

TREND Statement Checklist

| Paper Section/ Topic | Item No | Descriptor | Reported? | |
|---|---------|--|-----------|--|
| | | | √ | Section |
| Title and Abstract | | | | |
| Title and Abstract | 1 | • Information on how unit were allocated to interventions | √ | Abstract |
| | | • Structured abstract recommended | √ | |
| | | • Information on target population or study sample | √ | |
| Introduction | | | | |
| Background | 2 | • Scientific background and explanation of rationale | √ | Introduction |
| | | • Theories used in designing behavioral interventions | | Not applicable |
| Methods | | | | |
| Participants | 3 | • Eligibility criteria for participants, including criteria at different levels in recruitment/sampling plan (e.g., cities, clinics, subjects) | √ | Patients and Methods-Patients |
| | | • Method of recruitment (e.g., referral, self-selection), including the sampling method if a systematic sampling plan was implemented | √ | |
| | | • Recruitment setting | √ | |
| | | • Settings and locations where the data were collected | √ | |
| 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? | √ | Patients and Methods-Testing devices, Protocol |
| | | ○ Delivery method: how was the content given? | √ | Patients and Methods-Protocol |
| | | ○ Unit of delivery: how were the subjects grouped during delivery? | | Not applicable |
| | | ○ Deliverer: who delivered the intervention? | √ | Patients and Methods-Protocol |
| | | ○ Setting: where was the intervention delivered? | √ | |
| | | ○ 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? | | Not applicable |
| ○ Activities to increase compliance or adherence (e.g., incentives) | | Not applicable | | |
| Objectives | 5 | • Specific objectives and hypotheses | √ | Introduction |
| Outcomes | 6 | • Clearly defined primary and secondary outcome measures | | Not applicable |
| | | • Methods used to collect data and any methods used to enhance the quality of measurements | √ | Patients and Methods-Protocol |
| | | • Information on validated instruments such as psychometric and biometric properties | √ | Patients and Methods-Testing devices |
| Sample Size | 7 | • How sample size was determined and, when applicable, explanation of any interim analyses and stopping rules | | Not applicable |
| Assignment Method | 8 | • Unit of assignment (the unit being assigned to study condition, e.g., individual, group, community) | | Not applicable |
| | | • Method used to assign units to study conditions, including details of any restriction (e.g., blocking, stratification, minimization) | | Not applicable |
| | | • Inclusion of aspects employed to help minimize potential bias induced due to non-randomization (e.g., matching) | | Not applicable |

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| Blinding (masking) | 9 | <ul style="list-style-type: none"> Whether or not participants, those administering the interventions, and those assessing the outcomes were blinded to study condition assignment; if so, statement regarding how the blinding was accomplished and how it was assessed. | | Not applicable |
| Unit of Analysis | 10 | <ul style="list-style-type: none"> Description of the smallest unit that is being analyzed to assess intervention effects (e.g., individual, group, or community) | | Not applicable |
| | | <ul style="list-style-type: none"> 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) | | Not applicable |
| Statistical Methods | 11 | <ul style="list-style-type: none"> Statistical methods used to compare study groups for primary methods outcome(s), including complex methods of correlated data | √ | Patients and Methods-Statistical Analysis |
| | | <ul style="list-style-type: none"> Statistical methods used for additional analyses, such as a subgroup analyses and adjusted analysis | √ | |
| | | <ul style="list-style-type: none"> Methods for imputing missing data, if used | | Not applicable |
| | | <ul style="list-style-type: none"> Statistical software or programs used | √ | Patients and Methods-Statistical Analysis |
| Results | | | | |
| Participant flow | 12 | <ul style="list-style-type: none"> Flow of participants through each stage of the study: enrollment, assignment, allocation, and intervention exposure, follow-up, analysis (a diagram is strongly recommended) | | |
| | | <ul style="list-style-type: none"> Enrollment: the numbers of participants screened for eligibility, found to be eligible or not eligible, declined to be enrolled, and enrolled in the study | √ | Results-Clinical features |
| | | <ul style="list-style-type: none"> Assignment: the numbers of participants assigned to a study condition | √ | |
| | | <ul style="list-style-type: none"> Allocation and intervention exposure: the number of participants assigned to each study condition and the number of participants who received each intervention | | Not applicable |
| | | <ul style="list-style-type: none"> 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 | √ | Results-Clinical features |
| | | <ul style="list-style-type: none"> Analysis: the number of participants included in or excluded from the main analysis, by study condition | √ | |
| | | <ul style="list-style-type: none"> Description of protocol deviations from study as planned, along with reasons | √ | Results-Clinical features |
| Recruitment | 13 | <ul style="list-style-type: none"> Dates defining the periods of recruitment and follow-up | √ | Patients and Methods-Patients |
| Baseline Data | 14 | <ul style="list-style-type: none"> Baseline demographic and clinical characteristics of participants in each study condition | √ | Results-Clinical features |
| | | <ul style="list-style-type: none"> Baseline characteristics for each study condition relevant to specific disease prevention research | | Not applicable |
| | | <ul style="list-style-type: none"> Baseline comparisons of those lost to follow-up and those retained, overall and by study condition | | Not applicable |
| | | <ul style="list-style-type: none"> Comparison between study population at baseline and target population of interest | | Not applicable |
| Baseline equivalence | 15 | <ul style="list-style-type: none"> Data on study group equivalence at baseline and statistical methods used to control for baseline differences | | Not applicable |

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| Numbers analyzed | 16 | <ul style="list-style-type: none"> 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 | | Results |
| | | <ul style="list-style-type: none"> Indication of whether the analysis strategy was “intention to treat” or, if not, description of how non-compliers were treated in the analyses | | Not applicable |
| Outcomes and estimation | 17 | <ul style="list-style-type: none"> 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 | | Not applicable |
| | | <ul style="list-style-type: none"> Inclusion of null and negative findings | | Not applicable |
| | | <ul style="list-style-type: none"> Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any | | Not applicable |
| Ancillary analyses | 18 | <ul style="list-style-type: none"> Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory | √ | Results – all sections |
| Adverse events | 19 | <ul style="list-style-type: none"> Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals) | | Not applicable |
| Discussion | | | | |
| Interpretation | 20 | <ul style="list-style-type: none"> 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 | √ | Discussion |
| | | <ul style="list-style-type: none"> Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations | | Not applicable |
| | | <ul style="list-style-type: none"> Discussion of the success of and barriers to implementing the intervention, fidelity of implementation | √ | Discussion |
| | | <ul style="list-style-type: none"> Discussion of research, programmatic, or policy implications | √ | |
| Generalizability | 21 | <ul style="list-style-type: none"> 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 | √ | Discussion |
| Overall Evidence | 22 | <ul style="list-style-type: none"> General interpretation of the results in the context of current evidence and current theory | √ | Discussion |

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