## Checklist for Initial Data Analysis (IDA) – longitudinal studies

Lusa L, Proust-Lima C, Schmidt CO, Lee KJ, le Cessie S, Baillie M, Lawrence F, Huebner M. Initial data analysis for longitudinal studies to build a solid foundation for reproducible analysis. PLoS ONE 2024, to appear. https://stratosida.github.io/

Торіс	Item	Features		
IDA screening domain	: Partic			
Time frame	P1	Provide number of time points and intervals at which measurements are taken,		
		using the time metric that best reflects the time of inclusion in the study		
		(typically time from enrollment, or calendar time in studies that involve long		
		enrollment times). Highlight the differences between the time of first		
		measurements and follow-up times.		
Time metric	P2	Describe the time metric and corresponding time points specified in the analysis		
D (: : )	D2	strategy, if different from the time metric described in P1.		
Participants	P3	Provide the number of participants who attended the assessment by time		
Entensiones Destisinatio	n Dusfile	metric(s).		
Extensions: Participatio Other time metrics				
	PE1	Use different time metric(s) to describe the time frame of the study, if applicable and appropriate, e.g. calendar time or measurement occasion.		
Data collection	PE2	Describe aspects of the data collection process that can have an impact on the		
		data, if applicable. For example, describe if baseline and longitudinal		
		measurements are different, possible changes in variable measurements though		
		time, etc.		
<b>IDA screening domain</b> Non-enrollment				
Non-enronment	M1	Describe the non-enrolled, i.e., the participants that were selected but did not enter the study (and the reasons, if available), if applicable.		
Drop-out	M2	Describe the participants who dropped out from the study during the follow-up		
Drop out	1012	(loss to follow-up and other possible reasons: death, withdrawal, missing by		
		design, if applicable).		
Intermittent visit	M3	Describe the participants that have missing data for some of the measurements		
missingness		(intermittent, occasional omission, but do not drop out out of the study).		
Variable (item)	M4	Provide the number and proportion of missing values for each variable at each		
missingness		time point as appropriate for fixed or time-varying variables. Describe		
		missingness stratifying the summaries by variables that might influence the		
		frequency of missing values, if relevant (for example: structural variables or		
D	2.65	levels of measurement).		
Patterns	M5	Describe patterns of missing values across variables at each time point and		
Entensiones Missing dat		across time points.		
Extensions: Missing dat Non-enrollment	ME1	Compare the characteristics of the participants that entered the study with those		
Non-emonnent	IVIL I	of the non-enrolled or with the characteristics of the target population, if		
		applicable and data are available.		
Probability of drop-	ME2	Estimate the probability of drop-out after inclusion, taking appropriately into		
out	111111	account the reasons for drop-out.		
Dropout effect on	ME3	Visualize mean profiles of a continuous outcome by time metric stratified by		
outcome	_	time to drop-out.		
Predictors of	ME4	Explore whether there are predictors of missingness by comparing complete vs		
missingness		incomplete cases or investigate predictors of time to dropout, as appropriate;		
		this can assist in understanding of the missing data mechanism.		
IDA screening domain: Univariate descriptions				
Description of the	U1	Summarize the outcome variable and the explanatory variables with numerical		
variables at baseline		and graphical summaries at baseline.		

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Description of the	U2	Summarize the outcome variable and the time-varying explanatory variables		
time-varying variables		also at later time points. This might require discretization of time intervals		
at later points		and/or the use of different time metrics.		
IDA screening domain: Multivariate descriptions				
Association at	V1	Visualize the association between each explanatory variable with the structural		
baseline		variables at baseline.		
Correlation at baseline	V2	Quantify association with pairwise correlation coefficients between all		
		explanatory variables in a matrix or heatmap at baseline.		
Interactions at	V3	Evaluate bivariate distributions of the variables specified in the analysis strategy		
baseline, if applicable		with an interaction term; include appropriate graphical displays.		
Extensions: Multivariate descriptions				
Stratification	VE1	Compute summary statistics and describe variation between strata defined based		
		on level of measurement, e.g. centers, providers, locations, or by structural		
		variables or other variables described as stratification variables in the analysis		
		strategy (at baseline, other time points/time intervals can be also included).		
Associations and	VE2	Associations and correlations between explanatory variables at time points later		
correlations at time-		than baseline to explore their possible change across time; his could be useful		
points beyond baseline		for the identification of auxiliary variables. Selection might bias the results.		
Sampling design	VE3	If relevant, identify the stratifying variables and/or the clusters used in the		
		sampling design; explore the distribution of the number of clusters (by		
		stratification variables).		
IDA screening domain: Longitudinal aspects				
Profiles	L1	Summarize changes and variability of variables within subjects, e.g. profile		
		plots (spaghetti-plots) for groups of individuals.		
Trends	L2	Describe numerically or graphically longitudinal (average) trends of the outcome		
1101100		variable.		
Correlation and	L3	Estimate the strength of the within-participant correlation of the outcome		
variability	20	variable between time points and its variability across time points.		
Trends of time-	L4	Describe numerically or graphically the longitudinal trends of the time-varying		
varying explanatory		variables.		
variables				
Extensions: Longitudina	laspect			
Cohort/Period effects	LE1	If appropriate, summarize possible cohorts or period effects (for example, age		
		birth cohorts or period cohorts defined by the calendar time/wave of		
		measurement) on the outcome, and on the explanatory variables, to assess if the		
		variation of the outcome can occur because of these effects.		
		variation of the outcome can occur occause of these effects.		