## **S2 File. Study Quality Checklist**

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| Quality Assessment[[1]](#footnote-1) |  |
| **Study design score**[[2]](#footnote-2) |
| 1 | Study without a comparison group – No analysis of change |
| 2 | Inadequately controlled study – No analysis of change |
| 3 | Study without a comparison group – With analysis of change |
| 4 | Inadequately controlled study – With analysis of change |
| 5 | Controlled non-experimental study – No analysis of change |
| 6 | Controlled non-experimental study – With analysis of change |
| 7 | Randomised experiment targeting a risk factor |
| **Cluster sampling (cluster= facility, community, school or other)** |
| 3 | Total population or random sampling |
| 2 | Purposive sampling |
| 1 | Convenience sampling |
| 0 | Not reported |
| **Within-cluster sampling of participants** |
| 3 | Total population or random sampling |
| 2 | Purposive sampling |
| 1 | Convenience sampling |
| 0 | Not reported |
| **Response rates** |
| 2 | Response*or* retention rates ≥70% *or* differential attrition ≤10%[[3]](#footnote-3) |
| 1 | Response rate <70% *or* retention rate <70% *or* differential attrition >10% |
| 0 | Not reported |
| **Sample size score** |
| 1 | Sample size ≥400 (cohort studies) and ≥200 (experimental designs) |
| 0 | Sample size <400 (cohort studies) and <200 (experimental designs) |
| **Outcome measures (***complete for each different outcome***)** |
| 4 | Self-reported measure validated through biomarker |
| 3 | Self-reported measure validated through other measurements |
| 2 | Self-reported measure collected through special techniques to improve response rate (e.g. AUDIO-ACASI/ AMASI, ACASI/ AMASI, sealed envelopes, etc.) |
| 1 | Self-reported measure collected through regular techniques |
| 0 | Not reported |
| **Determinant measures (***complete for each different determinant***)** |
| 3 | Reliability coefficient ≥.75 and reasonable face validity *or* criterion or convergent validity coefficient ≥ 0.3 *or* more than one instrument or information source used |
| 2 | Use of an instrument used in previous studies with the same sample |
| 1 | Use of a non-validated measure |
| 0 | Not reported |
| **Determinant (causal risk factor/ risk factor) score (***complete for each different determinant***)** |
| 6 | Analysis with variation in the predictor and adequately balanced, with analysis of change |
| 5 | Analysis with variation in the predictor and adequately balanced, no analysis of change |
| 4 | Analysis with variation in the predictor but inadequately balanced, with analysis of change |
| 3 | Analysis without variation in the predictor, with analysis of change |
| 2 | Analysis with variation in the predictor but inadequately balanced, no analysis of change |
| 1 | Analysis without variation in the predictor, no analysis of change |

***Risk of Bias Assessment***

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| Risk of Bias criteria ↓/ Study Design 🡪 | Randomised controlled trials | Non-randomised controlled trials, pre-and post-test, experimental designs | Non-experimental longitudinal or cross-sectional designs | Notes |
| Random sequence generation (selection bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | For non-RCTs, consider sampling strategy. |
| Allocation concealment (selection bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear |
| Blinding of participants and personnel (performance bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear |  |
| Blinding of outcome assessment (outcome bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | For non-RCTs, consider outcome measurements methods. |
| Incomplete outcome data (attrition bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | For non-RCTs, consider reporting of (1) missing data, (2) response rate, and (3) retention rate. |
| Selective reporting (reporting bias) | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear |  |
| Other bias | [ ]  low risk[ ]  high risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | [ ]  high risk[ ]  low risk[ ]  not clear | Contamination in cluster RCTs. |

1. Scoring to be completed separately for each analyzed association between an outcome and an associated factor. [↑](#footnote-ref-1)
2. Items 1 and 2 will be scored for cross-sectional study designs. [↑](#footnote-ref-2)
3. Response rate scored for cross-sectional study designs. Retention rate scored for longitudinal study design*s.* [↑](#footnote-ref-3)