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 \(\subseteq \text{No} \subseteq \) If yes, please explain:
	п уез, рієдзе ехрідіп.

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 □
	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
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	s the Introduction:		
For the Introduction please indicate whether the requested information can be found in this section of the manuscript, in		describe the importa	nnce of the problem?	
a supplemental file, or whether the information is not relevant to the study. If the information is not relevant,		n manuscript □	In supplemental files \square	Not relevant □
please provide a brief explanation.	<u> </u>	lf not relevant, pleas	se explain:	
	• (describe theoretical	or practical implications of th	e problem?
		n manuscript □	In supplemental files \square	Not relevant □
		f not relevant, pleas	e explain:	
	1			

•	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 how the current rep	cts of this study have been re ort differs from these earlier re	ported upon previously and eports?
	In manuscript □	In supplemental files \square	Not relevant □
	If not relevant, pleas	se explain:	
<u> </u>			

,	
	 describe the specific hypotheses or objectives, such as theories or other means to derive hypotheses, if hypotheses were offered?
	In manuscript \square In supplemental files \square Not relevant \square
	If not relevant, please explain:
	o primary hypotheses?
	In manuscript \square In supplemental files \square Not relevant \square
	If not relevant, please explain:

·			
	○ secondary h	ypotheses?	
	In manuscript □	In supplemental files \square	Not relevant □
	If not relevant, pleas	e explain:	
	o planned exp	loratory analyses?	
	In manuscript □	In supplemental files \square	Not relevant \square
	If not relevant, pleas	e explain:	

describe how hypotheses and research design relate to one another?
In manuscript □ In supplemental files □ Not relevant □
If not relevant, please 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 □
	\circ other methods used to determine accuracy of parameter estimates? Yes \square No \square
	If yes, describe:
	o stopping rules or interim analyses?Yes □ No □
	If yes, describe:

Measures and covariates:		of all primary and secondary measucted but not included in this report	ures and covariates taken in the study,
	Measure name:	Definition:	
	What methods were used to contain the second contains the sec	collect data?	
	Were methods used to enhan	ce the quality of measurements?	
	o training and reliability Yes □ No □	of data collectors?	
	o use of multiple obser Yes □ No □	vations?	
	What are the known psychon	netric and biometric properties of ins	struments used 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 \square No \square
	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 or supplemental file in which the	requested in the questionnaire or provide e information can be found.
		see Instructions to Authors for n	need to deposit your data set in an approved nore information:
Participant flow:		nove through each stage of the stud rt, if appropriate— <u>see Figure 1 bel</u>	dy and how many were lost at each stage, if ow for an example)?
 Recruitment:	Please provide the date	tes defining the periods of recruitm	nent and repeated measures or follow-up.
	Period Recruitment:	Start Date:	End Date:
Missing data:	Did you experience proaffect the validity of fin Yes No		mptions 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?
DISCUSSION	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.
Statistics and data analysis:	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 of secondary hypothese		original hypotheses distinguished by primary and
In manuscript \square	In supplemental files \square	Not relevant □
If not relevant, please	e explain:	
Are post hoc explanation in manuscript	In supplemental files □	Not relevant □
If not relevant, please	ехріаіп:	
Are the similarities an	d differences between these re	sults and the work of others discussed?
In manuscript \square	In supplemental files \square	Not relevant □
If not relevant, please exp	olain:	

 Are results interpreted taking into account sources of potential bias and other threats to internal validity?
In manuscript □ In supplemental files □ Not relevant □ If not relevant, please explain:
 imprecision of measures? In manuscript □ In supplemental files □ Not relevant □ If not relevant, please explain:
the overall number of tests or overlap among tests? In manuscript □ In supplemental files □ Not relevant □
If not relevant, please explain:

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		•
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:
 Masking:	
	 Were participants, those administering the interventions, and those assessing the outcomes unaware of condition assignments? Yes□ No□
	If no, why not?
	If marking took place, how was it accomplished, and how was its success evaluated?
	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:
ii iio, picase 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 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 \(\text{No} \)
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)

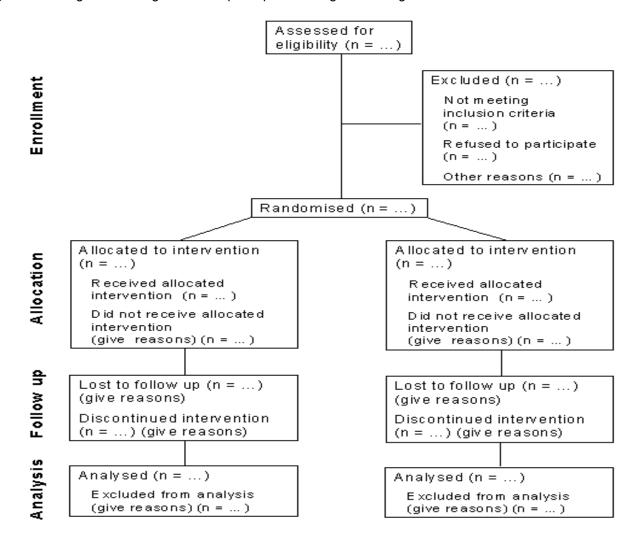
METHOD	In the Method section of a study that employed random assignment to experimental conditions, please provide the information requested below, regardless of whether it also appears in the manuscript or a supplemental file. If the information requested is irrelevant to the study, briefly explain 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□ 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.