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2008-04-22

CONSORT for Abstracts Checklist

Scope: Preferred Reporting Items for reporting randomized controlled trials in journal and conference abstracts.

Reference: See `source/variants/consort-abstracts.yml` for canonical link and provenance.

Instructions

- Use the boxes to confirm each reporting item.
- Add reviewer notes under each section as needed.

Checklist Items

- ☐ **Title:** Identification of the study as a randomized trial.
- ☐ **Authors:** Contact details for the corresponding author.
- ☐ **Trial design:** Description of the trial design (e.g., parallel, cluster, non-inferiority).
- ☐ **Methods:**
  - ☐ **Participants:** Eligibility criteria for participants and the settings where the data were collected.
  - ☐ **Interventions:** Interventions intended for each group.
  - ☐ **Objective:** Specific objective or hypothesis.
  - ☐ **Outcomes:** Clearly defined primary outcome for this report.
  - ☐ **Randomization:** How participants were allocated to interventions.
  - ☐ **Blinding (masking):** Whether or not participants, care givers, and those assessing the outcomes were blinded to group assignment.
- ☐ **Results:**
  - ☐ **Numbers randomized:** Number of participants randomized to each group.
  - ☐ **Recruitment:** Trial status.
  - ☐ **Numbers analyzed:** Number of participants analyzed in each group.
  - ☐ **Outcome:** For the primary outcome, a result for each group and the estimated effect size and its precision.

- ? **Harms:** Important adverse events or side effects.

- ? **Conclusions:** General interpretation of the results.
- ? **Trial registration:** Registration number and name of trial register.
- ? **Funding:** Source of funding.

Notes
Reviewer notes