

# Librarians can help address reporting concerns in the biomedical literature particularly, for systematic reviews – here's how!

CEC 6  
ICML + EAHIL 2017



CONSEJERÍA DE SALUD  
Agencia de Evaluación de Tecnologías  
Sanitarias de Andalucía (AETSA)



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# Common deficiencies in biomedical research reporting

# Deficiencies in biomedical research reporting

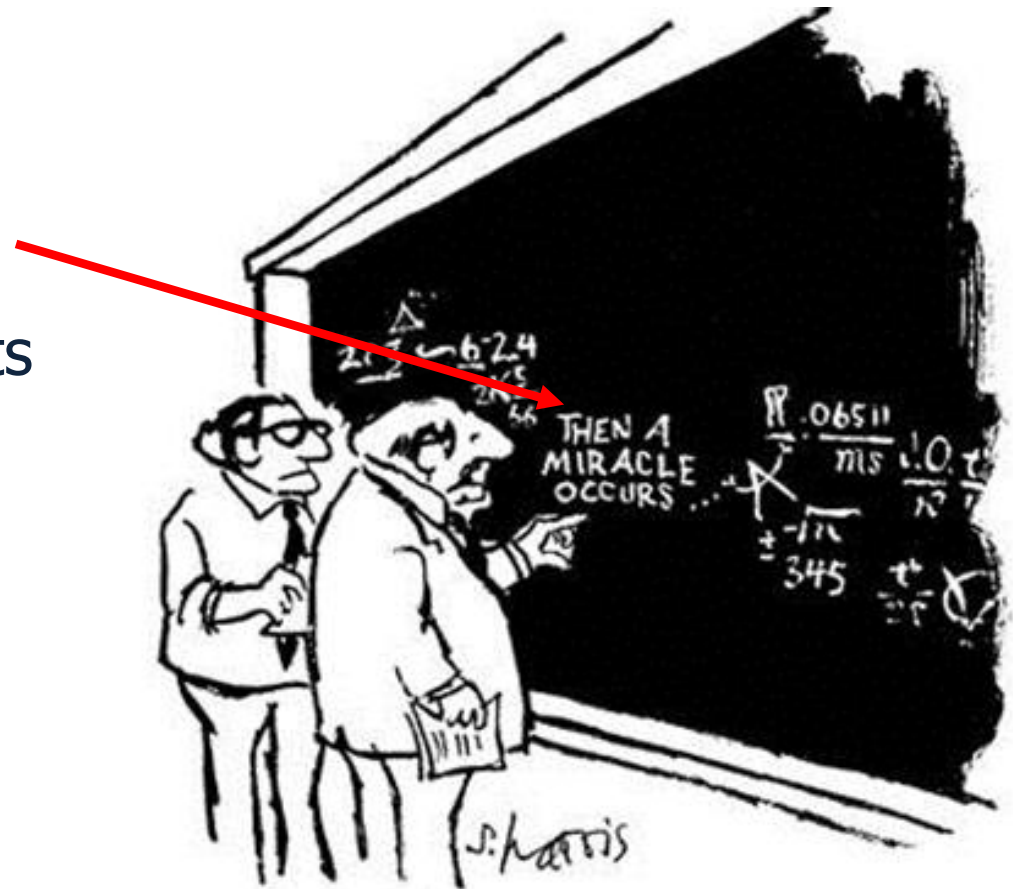
- A research article is often the only available record that a research study was conducted
- Scientific manuscripts should present sufficient data so that the reader can fully evaluate the information
- Readers need a clear understanding of exactly what was done and found

# 5 main areas where deficiencies have been identified

- Non-reporting (or delayed reporting) of studies
  - often studies with disappointing results
- Incomplete reporting
  - e.g. the omission of crucial aspects of the research methods, incomplete results or inadequate reporting of harms
- Selective reporting
  - e.g. selectively reporting patient outcomes or aspects of the analysis
- Misleading reporting
  - e.g. misinterpretation of study findings or using 'spin'
- Unacknowledged discrepancies between sources
  - E.g. where the publication conflicts with the study protocol or the information contained in the register

# Incomplete or unclear reporting

- Hundreds of published reviews show that key elements of *methods* and *findings* are commonly *missing* from journal reports



"I think you should be more explicit here in step two."

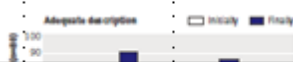
Impact of document type on reporting quality of clinical drug trials: a comparison of registry reports, clinical study reports, and journal publications

## What is missing from descriptions of treatment in trials and reviews?

Replicating non-pharmacological treatments in practice depends on how well they have been described in research studies, say **Paul Glasziou** and colleagues

Have you ever read a trial or review and wondered exactly how to carry out treatments such as a "behavioural intervention,"

receiving numerous requests for additional details from doctors and patients, the author of a randomised trial on graded exercise for



### ARTICLE

## Adequacy of Published Oncology Randomized Controlled Trials to Provide Therapeutic Details Needed for Clinical Application

Jennifer M. Duff, Helen Leather, Edmund O. Walden, Kourtnay D. LaPlant, Thomas J. George Jr

Manuscript received July 9, 2009; revised March 15, 2010; accepted March 16, 2010.

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**Background** Randomized-controlled trials (RCTs) improve clinical care through evidence-based medicine. However, RCT result reporting, but specific details of therapeutic administration and protocol design. We assess the reporting methodology in RCTs published in the medical literature.

**Methods** Ten essential elements of RCT reporting were identified and included a maximum number of cycles, premedication, growth factor support, patient adjustments for hematologic and organ-specific toxicity. All therapy-based RCTs published in the *New England Journal of Medicine (NEJM)*, *Journal of Clinical Oncology (JCO)*, *Journal of*

Clin Chem Lab Med 2012;50(3):411-413 © 2012 by Walter de Gruyter • Berlin • Boston, DOI 10.1515/cclm-2011-0904

## An appeal to medical journal editors: the need for a full description of laboratory methods and specimen handling in clinical study reports

elements.

27 from Cancer, 18% of complete data for toxicity (5/27; 18%), did not substantiate the results. Additional data necessary for

## Exercise prescription: a case for standardised reporting

Susan Carolyn Slade, Jennifer Lyn Keating

### ABSTRACT

**Background** Structured, regularly recommended to improve health takes many forms and varies in its frequency. The authors used exercise for chronic health conditions. Exercise programmes are described in reviews.

Two independent reviews of exercise reporting practices for exercise effects for material. Inclusion criteria characterised the effects of exercise with chronic health conditions. Views of studies of children and adolescent

## Empirical Evidence for Selective Reporting of Outcomes in Randomized Trials: Comparison of Protocols to Published Articles

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**Context** Selective reporting of outcomes within published studies based on the nature or direction of their results has been widely suspected, but direct evidence of such bias is currently limited to case reports.

**Objective** To study empirically the extent and nature of outcome reporting bias in a cohort of randomized trials.

**Design** Cohort study using protocols and published reports of randomized trials approved by the Scientific Ethical Committees for Copenhagen and Frederiksberg, Denmark, in 1994-1995. The number and characteristics of reported and unreported trial outcomes were recorded from protocols, journal articles, and a survey of trials. An outcome was considered incompletely reported if insufficient data were presented in the published articles for meta-analysis. Odds ratios relating the completeness of outcome reporting to statistical significance were calculated. Odds ratios of the extent and nature of outcome reporting bias were used to enhance strength, endurance, flexibility, function.

SELECTIVE PUBLICATION OF STUDIES with statistically significant results has received widespread recognition.<sup>1</sup> In contrast, selective reporting of favorable

These abbreviations. Use of session. Burden of cardiovascular disease, by physical therapy 57.

html#Resolutions). The four main goals are promo-

## Reporting of adverse events in randomized controlled trials of highly active antiretroviral therapy: systematic review

Michal Y. Chowers<sup>1,2\*</sup>, Bat Sheva Gottesman<sup>1,2</sup>, Leonard Leibovici<sup>1,3</sup>, Ulrike Pielmeier<sup>4</sup>, Steen Andreassen<sup>4</sup> and Mijal Paul<sup>1,3</sup>

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### RESEARCH ARTICLE

### Open Access

## Reporting and Interpretation of Randomized Controlled Trials With Statistically Nonsignificant Results for Primary Outcomes

Isabelle Boutron, MD, PhD  
Susan Dutton, MSc  
Philippe Ravaud, MD, PhD  
Douglas G. Altman, DSc

**Context** Previous studies indicate that the interpretation of trial results can be distorted by authors of published reports.

**Objective** To identify the nature and frequency of distorted presentation or "spin" (ie, specific reporting strategies, whatever their motive, to highlight that the experimental treatment is beneficial, despite a statistically nonsignificant difference for the primary outcome, or to distract the reader from statistically nonsignificant results) in published reports of randomized controlled trials (RCTs) with statistically nonsignificant results for primary outcomes.

**Data Sources** March 2007 search of MEDLINE via PubMed using the Cochrane Highly Sensitive Search Strategy to identify reports of RCTs published in December 2006.

## Electronic search strategies to identify reports of cluster randomized trials in MEDLINE: low precision will improve with adherence to reporting standards

Monica Taljaard<sup>1,2\*</sup>, Jessie McGowan<sup>1,3,4,5</sup>, Jeremy M Grimshaw<sup>1,6</sup>, Jamie C Brebault<sup>1,2</sup>, Andrew McRae<sup>7</sup>, Martin P Eccles<sup>8</sup>, Allan Donner<sup>7,9</sup>

### OPEN ACCESS

## Publication Bias in Antipsychotic Trials: A Efficacy Comparing the Published Literature and the Food and Drug Administration Database

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Gregory, United States of America. <sup>2</sup>Department of Health Care, Oregon Health & Science University Medical Center, Portland, Oregon, United States

of evidence-based medicine, yet a drug regulatory agencies, e.g., the US, which data in journal articles can be ch... extent to which it inflates apparent