## **TREND Statement Checklist**

Paper Section/ Topic	Item	Descriptor	Reported?	
	No		$\checkmark$	Pg#
Title and Abst	ract			
Title and	1	Information on how unit were allocated to interventions	Yes	3
Abstract		Structured abstract recommended		3
		Information on target population or study sample	Yes	3
Introduction	· II			•
Introduction Background	2	Scientific background and explanation of rationale	Yes	4
Васкугоина			N/A	
		Theories used in designing behavioral interventions	14//	
Methods				
Participants	3	Eligibility criteria for participants, including criteria at different levels in	Yes	6-7
		recruitment/sampling plan (e.g., cities, clinics, subjects)		
		Method of recruitment (e.g., referral, self-selection), including the	Yes	7
		sampling method if a systematic sampling plan was implemented		
		Recruitment setting	N/A	
		Settings and locations where the data were collected	Yes	Supp1:
Interventions	4	Details of the interventions intended for each study condition and how	Yes	6
		and when they were actually administered, specifically including:		
		Content: what was given?	Yes	6
		<ul> <li>Delivery method: how was the content given?</li> </ul>	Yes	6
		<ul> <li>Unit of delivery: how were the subjects grouped during delivery?</li> </ul>	N/A	
		Deliverer: who delivered the intervention?	Yes	6
		<ul> <li>Setting: where was the intervention delivered?</li> </ul>	Yes	6
		<ul> <li>Exposure quantity and duration: how many sessions or episodes or</li> </ul>	Yes	9
		events were intended to be delivered? How long were they intended to last?		
		Time span: how long was it intended to take to deliver the	N/A	
		intervention to each unit?		
		Activities to increase compliance or adherence (e.g., incentives)	N/A	·
Objectives	5	Specific objectives and hypotheses	Yes	6
Outcomes	6	Clearly defined primary and secondary outcome measures	Yes	7-8
		Methods used to collect data and any methods used to enhance the	Yes	7-8
		quality of measurements		
		Information on validated instruments such as psychometric and biometric	N/A	
		properties		
Sample Size	7	How sample size was determined and, when applicable, explanation of any	Yes	6
		interim analyses and stopping rules		
Assignment	8	<ul> <li>Unit of assignment (the unit being assigned to study condition, e.g.,</li> </ul>	Yes	6
Method		individual, group, community)		
		Method used to assign units to study conditions, including details of any	N/A	
		restriction (e.g., blocking, stratification, minimization)		
		Inclusion of aspects employed to help minimize potential bias induced due	N/A	
	1	to non-randomization (e.g., matching)		1

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Blinding	9	Whether or not participants, those administering the interventions, and	Yes	6
(masking)		those assessing the outcomes were blinded to study condition assignment; if so, statement regarding how the blinding was accomplished and how it was assessed.		
Unit of Analysis	10	Description of the smallest unit that is being analyzed to assess intervention effects (e.g., individual, group, or community)	Yes	6
		If the unit of analysis differs from the unit of assignment, the analytical method used to account for this (e.g., adjusting the standard error estimates by the design effect or using multilevel analysis)	N/A	
Statistical Methods	11	Statistical methods used to compare study groups for primary methods outcome(s), including complex methods of correlated data	N/A	
		Statistical methods used for additional analyses, such as a subgroup analyses and adjusted analysis	Yes	Supp10
		Methods for imputing missing data, if used	N/A	
		Statistical software or programs used	Yes	9
Results				
Participant flow	12	Flow of participants through each stage of the study: enrollment,	Yes	Fig 1
•		assignment, allocation, and intervention exposure, follow-up, analysis (a diagram is strongly recommended)	100	1 19 1
		<ul> <li>Enrollment: the numbers of participants screened for eligibility, found to be eligible or not eligible, declined to be enrolled, and enrolled in the study</li> </ul>	Yes	Fig 1
		<ul> <li>Assignment: the numbers of participants assigned to a study condition</li> </ul>	Yes	Fig 1
		<ul> <li>Allocation and intervention exposure: the number of participants assigned to each study condition and the number of participants who received each intervention</li> </ul>	Yes	Fig 1
		<ul> <li>Follow-up: the number of participants who completed the follow-up or did not complete the follow-up (i.e., lost to follow-up), by study condition</li> </ul>	Yes	Fig 1
		<ul> <li>Analysis: the number of participants included in or excluded from the main analysis, by study condition</li> </ul>	Yes	Fig 1
		Description of protocol deviations from study as planned, along with reasons	Yes	9
Recruitment	13	Dates defining the periods of recruitment and follow-up	Yes	6
Baseline Data	14	Baseline demographic and clinical characteristics of participants in each study condition	Yes	9-10
		Baseline characteristics for each study condition relevant to specific disease prevention research	N/A	
		Baseline comparisons of those lost to follow-up and those retained, overall and by study condition	N/A	
		Comparison between study population at baseline and target population of interest	Yes	10
Baseline equivalence	15	Data on study group equivalence at baseline and statistical methods used to control for baseline differences	N/A	

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Numbers analyzed	16	Number of participants (denominator) included in each analysis for each study condition, particularly when the denominators change for different outcomes; statement of the results in absolute numbers when feasible	Yes	9
		<ul> <li>Indication of whether the analysis strategy was "intention to treat" or, if not, description of how non-compliers were treated in the analyses</li> </ul>	Yes	9
Outcomes and estimation	17	For each primary and secondary outcome, a summary of results for each estimation study condition, and the estimated effect size and a confidence interval to indicate the precision	Yes	9-13
		Inclusion of null and negative findings	Yes	12
		<ul> <li>Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any</li> </ul>	N/A	
Ancillary analyses	18	Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory	Yes	Supp10
Adverse events	19	<ul> <li>Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals)</li> </ul>	Yes	12
DISCUSSION				
Interpretation	20	<ul> <li>Interpretation of the results, taking into account study hypotheses, sources of potential bias, imprecision of measures, multiplicative analyses, and other limitations or weaknesses of the study</li> </ul>	Yes	14
		Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations	Yes	15
		Discussion of the success of and barriers to implementing the intervention, fidelity of implementation	N/A	
		Discussion of research, programmatic, or policy implications	Yes	16
Generalizability	21	<ul> <li>Generalizability (external validity) of the trial findings, taking into account the study population, the characteristics of the intervention, length of follow-up, incentives, compliance rates, specific sites/settings involved in the study, and other contextual issues</li> </ul>	Yes	13-17
Overall Evidence	22	General interpretation of the results in the context of current evidence and current theory	Yes	13-17

*From:* Des Jarlais, D. C., Lyles, C., Crepaz, N., & the Trend Group (2004). Improving the reporting quality of nonrandomized evaluations of behavioral and public health interventions: The TREND statement. *American Journal of Public Health*, 94, 361-366. For more information, visit: <a href="http://www.cdc.gov/trendstatement/">http://www.cdc.gov/trendstatement/</a>