

Single Day Event

Implementation of the ICH E9(R1) Estimands Framework Using Data Standards

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This PHUSE project includes members from PHUSE collaborative partners and stakeholders and is currently ongoing, the final white paper will be shared for public review. Consequently, these recommendations are subject to change upon completion of the public review.

All examples contained within this presentation are examples of how to implement the E9(R1) estimands framework within data collection, tabulation and analysis following data standards. They should not be considered as examples of how to appropriately implement the estimands framework within an individual clinical study.



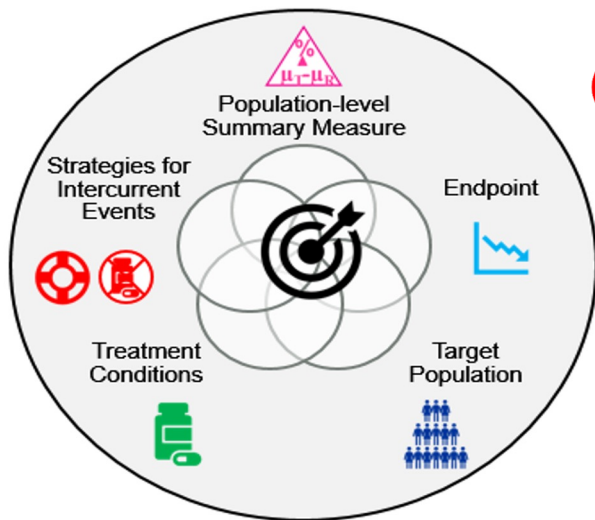
Introduction



What are Estimands?

Estimands

A precise description of the treatment effect reflecting the clinical question posed by the trial objective.
The estimand consists of 5 components:



Intercurrent Events

Events occurring after treatment initiation that affect either the interpretation or the existence of the measurements associated with the clinical question of interest



Trt discn
due to
Safety



Rescue Therapy



Trt discn
due to Lack
of Efficacy



Study Withdrawal



Death



Treatment
Switch

Analysis
Handling
Strategy



Treatment Policy

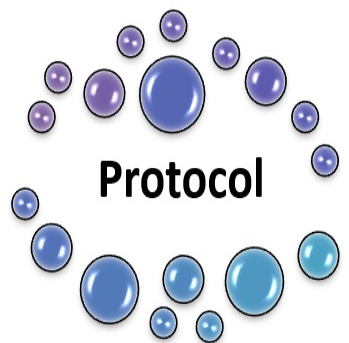
Composite

Hypothetical

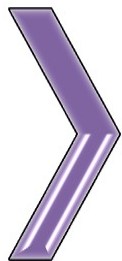
While on Treatment

Principal Stratum

Documentation to Data

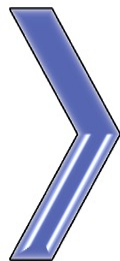


Describe the study objective in terms of the estimands framework (**WHAT**)



SAP

Statistical details on estimand (**WHAT detailed**), link estimands to their estimators (**HOW performed statistically**)

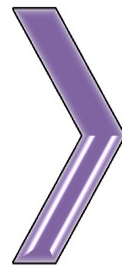


ADRG
Define.xml

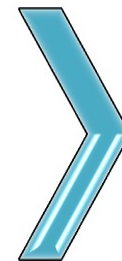
HOW the relevant aspects of estimands were **implemented** in the data. New section on estimands in CSDRG & ADRG



Get **sufficiently detailed answers** during collection



cSDRG



SDTM



Dedicated datasets and variables to document the traceability of estimands and impact in the data



Data Collection & Tabulation





Current Data Collection Practices

What was the subject's status?

- ☐ Progressive Disease (PROGRESSIVE DISEASE)
- ☐ Adverse Event (ADVERSE EVENT)
- ☐ Death (DEATH)
- ☐ Withdrawal by Subject (WITHDRAWAL BY SUBJECT)
- ☐ Physician Decision (PHYSICIAN DECISION)
- ☐ Non-Compliance With Study Drug (NON-COMPLIANCE WITH STUDY DRUG)
- ☐ Protocol Deviation (PROTOCOL DEVIATION)
- ☐ Study Terminated by IRB / ERB (STUDY TERMINATED BY IRB / ERB)
- ☐ Study Terminated by Sponsor (STUDY TERMINATED BY SPONSOR)
- ☐ Lost to follow up (LOST TO FOLLOW-UP)
- ☐ Pregnancy (PREGNANCY)

Need for Data Collection Enhancements

Accurate collection of intercurrent events is critical in defining estimands and constructing the estimators



Granular data collection of the reasons for treatment discontinuations



Data collection enhancements enable to use the most appropriate strategies to handle intercurrent events based on the underlying reasons



Commonly Observed Intercurrent Events



Intercurrent Events – Concomitant Medication

Suggested CRF for
Concomitant
Medication

Indicate if the subject took any concomitant medication/treatment. Record only one treatment per line. Provide full trade name of the medication/treatment Record specific reasons the medication was taken.	Were any concomitant medications taken? Not submitted	☐ Yes ☐ No
	What was the medication? CM.CMTRT	
	For what indication was the medication taken? CM.CMINDC	☐ ADVERSE EVENT. LINK TO ADVERSE EVENT: _____ ☐ MEDICAL HISTORY. LINK TO MEDICAL HISTORY: _____ ☐ CLINICAL EVENT. LINK TO CLINICAL EVENT: _____ ☐ PROPHYLAXIS FOR ANTIVIRAL ☐ PROPHYLAXIS FOR ANTIFUNGAL ☐ PROPHYLAXIS FOR INFECTION ☐ PROPHYLAXIS FOR INFUSION REACTION ☐ THROMBOPROPHYLAXIS ☐ PROPHYLAXIS FOR TUMOR LYSIS SYNDROME ☐ PROPHYLAXIS FOR COVID-19 ☐ VACCINATIONS ☐ REQUIRED CONCOMITANT MEDICATION FOR THE STUDY ☐ <STUDY INDICATION> ☐ RESCUE THERAPY ☐ BRIDGING THERAPY ☐ NON-THERAPEUTIC USE ☐ SUPPORTIVE CARE ☐ DIETARY SUPPLEMENT
	Start Date CM.CMSTDTC	
	Is the medication ongoing? CM.CMENRF or CMENRPT	☐ Yes
End date CM.CMENDTC		

SDTM Mapping
CM Domain

Row	STUDYID	DOMAIN	USUBJID	CMTRT	CMINDC	CMSTDTC	CMENRF	CMENDTC
1	ABC456	CM	456101	ASPIRIN	PROPHYLAXIS FOR INFECTION	2003-07-21	ONGOING	
2	ABC456	CM	456103	ANTACIDS	RESCUE THERAPY	2003-08-15		2003-09-01



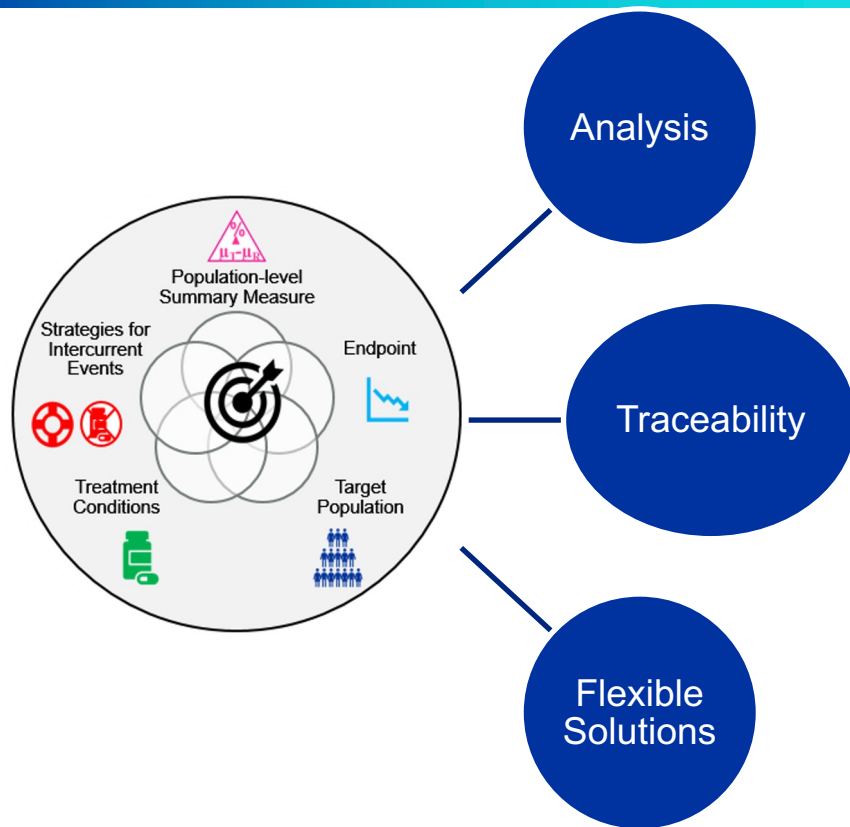
Data Collection & Tabulation - Summary



Data Analysis



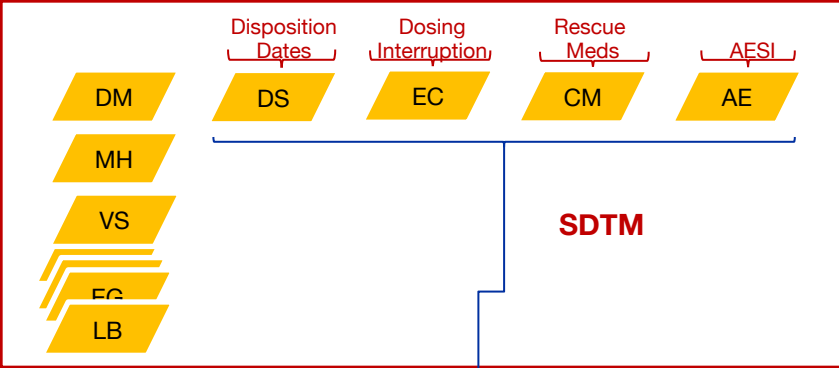
Estimands Impact on Analysis



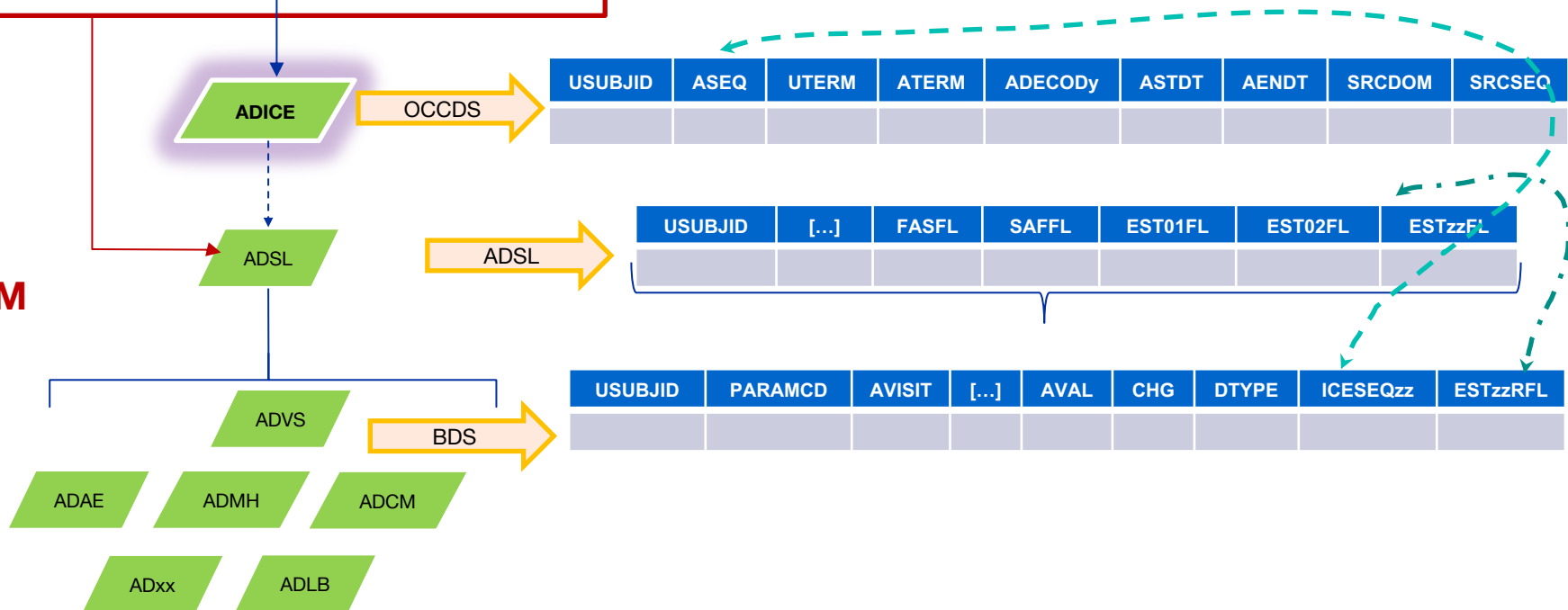
- Mapping intercurrent events
- Identifying subjects and data points for estimand-based analyses
- Enhanced ADaM dataset guidance is needed

- Documentation
- Estimands description and implementation

- Based on user needs
- Proposed examples will be offered in white paper



ADaM





NEW Intercurrent Events Dataset (ADICE)

- Documents intercurrent events across all estimands
- Facilitates traceability and inclusion of intercurrent events into other datasets
- OCCDS structure (one record per intercurrent event)
- This is an ***optional*** and supportive dataset to consolidate all intercurrent events in one place

USUBJID	ASEQ	ATERM	ADECODY	ASTDT(M)	AENDT(M)	SRCDOM	SRCSEQ

- Optional columns per estimand:
 - **ESTzzSTR**: Strategy (e.g., treatment policy) for handling the intercurrent event for estimand zz
 - **ESzzGRID**: Group multiple intercurrent events affecting a datapoint for estimand zz



NEW ADaM Dataset Variables

- ADSL (Subject-Level)

USUBJID	[...]	FASFL	SAFFL	EST01FL	EST02FL	ESTzzFL

- **ESTzzFL**: Subjects considered in all estimand zz estimations

- BDS (Basic Data Structure)

USUBJID	PARAMCD	AVISIT	[...]	AVAL	CHG	DTYPE	ICESEQzz	EST01RFL	EST02RFL	ESTzzRFL

- **ESTzzRFL**: Record-level datapoints considered in all estimand zz estimations
 - **ICESEQzz**: Links the intercurrent event(s) impacting the datapoint for estimand zz
 - Point to **ASEQ** of the single intercurrent event affecting the datapoint
 - Point to **ESzzGRID** of the multiple intercurrent events affecting the datapoint (advanced)
 - Note: if ADICE is not implemented: **ICEDOMzz** and **ICEVARzz** link to SDTM source
 - Similar for OCCDS and ADaM OTHER structures



Example – Major Depressive Disorder (MDD)

Treatment conditions	Assignment to drug X vs placebo, at the selected dose and frequency of administration, regardless of treatment discontinuation and as if other pharmacological treatments for MDD were not available
Target population	Patients with a diagnosis of MDD in a current major depressive episode with at least moderate symptom severity
Endpoint	Change from baseline to Week 8 in the total score of the 17-item version of Hamilton Depression Rating Scale (HDRS)
Population-level summary	Difference in treatment means



Example - Subject Journeys



Subject ID	BASELINE / WEEK 0	WEEK 1	WEEK 2	Unsch WEEK 2.1	WEEK 3	WEEK 4	WEEK 5	WEEK 6	WEEK 7	WEEK 8
ABC001	R/TI						SO x any			
ABC002	R/TI		TD (xAE/LoE)	SO x any						
ABC003	R/TI				SO x any			TD (xAE/LoE)		
ABC004	R/TI									

Outcome collected

No intercurrent event, all outcomes collected

New' outcome data following Hypothetical strategy for addressing the intercurrent event

R/TI= Randomisation/Treatment initiation

Intercurrent Event of interest:

TD (xAE/LoE): Treatment discontinuation (TD, due to any reason)

SO x any: Start other pharmacological treatments for MDD (SO)



Example - ADICE

USUBJID	ASEQ	ATERM	ADECOD1	ACAT1	ASTDT	AENDT	EST01STR	EST03STR	SRCDOM	SRCSEQ
ABC001	1	DULOXETINE	Starting other pharmacological treatments	Starting other pharmacological treatments	10APR2022		Hypothetical		CM	1
ABC002	1	ADVERSE EVENT	Treatment discontinuation due to adverse events	Treatment discontinuation	30MAR2022		Treatment Policy	Composite	DS	3
ABC002	2	FLUOXETINE	Starting other pharmacological treatments	Starting other pharmacological treatments	03APR2022		Treatment Policy		CM	1
ABC003	1	LORAZEPAM	Starting other pharmacological treatments	Starting other pharmacological treatments	06APR2022	20APR2022	Treatment Policy		CM	1
ABC003	2	LACK OF EFFICACY	Treatment discontinuation due to lack of efficacy	Treatment discontinuation	27APR2022		Treatment Policy	Composite	DS	3



Example - ADHDRS

USUBJID	PARAMCD	ADT	AVISIT	AVAL	BASE	CHG	ICESEQ01	CESEQ03	ANL01FL	EST01RF	EST03RFL
ABC001	HDRS	16MAR2022	BASELINE	25	25				Y	Y	Y
ABC001	HDRS	23MAR2022	WEEK 1	26	25	1			Y	Y	Y
ABC001	HDRS	30MAR2022	WEEK 2	24	25	-1			Y	Y	Y
ABC001	HDRS	06JUN2022	WEEK 3	23	25	-2			Y	Y	Y
ABC001	HDRS	13APR2022	WEEK 4	19	25	-6			Y	Y	Y
ABC001	HDRS	20APR2022	WEEK 5	18	25	-7	1		Y		Y
ABC001	HDRS	27APR2022	WEEK 6	23	25	-2			Y		Y
ABC001	HDRS	04MAY2022	WEEK 7	21	25	-4			Y		Y
ABC001	HDRS	25MAY2022	WEEK 8	16	25	-9			Y		Y
ABC002	HDRS	16MAR2022	BASELINE	31	31				Y	Y	Y
ABC002	HDRS	23MAR2022	WEEK 1	31	31	0			Y	Y	Y
ABC002	HDRS	30MAR2022	WEEK 2	30	31	-1	1	1	Y	Y	
ABC002	HDRS	03APR2022	WEEK 2 UNSCHEDULED	16	31	-15	2			Y	
ABC002	HDRS	06JUN2022	WEEK 3	30	31	-1			Y	Y	
ABC002	HDRS	13APR2022	WEEK 4	32	31	1			Y	Y	
ABC002	HDRS	20APR2022	WEEK 5	30	31	-1			Y	Y	
ABC002	HDRS	27APR2022	WEEK 6	34	31	3			Y	Y	
ABC002	HDRS	04MAY2022	WEEK 7	29	31	-2			Y	Y	



Data Analysis - Summary



Conclusion & Next Steps





Conclusion

Cross-functional
interaction is critical

Impacts protocol,
data collection and
data analysis

Different
implementation
approaches may be
appropriate

Need to
update/extend
existing data
standards

Consistent
implementation of
estimands is
beneficial



How Can You Help?

White paper now
available for
public review

Share with
colleagues

Provide us
feedback

Final Paper due
Q1 24

Share in your
organizations

Implement

Contact Information

Email: workinggroups@phuse.global

PHUSE Advance Hub:

<https://advance.phuse.global/display/WEL/Implementation+of+Estimands+%28ICH+E9+%28R1%29%29+using+Data+Standards>



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