**Double blind, you are the weakest link — goodbye!**

*Double blind* is the term researchers frequently use, and readers frequently accept, as a key marker of validity of a randomised controlled trial (RCT). Clinical trial experts and clinicians, when asked, all claim to “know” what *double blind* means; however, unfortunately it means diverse things to those questioned.\(^1\) The term lacks consistency in its use and interpretation — a critical flaw for any technical term if it is to be understood. In this editorial, we advocate abandoning the current blinding lexicon (ie, single, double, and triple blinding) and recommend transparent reporting of the blinding status of each group involved in the execution, monitoring, and reporting of clinical trials.

Blinding (or masking) in RCTs is the process of withholding information about treatment allocation from those who could potentially be influenced by this information. Blinding has long been considered an important safeguard against bias. Benjamin Franklin, in 1784, was probably the first to use blinding in scientific experimentation.\(^2\) Louis XVI commissioned Franklin to evaluate mesmerism, the most popular unconventional “healing fluid” of the eighteenth century.\(^3\) By applying a blindfold to participants, Franklin removed their knowledge of when mesmerism was and was not being applied. Blinding eliminated the intervention’s effects and established mesmerism as a sham.\(^4\) From this work, the scientific community recognised the power of blinding to enhance objectivity and it quickly became, and remains, a commonly used strategy in scientific research.

The groups who can potentially introduce bias into an RCT through knowledge of the treatment allocation are shown in the table.

<table>
<thead>
<tr>
<th>Group</th>
<th>Description</th>
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<tbody>
<tr>
<td>Healthcare providers</td>
<td>The physicians, nurses, physiotherapists, or other personnel who care for the participants during the study period or those who administer the interventions</td>
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<tr>
<td>Data collectors</td>
<td>Individuals who collect data for the study outcomes; data collection could include administering a questionnaire, taking a physical measurement, or eliciting symptoms</td>
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<tr>
<td>Judicial assessors of outcomes</td>
<td>The individuals who ultimately decide whether a participant has had the outcome, or outcomes, of interest</td>
</tr>
<tr>
<td>Data analysts</td>
<td>The individuals who conduct the data analysis</td>
</tr>
<tr>
<td>Data safety and monitoring committee</td>
<td>The committee that reviews the data to advise on the continuing safety of the trial and persistent uncertainty of the efficacy of the intervention being evaluated</td>
</tr>
<tr>
<td>Manuscript writers</td>
<td>The individuals who write versions of the manuscript before the breaking of the randomisation code: in a fully blinded study, one version is written with the assumption that group A received the experimental intervention, and the other is written with the assumption that group B received the experimental intervention</td>
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From this work, the scientific community recognised the power of blinding to enhance objectivity and it quickly became, and remains, a commonly used strategy in scientific research. The groups who can potentially introduce bias into an RCT through knowledge of the treatment allocation are shown in the table.

**Groups that can potentially be blinded in RCTs**

1. **Healthcare providers**
   - The physicians, nurses, physiotherapists, or other personnel who care for the participants during the study period or those who administer the interventions

2. **Data collectors**
   - Individuals who collect data for the study outcomes; data collection could include administering a questionnaire, taking a physical measurement, or eliciting symptoms

3. **Judicial assessors of outcomes**
   - The individuals who ultimately decide whether a participant has had the outcome, or outcomes, of interest

4. **Data analysts**
   - The individuals who conduct the data analysis

5. **Data safety and monitoring committee**
   - The committee that reviews the data to advise on the continuing safety of the trial and persistent uncertainty of the efficacy of the intervention being evaluated

6. **Manuscript writers**
   - The individuals who write versions of the manuscript before the breaking of the randomisation code: in a fully blinded study, one version is written with the assumption that group A received the experimental intervention, and the other is written with the assumption that group B received the experimental intervention

**Performance testing, differential timing or frequency of outcome measurements, and variable recording of outcomes.**\(^5\) Unblinded judicial assessors may introduce bias in their assessment of outcomes, and this is most likely during assessment of subjective outcomes.\(^6\) Unblinded data analysts have the potential to introduce systematic bias through decisions on patient withdrawals, post hoc selection of outcomes or analytic approaches, selection of time points that show the maximum or minimum effects, and many other decisions.\(^7\) Unblinded members of the data safety and monitoring committee may introduce bias at the time of interim analyses through their recommendations to stop or continue a study.\(^8\) Blinding of authors, although seldom done,\(^9\) may reduce the biases in the presentation and interpretation of results.

Case reports document individual examples of the biases described above.\(^7\) Unfortunately, suboptimum reporting of blinding status in RCTs reported as double blinded,\(^10\) the other study did not find any association between the reporting of double blinding and the estimate of effect.\(^11\) Who was actually blinded in these studies probably varied and is certainly open to question. Heterogeneity in who was blinded in the studies reported as double blinded may be responsible for these discrepant findings.

Although the true magnitude of bias introduced by unblinding remains (and is likely to remain) uncertain, clinicians should consider the blinding status of each group in assessing study validity. Unfortunately, suboptimum reporting of blinding status in full-text publications and secondary journals has hindered readers.\(^10\) Authors have commonly relied on conventional blinding terminology (single, double, and triple blinding) to convey blinding status.\(^11\) We have shown great variability in physician interpretations and textbook definitions of these terms.\(^12\) It is for this reason that we recommend, and the editors of...
ACP Journal Club and Evidence-Based Medicine have adopted, a strategy of abandoning the current blinding terminology for transparent reporting of the blinding status of the groups listed in the table. As a result of this policy, readers will be able to make more informed decisions about the validity of the studies on which they base their practice.

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Obituary

Chris Silagy

We were all grieved to hear that long time Associate Editor, Chris Silagy, died at the age of 41 from non-Hodgkin’s lymphoma on the 13th December, 2001. Chris was foundation director of the Monash Institute for Health Services Research, director of the Australasian Cochrane Centre, and chaired the international steering group of the Cochrane Collaboration from 1996 to 1998. Chris was a remarkable man, an influential family physician, and an inspirational leader in evidence-based primary health care. Despite repeated chemotherapy, a splenectomy, and a pleurodesis, he continued to work tirelessly until the day before his death, but also found time for family and friends, to take his wife ballooning, and to write his own obituary, which can be found at http://www.bmj.com:80/cgi/content/full/324/7328/53?maxtoshow=.

The Editors

Journals reviewed for this issue*

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*Approximately 60 additional journals are reviewed. This list is available on request.

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