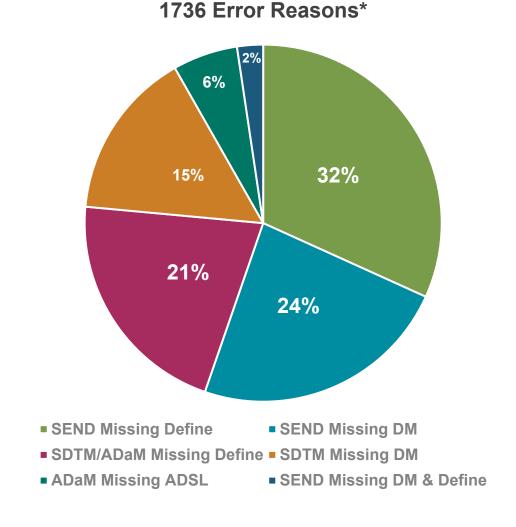


## Addressing the Most Common 1736 Error



The most common cause of 1736 errors is missing define.xml and/or dm.xpt files for non-clinical studies requiring SEND standardized datasets

	SEND	SDTM/ADaM
Missing Datasets	20	18
Missing Define	27	18
Both	2	-
Total	49 (58%)	36 (42%)







# Summary



# Summary: Addressing Top 3 Causes of TRC Errors

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	1734		1735	1736
Impact	All 1734 71% of Rejections	Comm. IND 52% of Rejections	17% of Rejections	12% of Rejections
Rule Summary	A dataset named ts.xpt with information on study start date must be present for each study in required sections		The correct STF file-tags must be used for all standardized datasets and corresponding define.xml files	Checks that DM datasets and define.xml files are submitted with SDTM and SEND (tabulations) data and an ADSL dataset and define.xml file are submitted with ADaM (analysis) data
1. Check if your study has an error	Self-Check Worksheet Sections 3 & 4		Self-Check Worksheet Section 5	Self-Check Worksheet Section 5
2. Correct the required, utili errors ts.xpt Cre		ed TS file is e the <i>Simplified</i> fion <i>Guide</i> or USE Utility	Ensure the correct STF file- tags for standardized datasets and define.xml files are used	Ensure that DM, ADSL datasets, and define.xml files are submitted with corresponding data

**Notes:** Metrics generated from data between September 15, 2021 - September 15, 2022 CDER began rejecting submissions with TRC errors on September 15, 2021



# Frequently Asked Questions







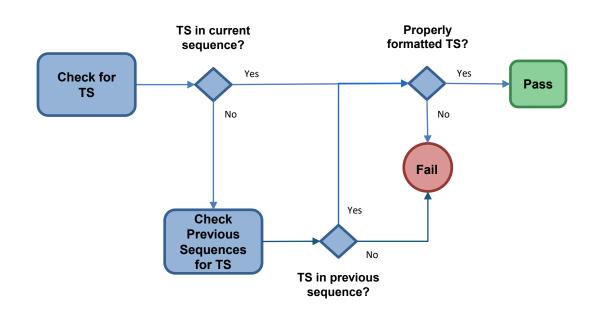
- It is not required to submit a ts.xpt with every submission
- Previously submitted ts.xpt must be standardized and properly formatted
- Validations check previously submitted files for Rule 1734 (ts.xpt) and Rule 1736 (dm.xpt, adsl.xpt, and define.xml)

#### **Properly Formatted TS:**

- ✓ Study ID (or SPREFID) matches STF Study ID<sup>^</sup>
- ✓ Study start date is provided (or TSVALNF = NA)
- ✓ Study start date is in a valid format

^Moving to Rule 1738, effective 3/16/2023







# How can multiple sets of data be submitted for the same study?

- Option 1: Submitting multiple sets of data under the same STF
  - Creating sub-folders for each set of data
  - > Assigning different leaf titles for each set of data
- Option 2: Submitting multiple sets of data under different STFs
  - ➤ Using different STFs for each sets of data treat each set of data as different studies



#### References



- Study Data Standards Resources
  - Providing Regulatory Submissions In Electronic Format Standardized Study Data: Guidance For Industry [June 2021]
  - FDA Data Standards Catalog [August 2022]
  - Study Data Technical Conformance Guide [March 2022]
  - Link: <a href="https://www.fda.gov/industry/fda-resources-data-standards/study-data-standards-resources">https://www.fda.gov/industry/fda-resources-data-standards/study-data-standards-resources</a>
- Study Data for Submission to CDER and CBER
  - Technical Rejection Criteria Self-Check Worksheet
  - Technical Rejection Criteria Self-Check Worksheet Instructions
  - Link: <a href="https://www.fda.gov/industry/study-data-standards-resources/study-data-submission-cder-and-cber">https://www.fda.gov/industry/study-data-standards-resources/study-data-submission-cder-and-cber</a>
- Providing Regulatory Submissions In Electronic Format Submissions Under Section 745a(a) Of The FD&C Act: Guidance For Industry
  - Link: <a href="https://www.fda.gov/regulatory-information/search-fda-guidance-documents">https://www.fda.gov/regulatory-information/search-fda-guidance-documents</a>