

SWAR 12: Contacting authors about additional study data – a randomised study comparing two strategies

Objective of this SWAR

To examine the effects of querying authors using personalised, study-specific questions in the email main text versus sending standardised data request forms as email attachments on response rate, data completeness, and reviewer time invested.

Study area: Data collection, Author contact

Sample type: Study authors

Estimated funding level needed: Unfunded

Background

Systematic review authors regularly come across eligible studies for which the description of methods is unclear or data are incomplete. Cochrane guidance recommends that reviewers contact study authors for clarification or to request additional data [1, 2]. In this context, review authors are advised to contact other authors if the corresponding author cannot be contacted or does not respond, to ask open-ended questions for descriptive information, and to provide a short data collection form (either uncompleted or partially completed) for specific data requests [1]. They are also encouraged to be judicious in the number of queries, considering the burden they impose on authors, to use user-friendly interfaces to display the data extracted to aid confirmation, and to be clear in describing the specific missing data being requested [3].

Previous empirical research has evaluated real or perceived barriers and facilitators to effective author contact. Barriers consist in uncertainties on how to conduct and report author contact in systematic reviews [3, 4], in difficulties locating study authors [4-6], and in authors not responding to emails, with observed response rates between 32 and 78% [4-8]. Contact with non-native speakers of English may be more difficult because of language barriers, but no evidence is available yet to support this supposition [7]. The number of items requested per author did not influence the probability of response [5]. Some authors found greater response rates when requesting clarification of methods than missing data [9], whereas others did not [7]. Finally, if conflicting data were obtained from the publications and from author contact, there was uncertainty on which source to trust [4, 10].

In terms of facilitators, email was a faster means of communication than postal mail (3 ± 3 versus 27 ± 30 days mean time to response), and generated more responses (hazard ratio: 2.5, 95% CI: 1.3 to 4.0) [5]. A combination of email with other methods, such as letter or telephone, was more effective than repeat emails, but also more time-consuming [5, 11]. Other reviewers received no additional data in response to telephone contact [7]. Short emails with the systematic review's protocol attached were as effective in eliciting a response as longer emails detailing the systematic review information [8]. Authors of more recent studies were more likely to be located and provide data [7]. The eminence of the email signatory did not elicit greater reply rates [12]. Previous research on the relative effectiveness of contacting corresponding versus other authors was inconclusive [9].

Our hypothesis is that the presentation of the data request may influence response rates and completeness of data provided. On one hand, study authors may perceive the workload to be lower when asked questions in the main text of an email than when receiving data collection forms as email attachments. A lower perceived workload might in turn elicit a higher response rate. On the other hand, review authors may anticipate a more complete response to standardised data collection forms than to free text in emails, because some items might be overlooked more easily in an email.

Therefore, we carried out a nested randomised controlled trial to investigate the effect of two different presentations of data requests to study authors. This study will be embedded in a systematic review of hospital volume-outcome relationships in total knee arthroplasty [13].

Interventions and comparators

Intervention 1: Personalised, study-specific questions in the email main text

Intervention 2: Structured data request form as an email attachment (partially pre-filled with available data for accuracy checking)

All authors of studies eligible for our systematic review (N=59 studies with 51 unique corresponding authors, potentially more after a search update) will be contacted by email, providing details of the systematic review including a brief description of research aims and the PROSPERO registration number and link to the PROSPERO record [13], as well as our contact details.

We will attempt to optimise response rates by searching for up-to-date author information online or in recent publications, by writing to non-responding authors in their mother tongue, and by contacting other authors if the corresponding author does not reply. We will send a maximum of four emails for each study, and the overall time to await replies will be limited to eight weeks. If study authors reply that they will send data later, we will send up to three email reminders to the same recipient if they do not send the data.

Access to all email correspondence and other study data will be limited to the SWAR team (Käthe Goossen, Tanja Rombey and Dawid Pieper). Results will be published as aggregate, anonymised data, so that personal identification of the study authors involved will not be possible.

Index Type:

Method for allocating to intervention or comparator

Randomisation

Outcome measures

Primary: Author reply received (yes/no, disregarding automated replies such as out-of-office messages)

Secondary: Complete data provided (yes/no, overall and stratified into (a) clarification of methods and (b) data missing in publication)

Time invested by reviewers in preparing the personalised emails and pre-filling the structured data request, as well as any follow-up communication with study authors (in hours).

Analysis plans

Response rates and data completeness will be compared using the odds ratio and corresponding 95% confidence interval; reviewer time invested will be compared using the mean and SD.

Possible problems in implementing this SWAR

Blinding of researchers will not be possible. To ensure that awareness of the study does not influence study authors in their decision on whether to reply and whether to provide data, we will contact them without making any reference to this SWAR. Blinding of outcome assessment was not considered necessary because the outcomes are objective.

The sample size is limited by the number of studies eligible in the systematic review. With 51 study authors, the study is powered to detect a 35-40% difference in reply rates between the groups (alpha=0.05, beta=0.20).

References

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Publications or presentations of this SWAR design

Käthe Goossen, Tanja Rombey, Charlotte M. Kugler, Karina K. De Santis, Dawid Pieper. Author queries via email text elicited high response and took less reviewer time than data forms - a randomised study within a review. *Journal of Clinical Epidemiology* (2021), <https://doi.org/10.1016/j.jclinepi.2021.02.006>

Examples of the implementation of this SWAR

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Date of idea: 15/JAN/2020

Revisions made by:

Date of revisions: