# LOST Information in Trials (LOST-IT) study

## Title and Abstract screening Form (version 9)

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<thead>
<tr>
<th>Screener initials:</th>
<th>Study ID:</th>
<th>Author, year:</th>
<th>Journal:</th>
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### 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

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**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:

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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