Archives of Scientific Psychology Reporting Questionnaire for Manuscripts Describing Primary Data Collections

JARS: ALL: These questions should be answered for all submitted manuscripts

MANUSCRIPT SECTION	Description
TITLE	Does the Title identify the variables and theoretical issues under investigation, as well as the relationship between them? Yes ■ No □ If no, please explain:
AUTHOR NOTE	Does the Author Note contain acknowledgment of special circumstances, for example:
For a review of what should be included in the Author Note, see the <i>Publication Manual of the American Psychological Association</i> : http://www.apastyle.org/manual/	use of data also appearing in previous publications, dissertations, conference papers? Yes No If yes, please explain:

sources of funding or other support? Yes □ No If yes, please explain:
 relationships that may be perceived as conflicts of interest? Yes □ No ■
If yes, please explain:

SCIENTIFIC ABSTRACT	Does the Scientific Abstract describe:
	the problem under investigation?
	Yes ■ No Ø
	If no, please explain:
	 participants or subjects, specifying pertinent characteristics; in animal research, including genus and species?
	Yes ■ No □
	If no, please explain:

 study method, including: sample size? Yes ■ No □ any apparatus used? Yes ■ No □ measures? Yes ■ No □ data-gathering procedures? Yes ■ No □ research design (e.g., experiment, observational study)? Yes ■ No □
If answered "no" for any of the study methods above, please explain:
 findings, including effect sizes and confidence intervals and/or statistical significance levels? Yes ■ No □

If no, please explain:
 conclusions and the implications or applications? Yes ■ No □
If no, please explain:

INTRODUCTION	Does the Introduction:
For the Introduction please indicate whether the requested information can be found in this section of the manuscript, in	describe the importance of the problem?
a supplemental file, or whether the information is not relevant to the study. If the information is not relevant,	In manuscript ■ In supplemental files □ Not relevant □
please provide a brief explanation.	If not relevant, please explain:
	 describe theoretical or practical implications of the problem?
	In manuscript ■ In supplemental files □ Not relevant □
	If not relevant, please explain:

•	review relevant scho	olarship in relation to previous	s work?
	In manuscript	In supplemental files \square	Not relevant □
	If not relevant, pleas	se explain:	
•	review if other aspe	cts of this study have been re	ported upon previously and
	how the current rep	cts of this study have been re ort differs from these earlier re	eports?
	In manuscript	In supplemental files \square	Not relevant □
	If not relevant, pleas	se explain:	

theories or o	c hypotheses or objectives, su	
offered? In manuscript ≡	In supplemental files □	Not relevant □
If not relevant, pleas	se explain:	
o primary hyp	otheses?	
In manuscript 🖬	In supplemental files □	Not relevant □
If not relevant, pleas	se explain:	

o secondary h	nvpotheses?	
In manuscript	In supplemental files □	Not relevant □
		NOT relevant L
If not relevant, pleas	e explain:	
o planned exp	oloratory analyses?	
In manuscript 🗏	In supplemental files □	Not relevant □
If not relevant, pleas	se explain:	
		· ·

•	describe how hypoth	neses and research design re	elate to one another?
	In manuscript	In supplemental files \square	Not relevant □
	If not relevant, pleas	se explain:	

METHOD	For the Method section, please provide the information requested below, regardless of whether it also appears in the rest of the manuscript or in supplemental files.
Participant or subject characteristics:	What were the eligibility and exclusion criteria for participants or subjects, including any restrictions based on demographic characteristics?
	What were the major demographic characteristics of participants or subjects as well as important topic-specific characteristics, or, in the case of animal research, the genus and species?
Sampling procedures:	What procedures were used for selecting participants, including
	o the sampling method

 the percentage of sample approached that participated % any self-selection, either by individuals or by nomination from others?
What were the settings and locations where data were collected?
Were any agreements and payments made to participants?
 Were IRB agreements obtained, ethical standards met, and safety monitored? Yes ■ No ☑ If no, please explain:

Sample size, power and precision:	What was the intended sample size? n =
	What was the actual sample size? n=
	How was sample size determined:
	o power analysis? Yes ☑ No ■
	 o other methods used to determine accuracy of parameter estimates? Yes □ No ■
	If yes, describe:
	 stopping rules or interim analyses? Yes □ No ■
	If yes, describe:

Measures and covariates:	Please provide the defin including measures	itions of all primary and sec collected but not included i		ariates taken in the study,
	Measure name:		Definition:	
	What methods were use	d to collect data?		
	Were methods used to e training and reli Yes ■ No □ use of multiple o Yes ■ No □	ability of data collectors?	urements?	
	What are the known psy	/chometric and biometric pr	operties of instruments use	ed in the study?
	Measure Name:	Property:	Result:	

Research design:	
	Were conditions manipulated ■or naturalistic □?
	If manipulated, please complete JARS:EXP (see below)
	If manipulated, were subjects randomly assigned to conditions? Yes ■ No ☑
	If randomly assigned, please complete JARS: RCT (see below)
	If not randomly assigned, please complete JARS:QED (see below)
Miscellaneous:	
	Are there any other aspects of the study's methods that are important for the interpretation or replication of its findings?

RESULTS		please provide the information is or supplemental file in which the	requested in the questionnaire or provide e information can be found.
	If your manuscript is account of the data repository. Pleases www.apa.org/pubs/journ	see Instructions to Authors for m	eed to deposit your data set in an approved nore information:
Participant flow:		nove through each stage of the stud ort, if appropriate— <u>see Figure 1 bel</u> a	dy and how many were lost at each stage, if ow for an example)?
Recruitment:	Please provide the da	ites defining the periods of recruitm	ent and repeated measures or follow-up.
	Period Recruitment:	Start Date:	End Date:
 Missing data:	Did you experience proaffect the validity of fin Yes □ No■		nptions and/or data distributions that could
	If yes, please describe	::	

Missing data
Is missing data a cause of concern in this data set?
Yes⊡ No■
 If missing data was a cause of concern, is there empirical evidence and/or theoretical arguments
for the causes of data that are missing (e.g., missing completely at random (MCAR), missing at
random (MAR), or missing not at random (MNAR))?
 If missing data was a cause of concern, is there empirical evidence and/or theoretical arguments for the causes of data that are missing (for example, missing completely at random (MCAR),
missing at random (MAR), or missing not at random (MNAR))?

If missing data was a cause of concern, what methods, if any, were used for addressing missing data?
For the Discussion section, please indicate whether the requested information can be found in this
section of the manuscript, in a supplemental file, or whether the information is not relevant to the study. If not relevant, please provide a brief explanation.
 Did you experience problems concerning statistical assumptions and/or data distributions that could affect the validity of findings?
Yes☑ No■
If yes, please describe:
For inferential statistics (NHST), please indicate the a priori Type 1 error rate adopted:

For each NHST conducted, regardless of whether significant results were obtained and regardless of whether or not reported in the text, please provide a log of the centrality (primary, secondary exploratory) of the analyses to the study's purpose, the analytic technique used, the direction, magnitude, degrees of freedom, and exact p-level associated with each test: Provided Test
For multivariable analytic systems (e.g., multivariate analyses of variance, regression analyses, structural equation modeling analyses, and hierarchical linear modeling)
provide the associated variance-covariance (or correlation) matrix or matrices:
describe any estimation problems (e.g., failure to converge, bad solution spaces), anomalous data points:
identify the statistical software program, if specialized procedures were used:
- identify the statistical contraint program, in openialized procedures were deed.

Is there a statement secondary hypotherical secondary hypotheri		Il original hypotheses distinguished by primary and
In manuscript	In supplemental files \square	Not relevant □
If not relevant, pl	ease explain:	
Are post hoc exp	lanations proposed?	
In manuscript	In supplemental files \square	Not relevant □
If not relevant, ple	ease explain:	
Are the similaritie	es and differences between these	results and the work of others discussed?
In manuscript	In supplemental files \square	Not relevant □
If not relevant, please	e explain:	

 Are results interprete sources of potent 	d taking into account	nal validity?
In manuscript ■ If not relevant, please	In supplemental files □ e explain:	Not relevant □
 imprecision of me In manuscript ■ 	easures? In supplemental files □	Not relevant □
If not relevant, please		
the overall number	er of tests or overlap among tests	s?
In manuscript ■ If not relevant, please	In supplemental files □	Not relevant □
ii not relevant, pleast	о одрішії.	

other limitations	or weaknesses of the study?	
In manuscript 🖃	In supplemental files \square	Not relevant □
If not relevant, pleas	se explain:	
Is the generalizability	y (external validity) of the finding	gs taken into account with regard to
 the target popula 	ation?	
In manuscript 🚍	In supplemental files \square	Not relevant □
If not relevant, pleas	se explain:	
other contextual	issues?	
In manuscript 🗏	In supplemental files \square	Not relevant □

If not relevant, please explain:
Is there discussion of implications for future research, program, or policy In manuscript ■ In supplemental files □ Not relevant □
If not relevant, please explain:

JARS: EXP: These questions should be answered for all studies with an experimental manipulation or intervention (in addition to the JARS: ALL Questionnaire)

METHODS	In the Method section of a study with an experimental manipulation or intervention, please provide the information requested below, regardless of whether it also appears in the manuscript or a supplemental file. If the information requested is irrelevant to the study, briefly explain why.
Experimental manipulations or interventions:	 Please provide the details about the experimental manipulations or interventions intended for each study condition, including control groups and specifically including the content of the specific experimental manipulations or interventions—a summary or paraphrasing of instructions (unless they are unusual or compose the manipulation, in which case they may be presented verbatim):
	the method of manipulation or intervention delivery—a description of apparatus and materials used and their function in the experiment:
	Identify specialized equipment by model and supplier:
	the deliverers, that is, who delivered the manipulations or interventions level of professional training:

level of training in specific manipulations or interventions:
 the number of deliverers and, in the case of interventions, the M, SD, and range of number of individuals/units treated by each:
the setting , that is, where the manipulations or interventions occurred:
 the exposure quantity and duration, that is, how many sessions, episodes, or events were intended to be delivered and how long they were intended to last:
the time span , that is, how long it took to deliver the intervention or manipulation to each unit:

	activities to increase compliance or adherence (e.g. incentives):
	the use of languages other than English and the translation method:
	and use of languages outer than English and the translation method.
Modrings	
Masking:	Were participants, those administering the interventions, and those assessing the outcomes unaware of
	condition assignments?
	Yes■ No 🖸
	If no, why not?
	If masking took place, how was it accomplished, and how was its success evaluated?

 Units of delivery and analysis:	Unit of delivery: How were participants grouped during delivery?
	 What was the smallest unit that was analyzed (and, in the case of experiments, that was randomly assigned to conditions) to assess manipulation or intervention effects (e.g., individuals, work groups, classes)?
	If the unit of analysis differed from the unit of delivery, please describe the analytical method used to account for this (e.g., adjusting the standard error estimates by the design effect or using multilevel analysis):

RESULTS	For the Results section, please indicate below the page number, table, or supplemental file in which the information can be found.
Participant flow:	What was the total number of groups (if the experimental manipulation or intervention was administered at the group level), and what was the number of participants assigned to each group?
Treatment fidelity:	What evidence is there that the deliverers of treatment adhered to the respective intervention manuals/guidelines?
	What evidence is there that the treatments were delivered competently?

Statistics and data analysis:	Were the analyses intent-to-treat□, complier average causal effect≡, or other or multiple ways□?
	Please explain:
Adverse events and side effects:	
	Please describe all important adverse events or side effects in each experimental or intervention:

DISCUSSION	For the Discussion section, please indicate below the page number, table, or supplemental file in which the information can be found.
	Do results discussed take into account the mechanism by which the manipulation or intervention was intended to work (causal pathways) or alternative mechanisms?
	Yes■ No⊠
	If no, please explain:
	 If an intervention is involved, is there discussion of the success of and barriers to implementing the intervention, and the fidelity of implementation?
	Yes ☑ No■
	If no, please explain:
	Is there a discussion of the generalizability (external validity) of the findings taking into account
	 the characteristics of the intervention?
	Yes■ No©

If no, please explain:
o how and what outcomes were measured?
Yes■ No⊡
If no, please explain:
o length of follow-up?
Yes☑ No■
 If no, please explain:
o incentives?
Yes⊠ No■
If we who are combate.
If no, please explain:

o compliance rates?
Yes⊠ No∎
If no, please explain:
 Is there discussion of the clinical or practical significance of outcomes and the basis for these interpretations?
Yes■ No⊠
If no, please explain:

JARS: RCT: These questions should be answered for all studies with an experimental manipulation or intervention that employed random assignment to experimental conditions (in addition to JAR:ALL and JARS: EXP)

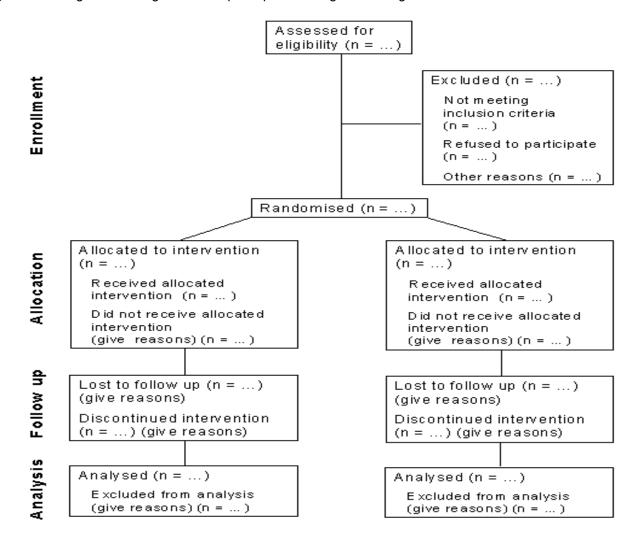
METHOD	In the Method section of a study that employed random assignment to experimental conditions, please provide the information requested below, regardless of whether it also appears in the manuscript or a supplemental file. If the information requested is irrelevant to the study, briefly explain why.
	What procedures were used to generate the random assignment sequence (including details of any restrictions—e.g., blocking, stratification)?
 Random assignment – concealment:	Was the sequence concealed until experimental or intervention sequence was assigned? Yes No No If no, why not?

Random assignment – implementation:	Who generated the assignment sequence?
	Who enrolled participants?
	Who assigned participants to groups?

JARS: QED: These questions should be answered for all studies with an experimental manipulation or intervention that did not employ random assignment to experimental conditions (in addition to JARS: All and JARS: EXP).

METHOD	
Assignment method:	What was the unit of assignment (the unit being assigned to study conditions—e.g., individual, group, community)?
	What was the method used to assign units to study conditions, including details of any restriction (e.g., blocking, stratification, minimization)?
	What procedures were employed to help minimize potential bias due to nonrandomization (e.g., matching, propensity score matching)?

Figure 1. Diagram showing the flow of participants through each stage of a randomized trial.



intervention (in addition to JARS: All).
Please provide below as detailed a description as possible of the research design used in the study or studies. This description should be at least as detailed than that expected in all APA journals. There is no restriction on length.