



Tracking the impact of research on policy and practice: a protocol for the use of clinical guidelines in research evaluation

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1. Title page

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2. Conflicts of Interest Notification Page

All authors have completed the Unified Competing Interest form at www.icmje.org/coi_disclosure.pdf (available on request from the corresponding author) and declare that (1) DK also holds the position of Research Leader at RAND Europe; and (2) DK, LA, KD, BS and IV have no non-financial interests that may be relevant to the submitted work.

For peer review only

3. Abstract

Introduction

In recent years, the medical research community has come under increasing pressure to demonstrate the value and extent of the impacts of its work . In response, funders have made considerable efforts to enhance their ability to track and understand the impact of their funded research. These efforts will provide evidence of the strategic significance and 'value' of investments into particular areas of research while also helping inform future funding strategies. However, determining the impact of research is challenging, particularly in the area of basic and fundamental research where the time lag between original research subsequent impacts on health can be long and the attribution difficult to disentangle.

Perhaps one of the most challenging areas of research impact evaluation is the quest to understand how and when research reaches policy and practice. Clinical guidelines, both national and international, bring together current high quality evidence concerning the prevention, diagnosis, prognosis and therapy of clinical problems. The inclusion of specific research results in the formulation of clinical guidelines presents us with an indication of research use in medical/clinical practice.

Methods and analysis

In 2009, the UK Medical Research Council, Wellcome Trust and Department of Health (England) commissioned a detailed analysis of research cited in two National Institute for Health and Clinical Excellence (NICE) UK clinical guidelines; Dementia and Chronic Obstructive Pulmonary Disease (COPD). The purpose was to explore the potential of using the guidelines as part of a toolkit to support impact tracking.

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Ethics

There were no ethical issues or considerations necessary for this study.

Dissemination

This paper presents an analysis of the research cited in clinical guidelines and considers how this type of information might have a more systematic role to play in enhancing our understanding of the pathways and time lags involved in research reaching policy and practice.

For peer review only

4. Introduction

Medical research has advanced rapidly in recent times in all areas from basic sciences (i.e. decoding the human genome) through the development of more precise diagnostic tools to novel treatments. At the same time, public interest in biomedical advances and the appetite for more effective treatments¹ are increasing in parallel. The demand for the results of biomedical research to lead to improvement in healthcare has never been higher^{2, 3, 4}.

Across the research world, and particularly in the biosciences, there has been a drive to better demonstrate and understand the impacts of research, essentially so that funds can be allocated to maximum effect. ¹ There remains a concern that the research community as a whole could be better at translating the findings of medical research into tangible health and health care benefits^{5, 6, 7}. Thus, the need to better understand research impact and the pathways to that impact is a key priority for research funders^{8, 9, 10, 11}. However, determining the impact of research is challenging, and particularly in basic and fundamental research where the time lag between original research subsequent impacts on health can be long and the attribution difficult to track. In addition, perhaps one of the most challenging areas has been trying to understand the impact of research on policy and practice.

National and international clinical guidelines are intended to bring together the best and most current evidence about the prevention, diagnosis, prognosis and therapy of clinical problems. Clinical guidelines are a form of systematic review and, in the UK, focus on the defined medical needs of the National Health Service (NHS). In the UK, clinical guidelines are provided by the National Institute for Health and Clinical Excellence (NICE¹²) and, since 2005, these recommendations have had legal standing in the NHS in England and Wales. The guidelines exist to

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2
3 help standardise and improve patient care and can help to introduce cost-
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5 efficiencies to the delivery of health care. The guidelines are evidence-based and
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7 their formulation brings pieces of important and influential research together. For
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9 a funder, if research they have supported is referenced as part of the evidence
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11 supporting a national and/or international clinical guideline, then it is a direct
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13 indication that a piece of research is influencing policy and practice. Hence clinical
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15 guidelines are potentially an attractive resource to support impact tracking and
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17 assessment^{13, 14}.

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21 For those engaged in evaluation, historically it has been difficult to extract
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23 information from a guideline in a way that helps support analysis of the
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25 references and funding sources: simply put, UK clinical guidelines are not
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27 designed to support the requirements of funders trying to track the impact of
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29 their support. However, work is underway at the National Centre for
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31 Biotechnology Information (NCBI, part of the National Library of Medicine, NIH),
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33 to digitise the content of major international clinical guidelines to encourage wider
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35 access to their content and enable greater ability to mine their content and allow
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37 automated links to individual cited research via databases such as PubMed.

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40 In 2009, the UK Medical Research Council (MRC), Wellcome Trust and National
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42 Institute of Health Research (NIHR) – who among them commit nearly £2bn
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44 annually to support biomedical and applied health research - commissioned a
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46 detailed analysis of the research cited on a small number of UK clinical guidelines
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48 to explore the potential of the information in broader research impact tracking.
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50 The objectives of this research were threefold. First, this study explored the
51
52 feasibility of *extracting* the funding source of the research papers cited on a
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54 guideline. Second, it identified *who funded* the research cited in the selected
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56 clinical guidelines. Third, it explored the extent to which there are shared
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58 characteristics of the publications cited in these guidelines.
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5 The then-current NICE guidelines for the management and treatment of two
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7 disease areas were selected: Dementia (2006) and Chronic Obstructive
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9 Pulmonary Disease (2004)^{15, 16}. These guidelines were of interest for the purposes
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11 of this analysis since (a) they had been available unchanged for several years,
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13 and (b) there was a likelihood that all three project sponsor funders would have
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15 funded some of the underlying research evidenced in the guidelines. The two
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17 guidelines were also in quite different clinical areas, so we wanted to see if there
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19 were differences in the process and/or adoption of research into practice. For
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21 each guideline, all cited research was examined to pick out its characteristics
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23 (e.g. age; bibliometric indicators) and identify any funding attributions.
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5. Methods

Data extraction

The first step was to extract a list of publications from each of the guidelines and export them into a Microsoft Excel spreadsheet. This was performed automatically using bespoke RAND Europe computer scripts, based on the PERL scripting language. Here we briefly describe the methodology, since a full description is available elsewhere¹⁷. A total of 744 references were extracted from the Dementia guideline and 446 for the COPD guideline.

Data cleaning

The extracted bibliographic references were cleaned and structured to permit analyses of funding source and paper performance indicators. Any references identified as non-academic, peer-reviewed publications (e.g. references to a website, grey literature) and all publications before 1980 were removed, since these could not be investigated using the Web of Science.

After extraction and initial cleaning, a total of 616 references were found for the Dementia guideline (79.4% of the original 776 references) and 412 references for the COPD guideline (83.9% of the original 491 references).

Data processing

For the funding analyses, the extracted publications were searched for in Web of Science¹⁸ to find the institution and country affiliations of the authors listed. One aim was to identify the publications with at least one UK author on the assumption that this would facilitate further funding analysis. Another aim was to use all the extracted publications for further bibliometric analysis (see ¹⁹ for a similar methodology).

From the Dementia guideline, 494 out of the 616 extracted publications

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2
3 (80.1%) were found in the Web of Science. While from the COPD guideline, 335
4 out of 412 publications were available (81.3%). Any publications not found in the
5 Web of Science were processed individually through a search methodology
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7 utilising the publication libraries of RAND Europe and Cambridge University. All
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9 search processes were duplicated by a second researcher to eliminate errors.
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14 This methodology used a simple search of both the RAND library by article title,
15 using Google Scholar, and the Cambridge University online library and free access
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17 journals, through the Google search engine. If the author affiliation and country
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19 remained unidentified, then the RAND library was searched by journal, followed
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21 by browsing for the article using the reference data available. In addition, the title
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23 could be searched for by keyword within the journal. Finally, where possible,
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25 Cambridge University print holdings were searched to find any articles that were
26
27 not accessible online.
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30 31 **Funder acknowledgement** 32

33 Where publications had at least partial UK attribution, the funding source was
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35 searched for in the Research Outputs Database (ROD)^{20, 21}. This is a database
36
37 housed at the Wellcome Trust recording the funding sources for UK and Irish
38
39 publications in the biomedical sciences for the period 1988–2001. Funding
40
41 acknowledgements were found for around one-third of publications using this
42
43 database. For the remaining publications, which fell outside of the appropriate
44
45 date range or were not found in the ROD, the full text of the publication was
46
47 found and funding acknowledgements recorded directly, where available.
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51 Funding source references were then standardised and categorised by broad
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53 sector. UK funders acknowledged on cited papers were categorised into the
54
55 following categories: industry, not-for-profit, hospital trust, government
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3 department, government agency (not controlled by ministries), local or regional
4 authority, foundation, none given and unknown.
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9 **Bibliometric & paper characteristic analysis**

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11 Author affiliations and country locations were identified for 595 of the 616 (97%)
12 Dementia guideline publications and 402 of the 412 (98%) COPD guideline
13 publications. These affiliated papers form the basis of the following descriptive
14 and bibliometric analysis.
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21 The citation impact of the publications referenced in both clinical guidelines was
22 analysed by sector using the concept of citation profiling. This is based on a
23 normalising technique called ReBased Impact (RBI), which takes account of the
24 field in which a paper appears and the date since its publication to effectively
25 provide a proxy measure for the 'quality' of each paper. The world average
26 RBI=1; the most highly cited articles have an RBI >8²².
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35 Whole counts were used throughout this analysis; if more than one funding
36 source is cited in a publication, this was recorded as *one* publication for each
37 funding source.
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6. Results

Attribution by funding organisation

In a large proportion of cited papers, no funding acknowledgement was listed.

Nearly half (104 of the 228; 46%) of publications in the Dementia guideline with a UK author did not acknowledge any funder. Funding information was available for 117 papers, with the full text of 7 papers inaccessible and, hence, missing the funding information. For the 148 publications in the COPD guideline with at least one UK author, 60 included no funding acknowledgement (41%). Funding information was available for 81 publications. The full text of 7 publications was not accessible and therefore no funding data was available.

Examination of the funding acknowledgements for the Medical Research Council (MRC), Department of Health (England), NHS and the Wellcome Trust revealed that these funders were overtly linked to only a small proportion of papers cited on the guidelines (see figure 1 – Numbers of publications by funding source).

Attributions by funding organisation over time

To determine whether the practice of acknowledgement of funding has improved over time, funding acknowledgement was analysed by year of publication.

Although it appears that more recent publications have more complete funding acknowledgements than older ones, over the whole period, and for both guidelines, there was no clear statistical relationship between the age of publication and the presence or absence of a funding acknowledgement (see figure 2 – Funding acknowledgement by year).

The clinical guidelines, on the whole, cited recent research; the majority of research papers cited in these two UK clinical guidelines were published after 2000 - that is, within five years of the release of these guidelines. Although the

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3 average duration between the publication date of papers cited and the publication
4 date of the citing guideline was 5 years for Dementia and 3 years for COPD.
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7 8 **Attribution by funding sector** 9

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11 Industry was not as prominent a funding source for publications cited in the
12 Dementia guideline, where acknowledgements were distributed across a range of
13 funding sources across sectors. In the COPD guideline, industry was the most
14 frequently linked funder after 'none given'.
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19 20 21 **Attribution by country** 22

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24 Of the 616 publications extracted from the Dementia guideline, 228 (37.2%) had
25 at least one UK-based author; while from the COPD guideline, 148 publications
26 (35.9%) had at least one UK-based author. Researchers based in the UK and US
27 combined were linked to the majority of papers cited in both guidelines.
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34 Despite this dominance of the UK and USA-based researchers in the cited papers,
35 many other countries contributed to these publications. Papers cited in the
36 Dementia and COPD guidelines were linked to authors from 37 and 36 countries,
37 respectively (see figure 3 – Contribution of the most active countries).
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42 43 44 **Attribution by research sector** 45

46 The three research funders use different means of disbursing their money.
47 Funding includes grants to universities and hospitals, alongside direct support for
48 intramural research. Publication analysis by associated institutions revealed that
49 researchers with university addresses, followed by those with hospital addresses,
50 were linked to the bulk of papers cited on both guidelines. More than 80% of
51 publications cited in the two guidelines involved authors based at universities.
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3 The scientific contribution from other types of publicly funded institutions, as well
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5 as from non-profit institutions was low.
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9 As described, nearly 20% of the publications cited in the COPD guideline involved
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11 authors from industry - a slightly higher proportion than in the Dementia
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13 guideline.
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15 16 17 **Citation quality**

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19 Clinical guideline drafting committees are obliged to base their recommendations
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21 on the 'best' research available. The citation impact of the publications referenced
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23 in both clinical guidelines was analysed by sector using the concept of citation
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25 profiling.
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29 Overall papers cited across both guidelines had high RBIs. Papers linked to
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31 universities, companies (industry) and publicly-funded research institutions were
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33 particularly highly cited. At the time of the analysis, for the COPD guideline, cited
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35 papers linked to publicly-funded research institutions had $RBI > 8$; for the
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37 dementia guideline papers linked to universities, companies and publicly-funded
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39 research institutions all had particularly impressive RBIs (see figure 4 – Citation
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41 score by institutional sector).
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7. Discussion

Funding attribution

Perhaps the greatest challenge in research impact assessment is dealing with attribution. Attribution of research outcomes and impacts to a specific funder is complex and, in many cases, improbable. This arises from most medical research receiving funding from multiple sources, involving a host of researchers (often across institutions) and being incremental such that considerable time elapses between original research and impact on health. The tide has turned on this issue and funding bodies are increasingly working together to identify where their funding has made a difference and contributed to an outcome or impact. Exclusively 'claiming' impact ignores the complexities and reality of scientific research and we are more interested in noting our contribution alongside others and learning from this.

Nevertheless, there is much we can do to help us better understand the connection between funding inputs and changes in medical practice. This research project was intended to flesh out some of the issues that we face in trying to link research funding to research output (i.e. research papers) and specific outcomes (clinical guidelines).

As described, there was some variability in the quality of acknowledgement information provided on papers. In our analysis, we did not explore whether the differences in acknowledgement quality and completeness varied according to the nature of the paper (i.e. underpinning versus more applied research). A study of 43 UK clinical guidelines (and associated Health Technology Assessments) related to cancer demonstrated that the number of funding sources acknowledged in papers varied with the 'basicness' of the publications: "*the more clinical papers have fewer (funding) sources and the more basic papers have more*"¹⁴.

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5 An interesting direction for future research work on clinical guidelines would be to
6 investigate whether the high proportion of cited publications with at least one UK
7 author in UK clinical guidelines could be the result of a specific funding strategy.
8 That is, some research could be being funded to help create evidence for the
9 development of clinical guidelines. This specific funding strategy might explain
10 partially why UK-authored publications are over-cited in UK guidelines, relative to
11 the share of publications.
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20 The advice given to funding recipients by UK funders regarding how they should
21 be acknowledged in publications, and the extent to which these requirements
22 have been enforced, has varied significantly over the last two decades. While
23 most funders have included a requirement for acknowledgement in their terms
24 and conditions of award, it has only been since 2008 that there has been
25 published guidance about a standard format²³. This may explain why industry is
26 fairly highly cited across both guidelines as part of a researcher's more stringent
27 contractual obligations. However, it is worth noting that the extent to which
28 researchers are following the standard format is unknown. Furthermore, the
29 reality remains that, given the incremental nature of much research, it is not easy
30 to precisely attribute a publication to its source of funding.
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44 One broader question is how to ensure that research information accessible is
45 required to avoid spending funds on research that either cannot be used or may
46 be duplication²⁴.
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52 **Temporal issues**

53 We did find a correlation between the publication of the clinical guidelines and the
54 dates of the papers cited within it. These results corroborate earlier research¹⁴.
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3 Other bibliometric studies on UK clinical guidelines also found that a significant
4 share of publications cited are published within 10 years before the release of
5 these guidelines^{25, 26}.
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10 Therefore, this finding was not unexpected, since clinical guidelines should be
11 based on new evidence, although on occasion the newest evidence may not be
12 the best. However, it would be interesting to investigate whether the most recent
13 publications cited are the outcome of research specifically funded for to support
14 the development of a clinical guideline, thus potentially explaining the peak in
15 publications cited in the two guidelines a few years before their release.
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21 22 **National contributions**

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24 The UK's contribution is high in both guidelines, since its world share in medical
25 research measured by its share in the total number of publications published in
26 the field amounted to approximately 8.6% in 2006²⁷. This high proportion of
27 papers linked to UK-based authors echoes the finding of previous studies.
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31 Interestingly, our analysis – where around a third of papers 35% of papers are
32 linked to UK-based authors - reveals a higher proportion than we have seen in
33 other analyses.
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41 42 **Conclusions and recommendations**

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44 Having greater access to the work cited on clinical guidelines would present new
45 opportunities for funders and the research community alike to better understand
46 some of the mechanisms that take research from the bed to the bedside. While
47 this would be but one tool in the research impact evaluator's toolkit, it would be
48 one that could be relatively easy to harness if both access and acknowledgements
49 were improved. As described, work is underway between the UK NICE and the
50 NCBI, and in the future, it is envisioned that other guideline providers will make
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3 their content available in much more structured and accessible formats to permit
4 analyses of this nature.
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9 However, to bring clarity to the tracking of research inputs through to published
10 output, the age old problem of ensuring accurate and complete acknowledgement
11 information on peer-reviewed published work needs to be addressed. This
12 requires a change in the culture of how researchers use acknowledgements and
13 greater liaison with publishers and information providers who could do much to
14 enhance the quality and completeness of funding data as a paper is submitted for
15 publication. Clarity and perhaps demarcation between the requirements of an
16 'acknowledgement' section on a paper and description of the 'funding' that has
17 supported the work would seem an easy step to help remedy this²⁸. This is
18 particularly pertinent as information providers such as Thomson Reuters and
19 Elsevier are now developing complex reporting and analytical tools that provide
20 an ability to scrutinise the characteristics of published work – including who is
21 described as funding the work - in new and exciting ways.
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36 Moves to address the definition of an 'acknowledgement' may also help to
37 address the issue of 'over-authorship' and author inflation that has been seen in
38 recent years. This is thought to be fuelled in part by the drive towards impact
39 assessment through national research allocation formulae; such as, the UK
40 Research Assessment Exercise²⁹. Some contributors listed as 'authors' might be
41 more appropriately 'thanked'. Many journals now contain a section that asks for
42 a description of 'contributions' – these often do not contain all those listed in the
43 author list so there may be a place for a more defined 'acknowledgement as
44 thanks' section.
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55 Funders should also ensure clarity around their requirements for a 'funding
56 acknowledgement'. Complexities arise for example, when research takes place in
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3 buildings funded by a specific donor or when a specific piece of research uses a
4 piece of equipment funded by a specific donor. And for how long after funding
5 has been received by a researcher should they continue to provide a funding
6 acknowledgement?
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12 We found that there is great potential for national and international guidelines to
13 be used as sources of information on the impact of research on practice; the
14 challenge is to be able to harness that information in an efficient way – so that
15 we are able to use this information to feed into future research strategy and
16 thereby make the research cycle more effective.
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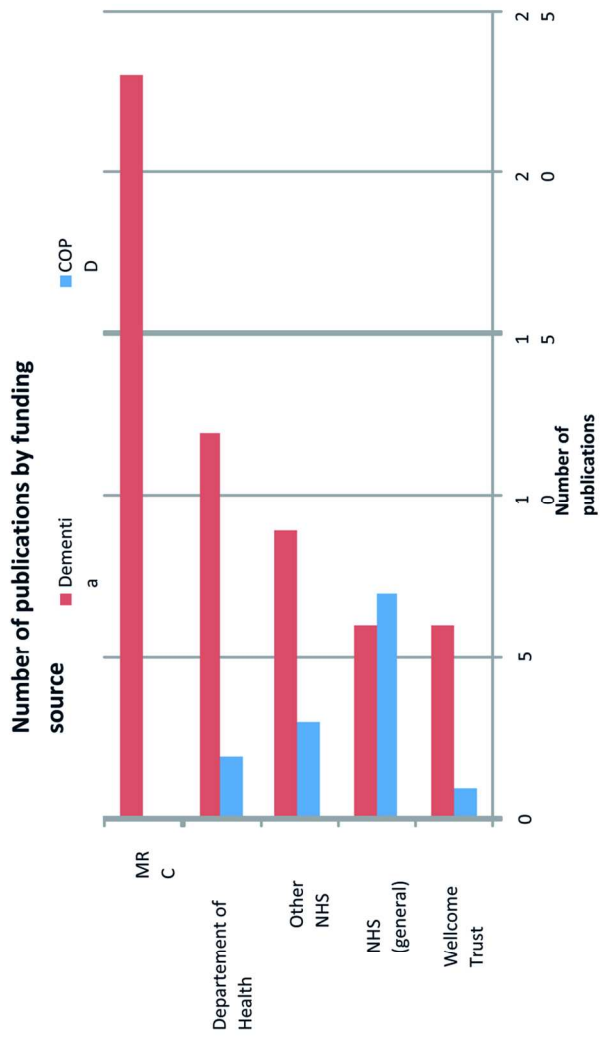
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Contributorship

All authors (DK, LA, KD, BS and IV) contributed extensively to all parts of this paper and commented on the manuscript at all stages. All authors were involved in the design of the study with some guidance on direction and feasibility from RAND Europe. All of the authors discussed the interpretation of the results of the study, and agreed the outline content for this publication. DK wrote the initial draft of the paper which was then reviewed and further shaped by all of the authors. BS prepared the final version of the paper and related information for submission for publication.

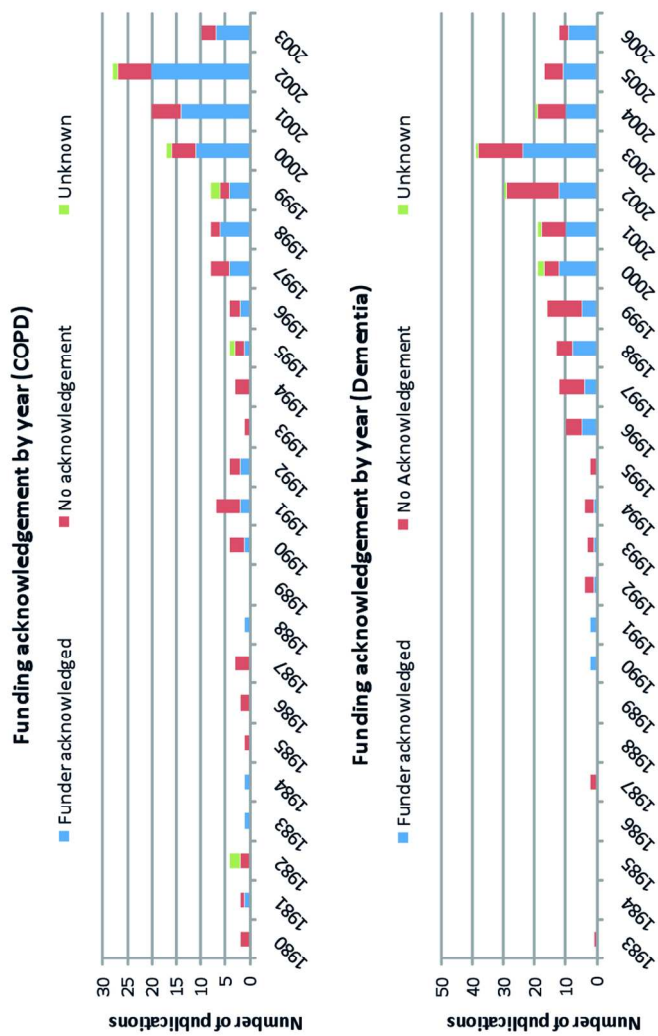
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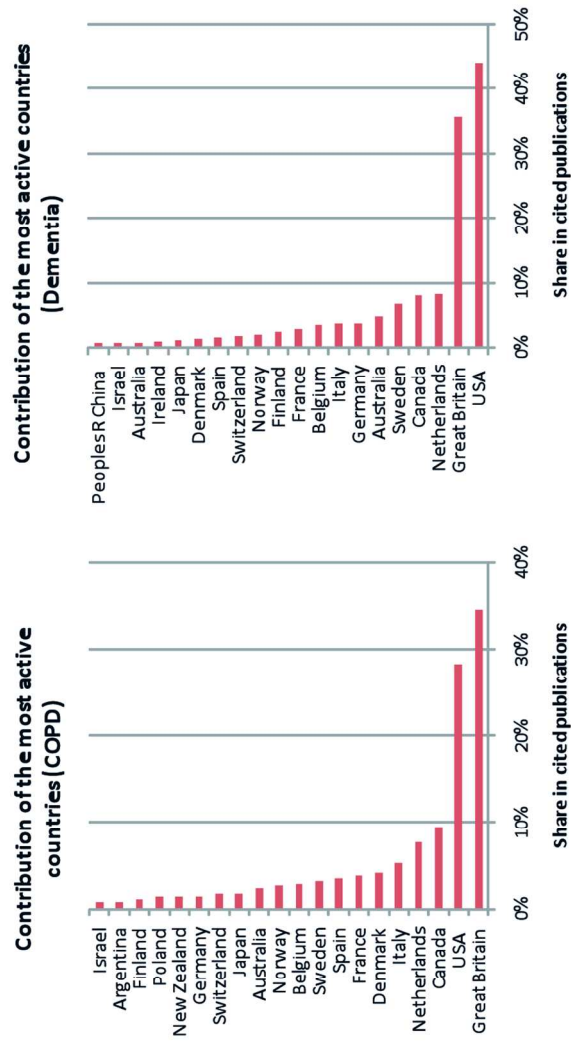
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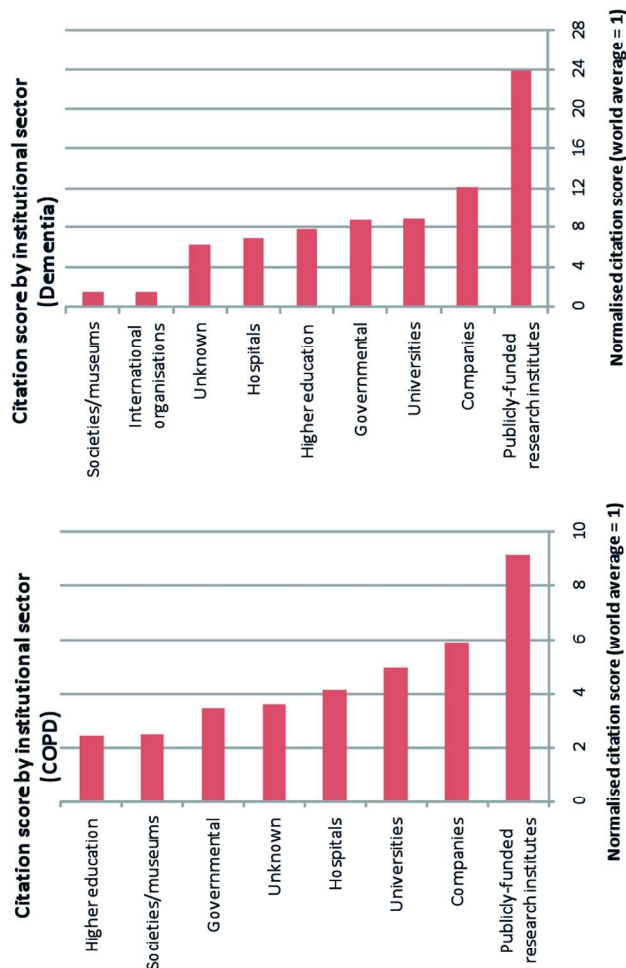
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TRACKING THE IMPACT OF RESEARCH ON POLICY & PRACTICE : THE USE OF CLINICAL GUIDELINES IN RESEARCH EVALUATION.

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**Tracking the impact of research on policy and practice:
investigating the feasibility of using citations in clinical
guidelines for research evaluation.**

Journal:	<i>BMJ Open</i>
Manuscript ID:	bmjopen-2012-000897.R1
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Date Submitted by the Author:	02-Mar-2012
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Primary Subject Heading:	Medical management
Secondary Subject Heading:	Communication, Health informatics, Evidence based practice, Health policy
Keywords:	GENERAL MEDICINE (see Internal Medicine), Health informatics < BIOTECHNOLOGY & BIOINFORMATICS, Protocols & guidelines < HEALTH SERVICES ADMINISTRATION & MANAGEMENT

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3. Abstract

Objectives

To investigate the feasibility of using research papers cited in clinical guidelines as a way to track the impact of particular funding streams or sources.

Setting

In recent years, medical research funders have made efforts to enhance the understanding of the impact of their funded research, to provide evidence of the 'value' of investments in particular areas of research. One of the most challenging areas of research evaluation is around impact of policy and practice.

In the UK, the National Institute of Health and Clinical Excellence (NICE) provide clinical guidelines, which bring together current high quality evidence on the diagnosis and treatment of clinical problems. Research referenced in these guidelines is an indication that of its potential to have real impact on health policy and practice.

Design

This study is based on analysis of the authorship and funding attribution of research cited in two NICE clinical guidelines; dementia and chronic obstructive pulmonary disease (COPD).

Results

Analysis identified that around a third of papers cited in the

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2
3 **two NICE guidelines had at least one author based in the UK.**
4 **In both cases about half of these UK attributed papers**
5 **contained acknowledgements which allowed the source of**
6 **funding for the research to be identified.**
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12 **The research cited in these guidelines was found to have**
13 **been supported by a diverse set of funders from different**
14 **sectors. The study also investigated the contribution of**
15 **research groups based in Universities, industry and the**
16 **public sector.**
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22 23 24 **Conclusions**

25 **The study found that there is great potential for guidelines to**
26 **be used as sources of information on the quality of the**
27 **research used in their development, and that it is possible to**
28 **track the source of the funding of the research. The challenge**
29 **is in harnessing the relevant information to track this in an**
30 **efficient way.**
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4. Introduction

Medical research has advanced rapidly in recent times in all areas from basic sciences (i.e. decoding the human genome) to the development of more precise diagnostic tools and novel treatments. At the same time, public interest in biomedical advances and the appetite for more effective treatments¹ are increasing in parallel. The demand for the results of biomedical research to lead to improvement in healthcare has never been higher^{2, 3, 4}.

Across the research world, and particularly in the biosciences, there has been a drive to better demonstrate and understand the impacts of research, essentially so that funds can be allocated to maximum effect.¹ There remains a concern that the research community as a whole could be better at translating the findings of medical research into tangible health and healthcare benefits^{5, 6, 7}. Thus, the need to better understand research impact and in particular the pathways to that impact is a key priority for research funders^{8, 9, 10, 11}. However, determining the impact of research is challenging, particularly in basic and fundamental research where the time lag between original research and subsequent impacts on health can be long and the attribution difficult to track. In addition, perhaps one of the most challenging areas has been trying to understand the nature of the pathways to, and subsequent impact of research on policy and practice.

National and international clinical guidelines are intended to bring together the best and most current evidence about the prevention, diagnosis, prognosis and therapy of clinical problems. Clinical guidelines are a form of systematic review and, in the UK, focus on the defined medical needs of the National Health Service (NHS). It should be noted that clinical guidelines are not standards of care but are recommendations to the non-specialist or GP. In the UK, clinical guidelines are provided by the National Institute for Health and Clinical Excellence (NICE¹²) and, since 2005, these have had legal standing in the NHS in England and Wales.

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3 As such the results of this particular study are potentially limited to the NHS,
4 however, the methodology could possibly be applied to guidelines governed by
5 other bodies. The guidelines exist to help standardise and improve patient care
6 and can help to introduce cost-efficiencies to the delivery of healthcare. The
7 guidelines are evidence-based and their formulation brings pieces of important
8 and influential research together. For a funder, if research it has supported is
9 referenced as part of the evidence supporting a national and/or international
10 clinical guideline, then it is an indication that a piece of research is likely to be
11 influencing policy and practice. Hence clinical guidelines are potentially an
12 attractive resource to support impact tracking and assessment^{13, 14}.
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24 For those engaged in evaluation, historically it has been difficult to extract
25 information from a guideline in a way that helps support analysis of the
26 references and funding sources: simply put, UK clinical guidelines are not
27 designed to support the requirements of funders trying to track the impact of
28 their support. However, work is underway at the National Centre for
29 Biotechnology Information (NCBI, part of the National Library of Medicine, NIH),
30 to digitise the content of major international clinical guidelines to encourage wider
31 access to their content and enable greater ability to mine their content and allow
32 automated links to individual cited research papers via databases such as
33 PubMed.
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46 In 2009, the UK Medical Research Council (MRC), the Wellcome Trust and the
47 National Institute of Health Research (NIHR), who among them commit nearly
48 £2bn annually to support biomedical and applied health research, commissioned a
49 detailed analysis of the research cited on a small number of UK clinical guidelines
50 to explore the potential of the information in broader research impact tracking.
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52 The objectives of this research were threefold. First, this study explored the
53 feasibility of *extracting* the funding source of the research papers cited on a
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3 guideline. Second, it identified *who funded* the research cited in the selected
4 clinical guidelines. Third, it explored the extent to which there are shared
5 characteristics of the publications cited in these guidelines.
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11 The then-current NICE guidelines for the management and treatment of two
12 disease areas were selected: dementia (2006) and chronic obstructive pulmonary
13 disease (2004)^{15, 16}. These guidelines were of interest for the purposes of this
14 analysis since (a) they had been available unchanged for several years, and (b)
15 there was a likelihood that all three project sponsor funders would have funded
16 some of the underlying research evidenced in the guidelines. The two guidelines
17 were also in quite different clinical areas, so we wanted to see if there were
18 differences in the process and/or adoption of research into practice. For each
19 guideline, all cited research was examined to pick out its characteristics (e.g. age,
20 bibliometric indicators) and identify any funding attributions.
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5. Methods

Data extraction

The first step was to extract a list of publications from each of the guidelines and export them into a Microsoft Excel spreadsheet. This was performed automatically using bespoke RAND Europe computer scripts, based on the PERL scripting language. Here we briefly describe the methodology, since a full description is available elsewhere¹⁷. A total of 744 references were extracted from the dementia guideline and 446 for the COPD guideline.

Data cleaning

The extracted bibliographic references were cleaned and structured to permit analyses of funding source and paper performance indicators. Any references identified as non-academic or peer-reviewed publications (e.g. references to a website, grey literature), and all publications before 1980 were removed, since these could not be investigated using the Web of Science.

After extraction and initial cleaning, a total of 616 references were found for the dementia guideline (79.4% of the original 776 references) and 412 references for the COPD guideline (83.9% of the original 491 references).

Data processing

For the funding analyses, the extracted publications were searched for in Web of Science¹⁸ to find the institution and country affiliations of the authors listed. One aim was to identify the publications with at least one UK author on the assumption that this would facilitate further funding analysis. Another aim was to use all the extracted publications for further bibliometric analysis (see¹⁹ for a similar methodology).

From the dementia guideline, 494 out of the 616 extracted publications

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3 (80.1%) were found in the Web of Science. While from the COPD guideline, 335
4 out of 412 publications were available (81.3%). Any publications not found in the
5 Web of Science were processed individually through a search methodology
6 utilising the publication libraries of RAND Europe and Cambridge University. All
7 search processes were duplicated by a second researcher to eliminate errors.
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14 This methodology used a simple search of both the RAND library by article title,
15 using Google Scholar, and the Cambridge University online library and free access
16 journals, through the Google search engine. If the author affiliation and country
17 remained unidentified, then the RAND library was searched by journal, followed
18 by browsing for the article using the reference data available. In addition, the title
19 could be searched for by keyword within the journal. Finally, where possible,
20 Cambridge University print holdings were searched to find any articles that were
21 not accessible online.
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31 **Funder acknowledgement**

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33 Where publications had at least partial UK attribution, the funding source was
34 searched for in the Research Outputs Database (ROD)^{20, 21}. This is a database
35 housed at the Wellcome Trust recording the funding sources for UK and Irish
36 publications in the biomedical sciences for the period 1988–2001. Funding
37 acknowledgements were found for around one-third of publications using this
38 database. For the remaining publications, which fell outside of the appropriate
39 date range or were not found in the ROD, the full text of the publication was
40 found and funding acknowledgements recorded directly, where available.
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51 Funding source references were then standardised and categorised by broad
52 sector. UK funders acknowledged on cited papers were categorised into the
53 following categories: industry, not-for-profit, hospital trust, government
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3 department, government agency (not controlled by ministries), local or regional
4 authority, foundation, none given and unknown.
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8 **Bibliometric and paper characteristic analysis**

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10 Author affiliations and country locations were identified for 595 of the 616 (97%)
11 dementia guideline publications and 402 of the 412 (98%) COPD guideline
12 publications. These affiliated papers form the basis of the following descriptive
13 and bibliometric analysis.
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19 The citation impact of the publications referenced in both clinical guidelines was
20 analysed by sector using the concept of citation profiling. This is based on a
21 normalising technique called ReBased Impact (RBI), which takes account of the
22 field in which a paper appears and the date since its publication to effectively
23 provide a proxy measure for the 'quality' of each paper. The world average RBI is
24 1; the most highly cited articles have an RBI $>8^{22}$
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31 Whole counts were used throughout this analysis; if more than one funding
32 source is cited in a publication, this was recorded as *one* publication for each
33 funding source.
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6. Results

Attribution by funding organisation

In a large proportion of cited papers, no funding acknowledgement was listed.

Nearly half (104 of the 228; 46%) of publications in the dementia guideline with a UK author did not acknowledge any funder. Funding information was available for 117 papers, with the full text of 7 papers inaccessible and, hence, missing the funding information. For the 148 publications in the COPD guideline with at least one UK author, 60 included no funding acknowledgement (41%). Funding information was available for 81 publications. The full text of 7 publications was not accessible and therefore no funding data was available.

Examination of the funding acknowledgements for the Medical Research Council (MRC), the Department of Health (England), the NHS and the Wellcome Trust revealed that these funders were overtly linked to only a small proportion of papers cited in the guidelines (see figure 1 – Numbers of publications by funding source).

Attributions by funding organisation over time

To determine whether the practice of acknowledgement of funding has improved over time, funding acknowledgement was analysed by year of publication.

Although it appears that more recent publications have more complete funding acknowledgements than older ones, over the whole period, and for both guidelines, there was no clear statistical relationship between the age of publication and the presence or absence of a funding acknowledgement (see figure 2 – Funding acknowledgement by year).

The clinical guidelines, on the whole, cited recent research; the majority of research papers cited in these two UK clinical guidelines were published after

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3 2000, that is, within five years of the release of these guidelines. Although the
4 average duration between the publication date of papers cited and the publication
5 date of the citing guideline was 5 years for dementia and 3 years for COPD.
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8 9 10 **Attribution by funding sector**

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13 Industry was not as prominent a funding source for publications cited in the
14 dementia guideline, where acknowledgements were distributed across a range of
15 funding sources across sectors. In the COPD guideline, industry was the most
16 frequently linked funder after 'none given'.
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20 21 22 **Attribution by country**

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25 Of the 616 publications extracted from the dementia guideline, 228 (37.2%) had
26 at least one UK-based author; while from the COPD guideline, 148 publications
27 (35.9%) had at least one UK-based author. Researchers based in the UK and US
28 combined were linked to the majority of papers cited in both guidelines.
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36 Despite this dominance of the UK and USA-based researchers in the cited papers,
37 many other countries contributed to these publications. Papers cited in the
38 dementia and COPD guidelines were linked to authors from 37 and 36 countries,
39 respectively (see figure 3 – Contribution of the most active countries).
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43 44 45 **Attribution by research sector**

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47 The three research funders use different means of disbursing their money.
48 Funding includes grants to universities and hospitals, alongside direct support for
49 intramural research. Publication analysis by associated institutions revealed that
50 researchers with university addresses, followed by those with hospital addresses,
51 were linked to the bulk of papers cited on both guidelines. More than 80% of
52 publications cited in the two guidelines involved authors based at universities.
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3 The scientific contribution from other types of publicly funded institutions, as well
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5 as from non-profit institutions, was low.
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9 As described, nearly 20% of the publications cited in the COPD guideline involved
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11 authors from industry - a slightly higher proportion than in the Dementia
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13 guideline.
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15 16 17 **Citation quality**

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19 Clinical guideline drafting committees are obliged to base their recommendations
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21 on the 'best' research available. The citation impact of the publications referenced
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23 in both clinical guidelines was analysed by sector using the concept of citation
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25 profiling.
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29 Overall papers cited across both guidelines had high RBIs. Papers linked to
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31 universities, companies (industry) and publicly-funded research institutions were
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33 particularly highly cited. At the time of the analysis, for the COPD guideline, cited
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35 papers linked to publicly-funded research institutions had RBI > 8; for the
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37 dementia guideline papers linked to universities, companies and publicly-funded
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39 research institutions all had particularly impressive RBIs (see figure 4 – Citation
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41 score by institutional sector).
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7. Discussion

Funding attribution

Perhaps the greatest challenge in research impact assessment is dealing with attribution. Attribution of research outcomes and impacts to a specific funder is complex and, in many cases, improbable. This arises from most medical research receiving funding from multiple sources, involving a host of researchers (often across institutions) and being incremental such that considerable time elapses between original research and impact on health. The tide has turned on this issue and funding bodies are increasingly working together to identify where their funding has made a difference and contributed to an outcome or impact. Exclusively 'claiming' impact ignores the complexities and reality of scientific research and we are more interested in noting our contribution alongside others and learning from this.

Nevertheless, there is much we can do to help us better understand the connection between funding inputs and changes in medical practice. This research project was intended to flesh out some of the issues that we face in trying to link research funding to research output (i.e. research papers) and specific outcomes (clinical guidelines).

As described, there was some variability in the quality of acknowledgement information provided on papers. In our analysis, we did not explore whether the differences in acknowledgement quality and completeness varied according to the nature of the paper (i.e. underpinning versus more applied research). A study of 43 UK clinical guidelines (and associated Health Technology Assessments) related to cancer demonstrated that the number of funding sources acknowledged in papers varied with the 'basicness' of the publications: "*the more clinical papers have fewer (funding) sources and the more basic papers have more*"¹⁴.

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5 An interesting direction for future research work on clinical guidelines would be to
6 investigate whether the high proportion of cited publications with at least one UK
7 author in UK clinical guidelines could be the result of a specific funding strategy.
8 That is, some research could be being funded to help create evidence for the
9 development of clinical guidelines. This specific funding strategy might explain
10 partially why UK-authored publications are over-cited in UK guidelines, relative to
11 the share of publications.
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20 The advice given to funding recipients by UK funders regarding how they should
21 be acknowledged in publications, and the extent to which these requirements
22 have been enforced, has varied significantly over the last two decades. While
23 most funders have included a requirement for acknowledgement in their terms
24 and conditions of award, it has only been since 2008 that there has been
25 published guidance about a standard format²³. This may explain why industry is
26 fairly highly cited across both guidelines as part of a researcher's more stringent
27 contractual obligations. However, it is worth noting that the extent to which
28 researchers are following the standard format is unknown. Furthermore, the
29 reality remains that, given the incremental nature of much research, it is not easy
30 to precisely attribute a publication to its source of funding.
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44 One broader question is how to ensure that research information is accessible is
45 required in order to avoid spending funds on research that either cannot be used
46 or may be duplication²⁴.
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52 **Temporal issues**

53 We did find a correlation between the publication of the clinical guidelines and the
54 dates of the papers cited within it. These results corroborate earlier research¹⁴.
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3 Other bibliometric studies on UK clinical guidelines also found that a significant
4 share of publications cited are published within 10 years before the release of
5 these guidelines^{19,25}
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10 Therefore, this finding was not unexpected, since clinical guidelines should be
11 based on new evidence, although on occasion the newest evidence may not be
12 the best. However, it would be interesting to investigate whether the most recent
13 publications cited are the outcome of research specifically funded to support the
14 development of a clinical guideline, thus potentially explaining the peak in
15 publications cited in the two guidelines a few years before their release.
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21 22 **National contributions**

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24 The UK's contribution is high in both guidelines, since its world share in medical
25 research measured by its share in the total number of publications published in
26 the field amounted to approximately 8.6% in 2006²⁶. This high proportion of
27 papers linked to UK-based authors echoes the finding of previous studies.
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29 Interestingly, our analysis – where around a third of papers are linked to UK-
30 based authors - reveals a higher proportion than we have seen in other analyses.
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40 41 **Conclusions and recommendations**

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43 Having greater access to the work cited on clinical guidelines would present new
44 opportunities for funders and the research community alike to better understand
45 some of the mechanisms that take research from the lab to the bedside. While
46 this would be but one tool in the research impact evaluator's toolkit, it would be
47 one that could be relatively easy to harness if both access and acknowledgements
48 were improved. As described, work is underway between the UK NICE and the
49 NCBI, and in the future, it is envisioned that other guideline providers will make
50 their content available in much more structured and accessible formats to permit
51 analyses of this nature.
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5 However, to bring clarity to the tracking of research inputs through to published
6 output, the age-old problem of ensuring accurate and complete acknowledgement
7 information on peer-reviewed published work needs to be addressed. This
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9 requires a change in the culture of how researchers use acknowledgements and
10 greater liaison with publishers and information providers who could do much to
11 enhance the quality and completeness of funding data as a paper is submitted for
12 publication. Clarity and perhaps demarcation between the requirements of an
13 'acknowledgement' section on a paper and description of the 'funding' that has
14 supported the work would seem an easy step to help remedy this²⁷. This is
15 particularly pertinent as information providers such as Thomson Reuters and
16 Elsevier are now developing complex reporting and analytical tools that provide
17 an ability to scrutinise the characteristics of published work – including who is
18 described as funding the work - in new and exciting ways.
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32 Furthermore, if the methodology of this paper is to be generalised both beyond
33 formal guidelines and to other healthcare delivery and research output systems
34 and metrics, then novel methods of identifying and tracking researcher and their
35 outputs, such via global identifier systems such as that proposed by the ORCID²⁸
36 (Open and Researcher Contributor ID) initiative, will be important.
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44 Moves to address the definition of an 'acknowledgement' may also help to
45 address the issue of 'over-authorship' and author inflation that has been seen in
46 recent years. This is thought to be fuelled in part by the drive towards impact
47 assessment through national research allocation formulae; such as the UK
48 Research Assessment Exercise²⁹. Some contributors listed as 'authors' might be
49 more appropriately 'thanked'. Many journals now contain a section that asks for a
50 description of 'contributions' – these often do not contain all those listed in the
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3 author list so there may be a place for a more defined 'acknowledgement as
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5 thanks' section.
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9 Funders should also ensure clarity around their requirements for a 'funding
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11 acknowledgement'. Complexities arise for example, when research takes place in
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13 buildings funded by a specific donor or when a specific piece of research uses a
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15 piece of equipment funded by a specific donor. And for how long after funding has
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17 been received by a researcher should they continue to provide a funding
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19 acknowledgement?
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22 We found that there is great potential for national and international guidelines to
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24 be used as sources of information to help further our understanding on the
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26 impact of research on practice; the challenge is to be able to harness that
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28 information in an efficient way – so that we are able to use this information to
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30 feed into future research strategy and thereby make the research cycle more
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32 effective.
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12 13 **Contributorship**

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15 All authors (DK, LA, KD, BS and IV) contributed extensively to all parts of
16 this paper and commented on the manuscript at all stages. All authors were
17 involved in the design of the study with some guidance on direction and
18 feasibility from RAND Europe. All of the authors discussed the
19 interpretation of the results of the study, and agreed the outline content
20 for this publication. DK wrote the initial draft of the paper which was then
21 reviewed and further shaped by all of the authors. BS prepared the final
22 version of the paper plus all related information; **and managed the submission and**
23 **revision of this article.**
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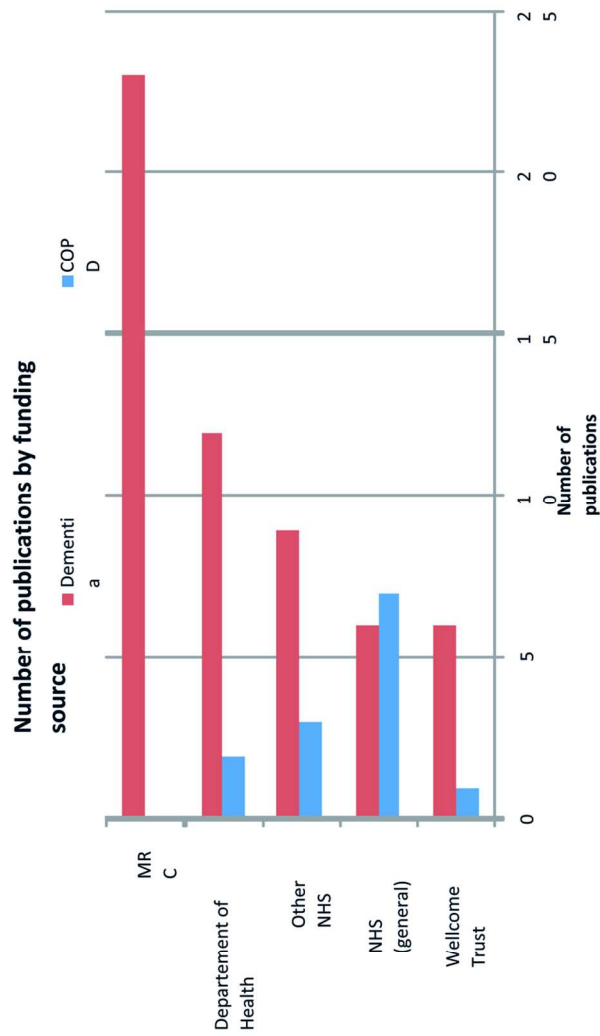
25 **Data sharing**

26 Unpublished data from this study is not currently available.
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28 **Funding**

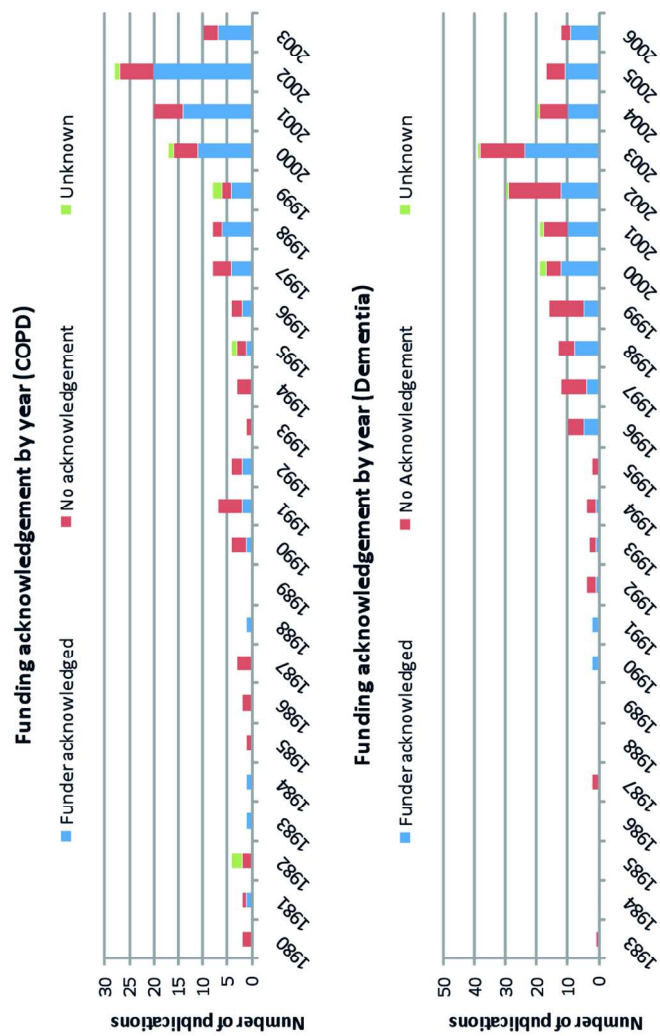
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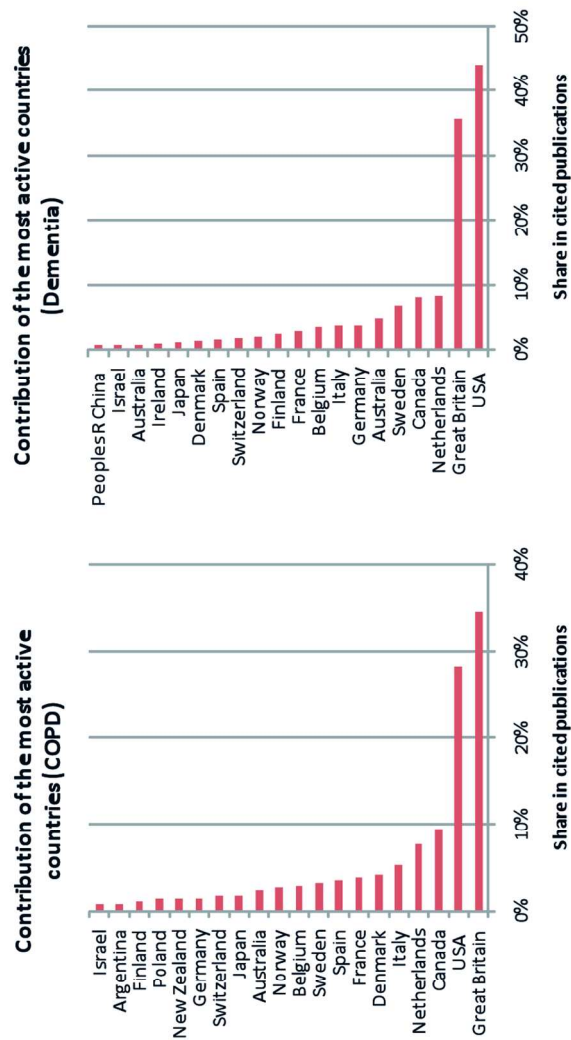
Numbers of publications by funding source.
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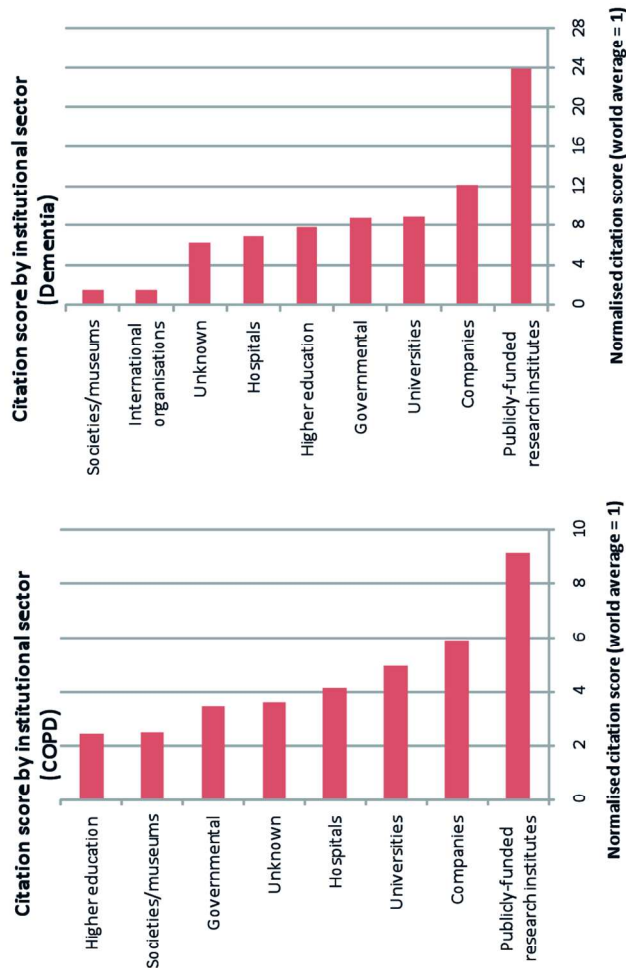
Funding acknowledgement by year.
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Contribution of the most active countries.
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ICMJE Form for Disclosure of Potential Conflicts of Interest

Instructions

The purpose of this form is to provide readers of your manuscript with information about your other interests that could influence how they receive and understand your work. The form is designed to be completed electronically and stored electronically. It contains programming that allows appropriate data display. Each author should submit a separate form and is responsible for the accuracy and completeness of the submitted information. The form is in four parts.

1. Identifying information.

Enter your full name. If you are NOT the corresponding author please check the box "no" and a space to enter the name of the corresponding author in the space that appears. Provide the requested manuscript information. Double-check the manuscript number and enter it.

2. The work under consideration for publication.

This section asks for information about the work that you have submitted for publication. The time frame for this reporting is that of the work itself, from the initial conception and planning to the present. The requested information is about resources that you received, either directly or indirectly (via your institution), to enable you to complete the work. Checking "No" means that you did the work without receiving any financial support from any third party -- that is, the work was supported by funds from the same institution that pays your salary and that institution did not receive third-party funds with which to pay you. If you or your institution received funds from a third party to support the work, such as a government granting agency, charitable foundation or commercial sponsor, check "Yes". Then complete the appropriate boxes to indicate the type of support and whether the payment went to you, or to your institution, or both.

3. Relevant financial activities outside the submitted work.

This section asks about your financial relationships with entities in the bio-medical arena that could be perceived to influence, or that give the appearance of potentially influencing, what you wrote in the submitted work. You should disclose interactions with ANY entity that could be considered broadly relevant to the work. For example, if your article is about testing an epidermal growth factor receptor (EGFR) antagonist in lung cancer, you should report all associations with entities pursuing diagnostic or therapeutic strategies in cancer in general, not just in the area of EGFR or lung cancer.

Report all sources of revenue paid (or promised to be paid) directly to you or your institution on your behalf over the 36 months prior to submission of the work. This should include all monies from sources with relevance to the submitted work, not just monies from the entity that sponsored the research. Please note that your interactions with the work's sponsor that are outside the submitted work should also be listed here. If there is any question, it is usually better to disclose a relationship than not to do so.

For grants you have received for work outside the submitted work, you should disclose support ONLY from entities that could be perceived to be affected financially by the published work, such as drug companies, or foundations supported by entities that could be perceived to have a financial stake in the outcome. Public funding sources, such as government agencies, charitable foundations or academic institutions, need not be disclosed. For example, if a government agency sponsored a study in which you have been involved and drugs were provided by a pharmaceutical company, you need only list the pharmaceutical company.

4. Other relationships.

Use this section to report other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work.

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Section 1. Identifying Information

1. Given Name (First Name) **BEVERLEY** 2. Surname (Last Name) **SHERBON** 3. Effective Date (07-August-2008) **21 - NOV - 2011**
4. Are you the corresponding author? Yes No
5. Manuscript Title **TRACKING THE IMPACT OF RESEARCH ON POLICY & PRACTICE : THE USE OF CLINICAL GUIDELINES IN RESEARCH EVALUATION.**
6. Manuscript Identifying Number (if you know it)

Section 2. The Work Under Consideration for Publication

Did you or your institution at any time receive payment or services from a third party for any aspect of the submitted work (including but not limited to grants, data monitoring board, study design, manuscript preparation, statistical analysis, etc...)?

Complete each row by checking "No" or providing the requested information. **If you have more than one relationship click the "Add" button to add a row. Excess rows can be removed by clicking the "X" button.**

The Work Under Consideration for Publication

Type	No	Money Paid to You	Money to Your Institution*	Name of Entity	Comments**	
1. Grant	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
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2. Consulting fee or honorarium	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
3. Support for travel to meetings for the study or other purposes	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
4. Fees for participation in review activities such as data monitoring boards, statistical analysis, end point committees, and the like	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
5. Payment for writing or reviewing the manuscript	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
6. Provision of writing assistance, medicines, equipment, or administrative support	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X



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The Work Under Consideration for Publication						
Type	No	Money Paid to You	Money to Your Institution*	Name of Entity	Comments**	
7. Other	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			ADD
						X
						ADD

* This means money that your institution received for your efforts on this study.

** Use this section to provide any needed explanation.

Section 3. Relevant financial activities outside the submitted work.

Place a check in the appropriate boxes in the table to indicate whether you have financial relationships (regardless of amount of compensation) with entities as described in the instructions. Use one line for each entity; add as many lines as you need by clicking the "Add +" box. You should report relationships that were present during the 36 months prior to submission.

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Relevant financial activities outside the submitted work						
Type of Relationship (in alphabetical order)	No	Money Paid to You	Money to Your Institution*	Entity	Comments	
1. Board membership	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
2. Consultancy	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
3. Employment	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
4. Expert testimony	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
5. Grants/grants pending	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
6. Payment for lectures including service on speakers bureaus	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
7. Payment for manuscript preparation	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X



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Relevant financial activities outside the submitted work

Type of Relationship (in alphabetical order)	No	Money Paid to You	Money to Your Institution*	Entity	Comments	
8. Patents (planned, pending or issued)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			ADD
						X
						ADD
9. Royalties	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
10. Payment for development of educational presentations	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
11. Stock/stock options	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
12. Travel/accommodations/meeting expenses unrelated to activities listed**	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD
13. Other (err on the side of full disclosure)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			X
						ADD

* This means money that your institution received for your efforts.

** For example, if you report a consultancy above there is no need to report travel related to that consultancy on this line.

Section 4. Other relationships

Are there other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work?

No other relationships/conditions/circumstances that present a potential conflict of interest

Yes, the following relationships/conditions/circumstances are present (explain below):

At the time of manuscript acceptance, journals will ask authors to confirm and, if necessary, update their disclosure statements. On occasion, journals may ask authors to disclose further information about reported relationships.

Hide All Table Rows Checked 'No'

SAVE



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