Seventy-Five Trials and Eleven Systematic Reviews a Day: How Will We Ever Keep Up?

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Thirty years ago, and a quarter of a century after randomised trials had become widely accepted, Archie Cochrane reproached the medical profession for not having managed to organise a “critical summary, by specialty or subspecialty, adapted periodically, of all relevant randomised controlled trials” [1]. Thirty years after Cochrane’s reproach we feel it is timely to consider the extent to which health professionals, the public and policymakers could now use “critical summaries” of trials for their decision-making.

The Landscape

Keeping up with information in health care has never been easy. Even in 1753, when James Lind published his landmark review of what was then known about scurvy, he needed to point out that “…before the subject could be set in a clear and proper light, it was necessary to remove a great deal of rubbish” [2]. And 20 years later, Andrew Duncan launched a publication summarising research for clinicians, lamenting that critical information “…is scattered through a great number of volumes, many of which are so expensive, that they can be purchased for the libraries of public societies only, or of very wealthy individuals” [3]. We continue to live with these two problems—an overload of unfiltered information and lack of open access to information relevant to the wellbeing of patients.

A century later, the precursor of the US National Library of Medicine (NLM) began indexing the medical literature. Between 1865 and 2006, the index grew from 1,600 references to nearly 10 million [4]. Even with the assistance of electronic databases such as NLM’s MEDLINE, the problem of having to travel through and sift vast amounts of data has grown. As mountains of unsynthesised research evidence accumulate, we need to keep improving our methods for gathering, filtering, and synthesising it. Some of the key events in the story so far are shown on the timeline in Figure 1.

A legal regulatory framework overseen by the US Food and Drug Administration (FDA) requiring proof of efficacy of new drugs was introduced in 1962, and other countries followed suit. These developments made it inevitable that randomised trials would increasingly become an important component of the evidence base [5]. Government health technology assessment agencies were also established as policymakers sought to have more reliable evidence of the effects of other forms of health care interventions [6].

As the number of clinical trials grew, so too did the science of reviewing trials. Systematic reviews and meta-analyses seeking to make sense of multiple trials began to appear in a variety of health fields in the 1970s and 1980s (see Box 1). An important early example showed that postoperative radiotherapy after surgical treatment of breast cancer was associated with a previously unrecognised increased risk of death [7]. Another challenged beliefs about vitamin C and the common cold [8]. A third suggested a previously

Summary Points

- When Archie Cochrane reproached the medical profession for not having critical summaries of all randomised controlled trials, about 14 reports of trials were being published per day. There are now 75 trials, and 11 systematic reviews of trials, per day and a plateau in growth has not yet been reached.
- Although trials, reviews, and health technology assessments have undoubtedly had major impacts, the staple of medical literature synthesis remains the non-systematic narrative review. Only a small minority of trial reports are being analysed in up-to-date systematic reviews. Given the constraints, Archie Cochrane’s vision will not be achieved without some serious changes in course.
- To meet the needs of patients, clinicians, and policymakers, unnecessary trials need to be reduced, and systematic reviews need to be prioritised. Streamlining and innovation in methods of systematic reviewing are necessary to enable valid answers to be found for most patient questions. Finally, clinicians and patients require open access to these important resources.
unrecognised advantage of some forms of fetal monitoring during labour in reducing neonatal seizures [9].

By the mid-1980s, the need to minimise the likelihood of being misled by the effects of biases and the play of chance in reviews of research evidence was being made evident in articles [10–14] and textbooks [15]. In 1988, regularly updated electronic publication of systematic reviews and meta-analyses, along with bibliographies of randomised trials, began in the perinatal field [16,17]. This provided a model for the inauguration of the international Cochrane Collaboration in 1993 to prepare, maintain, and disseminate systematic reviews of the effects of health care interventions.

**Where Are We Now?**

Despite this progress, the task keeps increasing in size and complexity. We still do not know exactly how many trials have been done. For a variety of reasons, a large proportion of trials have remained unpublished [18,19]. Furthermore, many trials have been published in journals without being electronically indexed as trials, which makes them difficult to find. One of the first steps in being able to adequately review literature is that scientific contributions which predate digitalised information systems and trial indexing need to be “rediscovered and inserted into the memory system” [20]. Through the 1990s, to identify possible reports of controlled trials, the Cochrane Collaboration mobilised thousands of volunteers around the globe to comb the major databases, and to hand-search nondigitised health literature, unpublished conference proceedings, and books. The result of this collaborative effort is the Cochrane Controlled Trials Register (CCTR) (now called the Cochrane Central Register of Controlled Trials).

The differences between the numbers of trial records in MEDLINE and CCTR (see Figure 2) have multiple causes. Both CCTR and MEDLINE often contain more than one record from a single study, and there are lags in adding new records to both databases. The NLM filters are probably not as efficient at excluding non-trials as are the methods used to compile CCTR. Furthermore, MEDLINE has more language restrictions than CCTR. In brief, there is still no single repository reliably showing the true number of randomised trials. Similar difficulties apply to trying to estimate the number of systematic reviews and health technology assessments (HTAs).

In Figures 2 and 3 we use a variety of data sources to estimate the numbers of trials and systematic reviews published from 1950 to the end of 2007 (see Text S1). The number of trials continues to rise: although the data from CCTR suggest some fluctuation in trial numbers in recent years, this may be misleading because the Cochrane Collaboration virtually halted additions to CCTR as it undertook a review and internal restructuring that lasted a couple of years.
Box 1. Early Systematic Reviews of the Effects of Health Care Interventions


Figure 2. The number of published trials, 1950 to 2007. CCTR is the Cochrane Controlled Trials Registry; Haynes filter uses the “narrow” version of the Therapy filter in PubMed:ClinicalQueries; see Text S1.
doi:10.1371/journal.pmed.1000326.g002
Even though these figures must be seen as more illustrative than precise, multiple data sources tell the same story: astonishing growth has occurred in the number of reports of clinical trials since the middle of the 20th century, and in reports of systematic reviews since the 1980s—and a plateau in growth has not yet been reached. With a median of perhaps 80 participants per trial, the number of people being enrolled in trials is likely to be more than 2,000,000 per year [21]. Prospective trial registration establishes a new genre of evidence repository; trials are registered in these databases at inception, theoretically enabling an overview of all published and unpublished trials.

In 2004, the International Committee of Medical Journal Editors (ICMJE, http://www.icmje.org/) announced that their journals would no longer publish trials that had not been prospectively registered [22]. Before this announcement, an average of 30 trials a week were being prospectively registered around the world. Once the journal editors’ deadline came into force, more than 200 ongoing trials per week were being registered [23]. In 2007, the US Congress made detailed prospective trial registration legally mandatory [24]. As WHO’s international clinical trials platform develops, it will become possible to generate a more realistic picture of how many trials are being done. This registry draws together standardised core data from all the trial registries meeting specified quality criteria. Registering full protocols and reporting trial results in these registries are the next frontiers.

How Close Are We to Archie Cochrane’s Goal?

In 1986 and 1987, Goldschmidt and Mulrow showed how great the potential is for error in reviews of health literature that were not conducted systematically [9,10]. Looking at data such as those in Figure 3 could provide the comforting illusion that systematic reviews have displaced other less reliable forms of information. However, as Figure 4 shows, this is far from the case. The growth has been even more remarkable in non-systematic (“narrative”) reviews and case reports. Journal publishing of non-systematic reviews, and the emergence of many journals whose sole product is non-systematic reviews, has far outstripped the growth of systematic reviews and HTAs, as impressive as the latter has been. And the number of case reports—which can also provide important new information such as adverse effects—is far higher than the number of trials or systematic reviews. Trials, systematic reviews, and HTAs have undoubtedly had major impacts, including on clinical guidelines: they are more likely to be cited and read than other study types [25]. However, the staple of medical literature synthesis remains the non-systematic narrative review.

Furthermore, we are a long way from having all relevant trials incorporated into good systematic reviews. The workload involved in producing reviews is increasing, and the bulk of systematic reviews are now many years out of date [26]. The median number of trials contained within individual systematic reviews has been

![Figure 3. The number of systematic reviews in health care, 1990 to 2007. INAHTA is International Network of Agencies for Health Technology Assessment; the Montori systematic review filter is detailed in Text S1. doi:10.1371/journal.pmed.1000326.g003](image-url)
variously estimated at between six and 16
(Cochrane reviews now include an average
of over 12 trials per review [27,28]; M
Clarke, personal communication), but
many reviews have covered much the
same territory. Thus, in the 30 years since
systematic reviews began in earnest, with
around 15 years of intensified and large-
scale reviewing effort, only a minority of
trials have been assessed in systematic
reviews. Given the triple constraint posed
by the growth in trials, the increasing
complexity of review methods, and current
resources, Archie Cochrane’s vision will
not be achieved without some serious
changes in course—in particular, with a
greater concentration on Cochrane’s use
of the word “relevant”.

Where to Now?

First, we need to prioritise effectively
and reduce avoidable waste in the pro-
duction and reporting of research evidence
[29]. This has implications for trials as well
as systematic reviews. Some funders and
others will now not consider supporting a
trial unless a systematic review has shown
the trial to be necessary [30]. It is essential
that this requirement be more widely
adopted. And it is essential that reviews
address questions that are relevant to
patients, clinicians and policymakers.

Second, we may need to choose be-
tween elaborate reviews of a quarter of the
questions clinicians and patients have or
“leaner” reviews of most of what we want
to know. The methodological standards
for systematic reviewing have been in-
creasing over time [28], and the evolution
of standards in the Cochrane Collabora-
tion and in HTA has been remarkable.

The increase in steps and reporting
required is reflected in the length of
reviews. Early Cochrane reviews could
typically be printed out in 10 or 20 pages,
even when they incorporated several trials.
Today, it is not unusual for a review by a
health technology agency to run to several
hundred pages. Often the reviews are
longer than the combined length of the
reports of all the included trials.

A contributing factor here is the in-
creasing expectation for reviews to include
study types other than randomised trials.
This will often be essential for detecting
less common adverse effects. However, the
inclusion of all study types to answer all
questions about the effects of treatments
would not necessarily provide better
quality information in every instance—
while it would unquestionably increase the
time and resource requirement for re-
views. While it is vital that reviews are
scientifically defensible, burdening those
preparing them with excessive require-
ments could result in having valid answers
to relatively few questions.

In particular, we need leaner and more
efficient methods of staying up-to-date
with the evidence. Using current methods,
the Cochrane Collaboration has not been
able to keep even half of its reviews up-to-
date [31], and other organisations are in a
similar predicament [32]. We need to
develop innovative methods to reduce the
labour of updating, and provide what
clinicians and patients need; an assurance
that a conclusion is not out of date, even if
not every later trial is included within
every analysis. It is also the responsibility
of reviewer authors and journal editors to
ensure that every new systematic review
places itself clearly in context of other

Figure 4. The rise in non-systematic reviews, case reports, trials, and systematic reviews, 1950 to 2007 (as identified in MEDLINE).
doi:10.1371/journal.pmed.1000326.g004
systematic reviews and HTAs. It will be to little avail to the average clinician, patient, and information provider, however, if the resulting knowledge is not comprehensible and openly accessible.

Finally, although more funding for evaluative clinical research internationally remains a priority, more international collaboration could result in better use being made of resources for systematic reviewing and HTAs. While multiple reviews on topics can provide a rounded picture of an area as well as a de facto form of updating when the reviews are conducted several years apart, there is also considerable duplication of review effort.

In November 2009, an international meeting in Cologne formed a new collaboration called “KEEP Up,” which will aim to harmonise updating standards and aggregate updating results. This should reduce the workload and enable organisations to be alerted when there are important shifts in evidence. Initiated and coordinated by the German Institute for Quality and Efficiency in Health Care (IQWiG) and involving key systematic reviewing and guidelines organisations such as the Cochrane Collaboration, Duodecim, the Scottish Intercollegiate Guidelines Network (SIGN), and the National Institute for Health and Clinical Excellence (NICE), this effort will provide a platform for tackling practical and methodological issues involved in keeping up-to-date.

There is nevertheless a risk that the increasing burdens placed on the methods of systematic reviewing could make the goal of keeping up-to-date with the knowledge won from trials recede ever more quickly into the distance. Perhaps one of the first questions we should ask whenever an additional process or more demanding methodology for systematic reviewing is proposed is this: Will this development serve or hinder our ability to better understand and communicate enough results from trials? In 1979, Archie Cochrane argued that we needed critical summaries to keep up with the crucial knowledge those trials were generating; there were perhaps 14 trials a day being published. Thirty years later, it would be just as hard to keep up with the systematic reviews. Every day there are now 11 systematic reviews and 75 trials, and there are no signs of this slowing down: but there are still only 24 hours in a day.

## Supporting Information

### Text S1 Search methods.

Found at: doi:10.1371/journal.pmed.1000326.s001 (0.03 MB DOC)

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### Acknowledgments

We are grateful to Sigrid Droste from the German Institute for Quality and Efficiency in Health Care (IQWiG) for her guidance on National Library of Medicine documentation on changes in reporting of publication types across time.

### Author Contributions

ICMJE criteria for authorship read and met: HB PG IC. Agree with the manuscript’s results and conclusions: HB PG IC. Designed the experiments/the study: HB. Analyzed the data: HB. Collected data/did experiments for the study: HB. Wrote the first draft of the paper: HB. Contributed to the writing of the paper: PG IC. Conceived the idea, participated in searching, data collection, and analysis for both the historical and publication trends: HB. Guaranteed for the article: HB. Search and analyzed publication trends: PG. Participated in the searching and analysis of historical trends: IC.