

LOST Information in Trials (LOST-IT) study

Title and Abstract screening Form (version 9)

Screener initials: Study ID: Author, year: _____, **2005**
 Journal: AIM BMJ JAMA Lancet NEJM

1. Eligible RCT? No **→ Exclude, stop here**
 Yes or unclear **→ Get Full Text, answer questions below**

2. Trial described as: Non-inferiority
 Equivalence
 Neither

3. Primary outcome clearly specified Yes, one: _____ (go to q5)
 No, multiple primary outcomes: _____
 _____ (go to q4)
 None specified (go to q4)

4. If multiple or no primary outcome specified, select one : _____

5. Primary outcome category # (refer to the box): _____ (e.g. II.3)

6. The primary outcome is a: Time to event outcome
 Continuous outcome (not time to event)
 Multinomial outcome
 Binary outcome reported as rate
 Binary outcome

7. Is it a composite endpoint? Yes
 No

8. Is it a patient important outcome? Yes
 No
 Not clear

9. Is the result statistically significant? Yes
 No
 Not clear

- I. Mortality
 - 1. all cause mortality
 - 2. disease specific mortality
- II. Morbidity
 - 1. cardiovascular major morbid events
 - 2. other major morbid events
 - 3. recurrence/relapse/remission of cancer and other chronic diseases
 - 4. renal failure requiring dialysis
 - 5. hospitalizations, medical and surgical procedures
 - 6. infections
 - 7. dermatological/ rheumatologic disorders
- III. Symptoms/Quality of life/Functional status
- IV. Surrogate outcomes

Please fill out this box for each study

Exclude
 Get full text
 3rd reviewer needed (no consensus between 2 reviewers)

If exclude, reason for exclusion:

Not RCT
 Not eligible RCT
 Other: