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		25
Abstract		Structured abstract recommended		23-36
		Information on target population or study sample		32
Introduction				
Background	2	Scientific background and explanation of rationale		38
		Theories used in designing behavioral interventions		62
Mathada				02
Methods Participants	3	Eligibility criteria for participants, including criteria at different levels in		
	_	recruitment/sampling plan (e.g., cities, clinics, subjects)		86
		Method of recruitment (e.g., referral, self-selection), including the		
		sampling method if a systematic sampling plan was implemented		102
		Recruitment setting		87
		Settings and locations where the data were collected		139
Interventions	4	• Details of the interventions intended for each study condition and how		
		and when they were actually administered, specifically including:		
		 Content: what was given? 		110
		 Delivery method: how was the content given? 		114
		 Unit of delivery: how were the subjects grouped during delivery? 		
		O Deliverer: who delivered the intervention?		100
		 Setting: where was the intervention delivered? 		123
		 Exposure quantity and duration: how many sessions or episodes or 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 		
		intervention to each unit?		135
		 Activities to increase compliance or adherence (e.g., incentives) 		
Objectives	5	Specific objectives and hypotheses		82
Outcomes	6	Clearly defined primary and secondary outcome measures		143
		 Methods used to collect data and any methods used to enhance the 		139
		quality of measurements		139
		• Information on validated instruments such as psychometric and biometric		
	_	properties		
Sample Size	7	• How sample size was determined and, when applicable, explanation of any		150
A	<u> </u>	interim analyses and stopping rules		
Assignment Method	8	 Unit of assignment (the unit being assigned to study condition, e.g., individual group, community) 		77
		individual, group, community)		
		 Method used to assign units to study conditions, including details of any restriction (e.g., blocking, stratification, minimization) 		
		 Inclusion of aspects employed to help minimize potential bias induced due 		
		to non-randomization (e.g., matching)		

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Blinding	9	Whether or not participants, those administering the interventions, and	
(masking)	5	those assessing the outcomes were blinded to study condition assignment;	
(masking)		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	139
		intervention effects (e.g., individual, group, or community)	107
		• 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	162
		estimates by the design effect or using multilevel analysis)	
Statistical	11	Statistical methods used to compare study groups for primary methods	150
Methods		outcome(s), including complex methods of correlated data	150
		Statistical methods used for additional analyses, such as a subgroup	
		analyses and adjusted analysis	
		Methods for imputing missing data, if used	
		Statistical software or programs used	167
Results			
Participant flow	12	Flow of participants through each stage of the study: enrollment,	
•		assignment, allocation, and intervention exposure, follow-up, analysis (a	174
		diagram is strongly recommended)	174
		• Enrollment: the numbers of participants screened for eligibility,	
		found to be eligible or not eligible, declined to be enrolled, and	170
		enrolled in the study	173
		 Assignment: the numbers of participants assigned to a study 	470
		condition	179
		 Allocation and intervention exposure: the number of participants 	
		assigned to each study condition and the number of participants	179
		who received each intervention	
		 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	173
		study condition	
		 Analysis: the number of participants included in or excluded from 	173
		the main analysis, by study condition	
		Description of protocol deviations from study as planned, along with	
		reasons	
Recruitment	13	Dates defining the periods of recruitment and follow-up	
Baseline Data	14	Baseline demographic and clinical characteristics of participants in each	173
		study condition	
		Baseline characteristics for each study condition relevant to specific	
		disease prevention research	
		Baseline comparisons of those lost to follow-up and those retained, overall and hu study as addition	
		and by study condition	
		 Comparison between study population at baseline and target population of interest 	
Baseline	15	 Data on study group equivalence at baseline and statistical methods used 	
equivalence	τı	• Data on study group equivalence at baseline and statistical methods used to control for baseline differences	
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Numbers	16	Number of participants (denominator) included in each analysis for each	
analyzed		study condition, particularly when the denominators change for different	
		outcomes; statement of the results in absolute numbers when feasible	
		• Indication of whether the analysis strategy was "intention to treat" or, if	
		not, description of how non-compliers were treated in the analyses	
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	158
		Inclusion of null and negative findings	
		Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any	
Ancillary	18	Summary of other analyses performed, including subgroup or restricted	200
analyses	10	analyses, indicating which are pre-specified or exploratory	
Adverse events	19	 Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals) 	
DISCUSSION			
Interpretation	20	• 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	279
		Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations	218
		• Discussion of the success of and barriers to implementing the intervention, fidelity of implementation	
		Discussion of research, programmatic, or policy implications	261
Generalizability	21	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	
Overall	22	General interpretation of the results in the context of current evidence	

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: <u>http://www.cdc.gov/trendstatement/</u>