



ORIGINAL ARTICLE

Evaluation of reproducible and transparent research practices in pulmonology



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Abstract

Background: Study reproducibility is valuable for validating or refuting results. Provision of reproducibility indicators, such as materials, protocols, and raw data in a study improve its potential for reproduction. Efforts to reproduce noteworthy studies in the biomedical sciences have resulted in an overwhelming majority of them being found to be unreplicable, causing concern for the integrity of research in other fields, including medical specialties. Here, we analyzed the reproducibility of studies in the field of pulmonology.

Methods: 500 pulmonology articles were randomly selected from an initial PubMed search for data extraction. Two authors scoured these articles for reproducibility indicators including materials, protocols, raw data, analysis scripts, inclusion in systematic reviews, and citations by replication studies as well as other factors of research transparency including open accessibility, funding source and competing interest disclosures, and study preregistration.

Findings: Few publications included statements regarding materials (10%), protocols (1%), data (15%), and analysis script (0%) availability. Less than 10% indicated preregistration. More than half of the publications analyzed failed to provide a funding statement. Conversely, 63% of the publications were open access and 73% included a conflict of interest statement.

Interpretation: Overall, our study indicates pulmonology research is currently lacking in efforts to increase replicability. Future studies should focus on providing sufficient information regarding materials, protocols, raw data, and analysis scripts, among other indicators, for the sake of clinical decisions that depend on replicable or refutable results from the primary literature.

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Key messages**What is the key question?**

Are practices to improve study replicability and transparency being applied in pulmonology research?

What is the bottom line?

Current research in pulmonology is lacking in efforts to improve study replicability.

Why read on?

Study replicability is a fundamental aspect of the scientific method and practices to ensure this should be improved upon for the betterment of research that could eventually lead to clinical decisions.

Introduction

Reproducibility—the ability to duplicate a study’s results using the same materials and methods as the original investigator—is central to the scientific method.¹ Study reproducibility establishes confidence in the efficacy of therapies, while results that contradict original findings may lead to overturning previous standards. Herrera-Perez et al. recently evaluated 396 medical reversals in which suboptimal clinical practices were overturned when randomized controlled trials yielded results contrary to current practices.² Given the evolving nature of evidence-based patient care, studies must be conducted in a way that fosters reproducibility and transparency. Further, materials, protocols, analysis scripts, and patient data must be made available to enable verification.

Efforts supporting reproducibility are becoming more widespread owing to the open science movement. In 2013, the Center for Open Science was established to “increase the openness, integrity, and reproducibility of scientific research”.³ The center sponsored two large-scale reproducibility efforts: a series of 100 replication attempts in psychology and a series of 50 landmark cancer biology study replication attempts. In the first, investigators successfully reproduced only 39% of the original study findings.⁴ In the second, efforts were halted after only 18 replications because of lack of information and materials from authors, insufficient funding, and insufficient time to perform all the experiments.⁵ The center also created the Open Science Framework, a repository in which authors may deposit study protocols, participant data, analysis scripts, and other materials needed for study reproduction. More recently, the center created Transparency and Openness Promotion Guidelines, which include eight transparency standards and provides guidance for funders and journals, and initiated the use of badges for journals that adopt reproducible practices.

Current estimates of study reproducibility are alarming. In the biomedical sciences, reproducibility rates may be as low as 25%.⁶ One survey of 1576 scientists found that 90% of respondents believed science was experiencing a reproducibility crisis; 70% reported not being able to reproduce another investigator’s findings, and more than half reported an inability to reproduce their own findings.⁷ The picture is even less clear in the clinical sciences. Ioannidis found that of 49 highly cited original research publications, seven were refuted by newer studies, and seven suggested higher efficacy than follow-up results; only 22 were successfully replicated.⁸ The National Institutes of Health and the

National Science Foundation have responded to this crisis by taking measures to ensure that studies funded by tax dollars are more reproducible. However, little is known about the extent to which reproducibility practices are used in clinical research.

In this study, we evaluated reproducible and transparent research practices in the pulmonology literature.¹¹ Our goals were (i) to determine areas of strength and weakness in current use of reproducible and transparent research practices and (ii) to establish a baseline for subsequent investigations of the pulmonology literature.

Methods

This observational study employed a cross-sectional design. We used the methodology of Hardwicke et al.,¹¹ with modifications. In reporting this study, we follow the guidelines for meta-epidemiological methodology research⁹ and the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA).¹⁰ This study did not satisfy the regulatory definition for human subjects research as specified in the Code of Federal Regulations and therefore was not subject to institutional review board oversight. We have listed our protocol, materials, and data on Open Science Framework (<https://osf.io/n4yh5/>).

Journal and publication selection

The National Library of Medicine catalog was searched by DT using the subject terms tag “Pulmonary Medicine[ST]” to identify pulmonary medicine journals on May 29, 2019. To meet inclusion criteria, journals had to be published in English and be MEDLINE indexed. We obtained the electronic ISSN (or linking ISSN) for each journal in the NLM catalog meeting inclusion criteria. Using these ISSNs, we formulated a search string and searched PubMed on May 31, 2019, to locate publications published between January 1, 2014, to December 31, 2018. We then randomly selected 500 publications for data extraction using Excel’s random number function (<https://osf.io/zxjd9/>). We used OpenEpi version 3.0 to conduct a power analysis to estimate sample size. Data availability was the primary outcome due to its importance for study reproducibility.⁹ The population size of studies published in MEDLINE-indexed journals from which we selected our random sample was 299,255 with a hypothesized frequency of 18.5% for the factor in the population (which was based upon data obtained by Hardwicke et al.¹¹); a confidence limit of 5%; and a design factor of 1. Based on these assumptions, our study would require a sample size of 232. To allow for the attrition of publications that would not meet inclusion criteria, we randomly sampled a total of 500 publications. Previous investigations, upon which this study is based, have included random samples of 250 publications in the social sciences and 150 publications in the biomedical sciences.

Extraction training

Prior to data extraction, two investigators (JN, CS) underwent training to ensure inter-rater reliability. The training

included an in-person session to review the study design, protocol, Google form, and location of the extracted data elements in two publications. The investigators were next provided with three additional publications from which to extract data. Afterward, the pair reconciled differences by discussion. This training session was recorded and deposited online for reference (<https://osf.io/tf7nw/>). Prior to extracting data from all 500 publications, the two investigators extracted data from the first 10, followed by a final consensus meeting. Data extraction for the remaining 490 publications followed, and a final consensus meeting was held to resolve disagreements. A third author (DT) was available for adjudication, if necessary.

Data extraction

The two investigators extracted data from the 500 publications in a duplicate and blinded fashion. A pilot-tested Google form was created from Hardwicke et al.,¹¹ with additions (see [Table 1](#) for a description of the indicators of reproducibility and transparency). This form prompted coders to identify whether a study had important information that needed to be reproducible (<https://osf.io/3nfa5/>). The extracted data varied by study design. Studies without empirical data (e.g., editorials, commentaries [without reanalysis], simulations, news, reviews, and poems) had only the publication characteristics, conflict of interest statement, financial disclosure statement, funding sources, and open access availability. Non-Empirical studies do not have the expectation of being reproduced, and as such do not contain the indicators used for this study. Empirical studies included clinical trials, cohort, case series, secondary analysis, chart review, and cross-sectional. We catalogued the most recent year and 5-year impact factor of the publishing journals. Finally, we expanded the funding options to include university, hospital, public, private/industry, or nonprofit. In order to look more in-depth at areas of pulmonology research, the journal and sub-specialty of each empirical study was analyzed.

Verification of Open Access Status of publications

We used Open Access Button (<http://www.openaccessbutton.org>) to identify publications as being publicly available. Both the journal title and DOI were used in the search to mitigate chances of missing an article. If Open Access Button could not locate an article, we searched Google and PubMed to confirm open access status.

Publication citations included in research synthesis and replication

For empirical studies, Web of Science was used to identify whether the publication was replicated in other studies and had been included in systematic reviews and/or meta-analyses. To accomplish these tasks, two investigators (CS, JN) inspected the titles, abstracts, and introductions of all publications in which the reference study was cited. This process was conducted in a duplicate, blinded fashion.

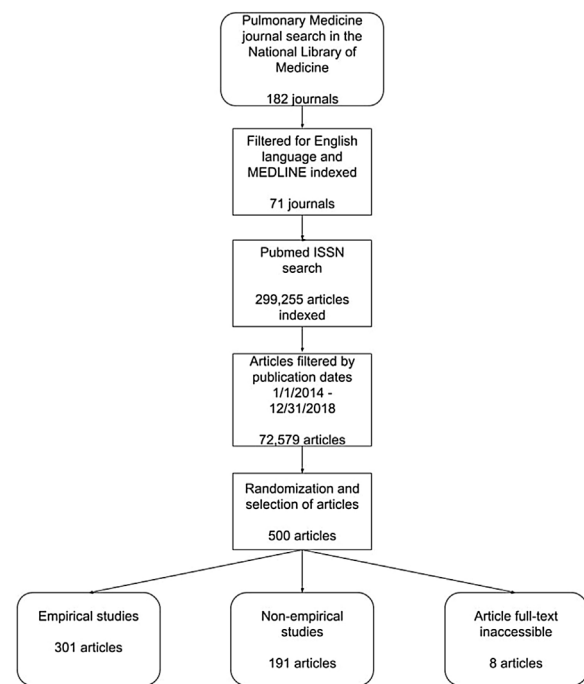


Figure 1 Article selection and filtering process.

Data analysis

We used Microsoft Excel to calculate descriptive statistics and 95% confidence intervals (95% CIs). The Wilson's Score for binomial proportions was used to create confidence intervals in this study.¹²

Role of the funding source

This study was funded through the 2019 Presidential Research Fellowship Mentor – Mentee Program at Oklahoma State University Center for Health Sciences. The funding source had no role in the study design, collection, analysis, interpretation of the data, writing of the manuscript, or decision to submit for publication.

Results

Study selection and article accessibility

Our PubMed search identified 299,255 publications. Limiting our search to articles published from January 1, 2014, to December 31, 2018, yielded 72,579 publications, from which 500 were randomly selected. Of these 500 publications, 312 were open access and 180 were behind a paywall. Eight publications could not be accessed by investigators and were thus excluded, leaving 492 for further analysis ([Fig. 1](#)). Characteristics of the included publications can be found in [Table 2](#).

Availability of reproducibility indicators

[Fig. 2](#) depicts an overview of our study results. A total of 238 empirical studies (excluding 56 case studies/case

Table 1 Indicators of reproducibility.

Reproducibility indicator	Number of studies	Role in producing transparent and reproducible science
Articles		
Availability of article (paywall or public access)	All (n = 500)	Widely accessible articles increase transparency
Funding		
Statement of funding sources	All included studies (n = 492)	Disclosure of possible sources of bias
Conflict of interest		
Statement of competing interests	All included studies (n = 492)	Disclosure of possible sources of bias
Evidence synthesis		
Citations in systematic reviews or meta-analyses	Empirical studies excluding systematic reviews and meta-analyses (n = 294)	Evidence of similar studies being conducted
Protocols		
Availability statement, and if available, what aspects of the study are included	Empirical studies excluding case studies (n = 245)	Availability of methods and analysis needed to replicate study
Materials		
Availability statement, retrieval method, and accessibility	Empirical studies excluding case studies and systematic reviews/meta-analysis (n = 238)	Defines exact materials needed to reproduce study
Raw data		
Availability statement, retrieval method, accessibility, comprehensibility, and content	Empirical studies excluding case studies (n = 245)	Provision of data collected in the study to allow for independent verification
Analysis scripts		
Availability statement, retrieval method, and accessibility	Empirical studies excluding case studies (n = 245)	Provision of scripts used to analyze data
Preregistration		
Availability statement, retrieval method, accessibility, and content	Empirical studies excluding case studies (n = 245)	Publicly accessible study protocol
Replication study		
Is the study replicating another study, or has another study replicated the study in question	Empirical studies excluding case studies (n = 245)	Evidence of replicability of the study

Bold values signifies to increase contrast between entries.

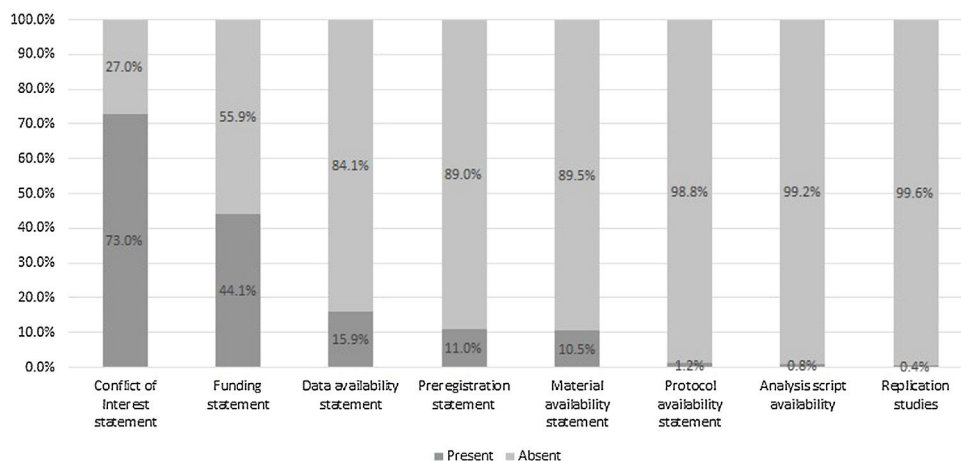


Figure 2 Proportion of studies with reproducibility indicators.

Table 2 Indicators of reproducibility in pulmonology studies.

Characteristics		N (%)	95% CI
Funding N = 492	University	8 (1.6)	0.5–2.7
	Hospital	5 (1.0)	0.1–1.9
	Public	65 (13.2)	10.2–16.2
	Private/industry	33 (6.7)	4.5–8.9
	Non-profit	11 (2.2)	0.9–3.5
	No statement listed	275 (55.9)	51.5–60.2
	Multiple sources	41 (8.3)	5.9–10.8
	Self-funded	1 (0.2)	0–0.6
	No funding received	53 (10.8)	8.0–13.5
Conflict of interest statement N = 492	Statement, one or more conflicts of interest	98 (19.9)	16.4–23.4
	Statement, no conflict of interest	261 (53.2)	48.8–57.5
	No conflict of interest statement	132 (26.9)	23.0–30.8
	Statement inaccessible	1 (0.2)	0–0.6
Data availability N = 245	Statement, some data are available	37 (15.1)	12.9–18.2
	Statement, data are not available	2 (0.8)	0–1.6
	No data availability statement	206 (84.1)	80.8–87.3
Material availability N = 238	Statement, some materials are available	24 (10.1)	7.4–12.7
	Statement, materials are not available	1 (0.4)	0–1.0
	No materials availability statement	213 (89.5)	86.8–92.2
Protocol availability N = 245	Full protocol	3 (1.2)	0.3–2.2
	No protocol	242 (98.8)	97.8–99.7
Analysis scripts N = 245	Statement, some analysis scripts are available	2 (0.8)	0–1.6
	Statement, analysis scripts are not available	0	0
	No analysis script availability statement	243 (99.2)	98.4–100
Replication studies N = 245	Novel study	244 (99.6)	99.0–100
	Replication	1 (0.4)	0–1.0
Cited by systematic review or meta-analysis N = 294	No citations	259 (88.1)	85.3–90.9
	A single citation	20 (6.8)	4.6–9.0
	One to five citations	14 (4.8)	2.9–6.6
	Greater than five citations	1 (0.3)	0–0.8
	Excluded in SR or MA	1 (0.3)	0–0.8
Preregistration N = 245	Statement present, preregistered	23 (9.4)	6.8–11.9
	Statement present, not pre-registered	4 (1.6)	0.5–2.7
	No preregistration statement	218 (89.0)	86.2–91.7
Frequency of reproducibility indicators in selected studies N = 301	Number of indicators present in study		
	0	49 (16.3)	–
	1	119 (39.5)	–
	2–5	133 (44.2)	–
	6–8	0	–
Open access N = 492	Found via open access button	215 (43.7)	39.4–48.0
	Yes-found article via other means	97 (19.7)	16.2–23.2
	Could not access through paywall	180 (36.6)	32.4–40.8

series, six meta-analyses, and one systematic review) were evaluated for material availability. The majority of studies offered no statement regarding availability of materials ($n=213$; 89.50% [95% CI, 86.81%–92.18%]). Twenty-four studies (10.08% [7.44%–12.72%]) had a clear statement regarding the availability of study materials. One study (0.42% [0%–0.99%]) included an explicit statement that the materials were not publicly available. Eighteen of the 24

materials files were accessible; the remaining six either led to a broken URL link or a pay-walled request form.

A total of 245 empirical studies (excluding 56 case studies/case series) were assessed for availability of protocols, raw data, and analysis scripts. Three studies provided access to a protocol (1.22% [0.26%–2.19%]). Data availability statements were more common, with 37 studies (15.10% [11.96%–18.24%]) including a statement that at least par-

tial data were available. Analysis scripts were found in two studies (0.82% [0.03%–1.61%]). More information on these metrics is presented in Supplemental Table 1 & 2.

Study preregistration

A total of 245 empirical studies (excluding 56 case studies/case series) were searched for a statement regarding study preregistration. Few studies included statements: 23 (9.39% [6.83%–11.94%]) declared preregistration, while four (1.63% [0.52%–2.74%]) explicitly disclosed that they were not preregistered. More information on preregistration is presented in Supplemental Table 1.

Study replication and citation analysis

Of 245 empirical studies analyzed, only one (0.41% [0%–0.97%]) reported replication of the methods of a previously published study. No studies were cited by a replication study. A total of 294 of the 301 empirical studies (excluding six meta-analyses and one systematic review) were evaluated to determine whether any had been included in a systematic review or meta-analysis. Twenty studies (6.80% [4.60%–9.01%]) were cited once in a systematic review or meta-analysis, 14 studies (4.76% [2.90%–6.63%]) were cited in two to five systematic reviews or meta-analyses, and one study (0.34% [0%–0.85%]) was cited in more than five systematic reviews or meta-analyses. One study (0.34% [0%–0.85%]) was explicitly excluded from a systematic review.

Conflict of interest and funding disclosures

All 492 publications were assessed for their inclusion of a conflict of interest statement and/or a funding statement. A majority ($n=359$; 73.08%) included a conflict of interest statement, with 261 declaring no competing interests (53.16% [48.78%–57.53%]). More than half of the publications failed to provide a funding statement ($n=275$; 55.89%; Table 2). In publications with a funding statement, public funding was the most common source ($n=65$; 13.21%).

Journal and sub-specialty characteristics

The total number of studies sampled from each journal is listed in Table 3 with the average number of reproducibility indicators with it. All 58 journals had at least one publication with empirical data and *The Annals of Thoracic Surgery* had the most with 33. The subspecialties of pulmonology are listed in Table 4 with the number of publications and average reproducibility indicators. Notable subjects were 102 in interventional pulmonology, 66 in obstructive lung disease, and 57 in critical care medicine. Publications over pulmonary hypertension averaged the most reproducibility indicators at 2.

Discussion

In this cross-sectional review of pulmonology publications, a substantial majority failed to provide materials,

Table 3 Number of studies per pulmonology subspecialty and mean number of reproducibility indicators.

Pulmonology subspecialty	Number of studies	Mean number of reproducibility indicators
Interventional pulmonology	102	0.98
Tobacco treatment	3	1
Lung transplantation	8	1.13
Sarcoidosis	4	1.25
Neuromuscular disease	3	1.33
Cystic fibrosis	9	1.44
Critical care medicine	57	1.47
Lung cancer	27	1.52
Obstructive lung disease	66	1.76
Interstitial lung disease	9	1.78
Sleep medicine	9	1.89
Pulmonary hypertension	4	2

participant data, or analysis scripts. Many were not preregistered and few had an available protocol. Reproducibility has been viewed as an increasingly troublesome area of study methodology.¹³ Recent attempts at reproducing preclinical^{14,15} and clinical studies have found that only 25%–61% of studies may be successfully reproduced.^{6,16} Within the field of critical care medicine, a recent publication found that only 42% of randomized trials contained a reproduction attempt with half of those reporting inconsistent results compared to the original.¹⁷ Many factors contribute to limited study reproducibility, including poor (or limited) reporting of study methodology, prevalence of exaggerated statements, and limited training on experimental design in higher education.¹⁸ In an effort to limit printed pages and increase readability, journals may request that authors abridge methods sections.¹⁹ Here, we briefly comment on selected indicators to present a balanced view of the perspectives of those in favor of reproducibility and transparency and those who resist enacting such changes.

First, data sharing allows for the independent verification of study results or reuse of that data for subsequent analyses. Two sets of principles exist. The first, known as FAIR, outlines mechanisms for findability, accessibility, interoperability, and reusability. FAIR principles are intended to apply to study data as well as the algorithms, tools, and workflows that led to the data. FAIR advocates that data be accessible to the right people, in the right way, and at the right time.²⁰ A second set of principles relate to making data available to the public for access, use, and share without licenses, copyrights, or patents.²¹ While we advocate for data sharing, we recognize that it is a complex issue. First, the process for making data available for others' use requires skills. Further, the process, which includes the construction of data dictionaries and data curation, is time consuming. Furthermore, concerns exist with regard to unrestricted access to data facilitating a culture of "research parasites," a term coined by Drazen and Longo²² that suggests that secondary researchers might exploit primary research data for publication. Drazen and Longo also cautioned that secondary authors might not understand the decisions made when

Table 4 Number of studies per journal and mean number of reproducibility indicators.

Journal title	Number of studies	Mean number of reproducibility indicators
Journal of cardiothoracic and vascular anesthesia	12	0.25
The annals of thoracic surgery	33	0.33
Respiration; international review of thoracic diseases	3	0.67
Respirology	3	0.67
The thoracic and cardiovascular surgeon	7	0.86
Respiratory investigation	1	1
Annals of thoracic and cardiovascular surgery	4	1
Canadian respiratory journal	2	1
Seminars in thoracic and cardiovascular surgery	1	1
Jornal Brasileiro de pneumologia	2	1
Journal of thoracic imaging	1	1
Respiratory care	3	1
Current allergy and asthma reports	1	1
European respiratory review	1	1
General thoracic and cardiovascular surgery	2	1
Journal of bronchology & interventional pulmonology	6	1.17
Annals of the American thoracic society	12	1.7
Respiratory physiology & neurobiology	4	1.25
Thoracic cancer	4	1.25
The journal of heart and lung transplantation	4	1.25
Heart, lung & circulation	7	1.29
The European respiratory journal	7	1.29
Lung	3	1.33
Pulmonary pharmacology & therapeutics	3	1.33
Journal of cystic fibrosis	6	1.33
American journal of physiology. lung cellular and molecular physiology	3	1.33
Interactive cardiovascular and thoracic surgery	8	1.38
The journal of thoracic and cardiovascular surgery	13	1.385
The clinical respiratory journal	5	1.4
Tuberculosis (Edinburgh, Scotland)	5	1.4
Journal of breath research	5	1.4
Experimental lung research	2	1.5
The journal of asthma	4	1.5
Clinical lung cancer	2	1.5
The international journal of tuberculosis and lung disease	5	1.6
Respiratory medicine	5	1.6
Chest	5	1.6
European journal of cardio-thoracic surgery	9	1.67
Thorax	7	1.71
Journal of cardiothoracic surgery	4	1.75
Respiratory research	9	1.78
Annals of allergy, asthma & immunology	10	1.8
American journal of respiratory and critical care medicine	5	1.8
Asian cardiovascular & thoracic annals	8	2
Journal of cardiopulmonary rehabilitation and prevention	1	2
Heart & lung : the journal of critical care	1	2
Sleep & breathing	7	2
Allergy and asthma proceedings	4	2
Chronic respiratory disease	2	2
International journal of chronic obstructive pulmonary disease	10	2
Multimedia manual of cardiothoracic surgery	1	2
The Lancet. Respiratory medicine	4	2
Pediatric pulmonology	8	2.13
Journal of aerosol medicine and pulmonary drug delivery	5	2.2

Table 4 (Continued)

Journal title	Number of studies	Mean number of reproducibility indicators
BMC pulmonary medicine	4	2.25
Journal of thoracic oncology	5	2.8
NPJ primary care respiratory medicine	1	3
COPD	2	3.5

defining parameters of the original investigations. Finally, the sensitive nature of some data causes concern among researchers.

Second, preregistering a study requires authors to provide their preliminary protocol, materials, and analysis plan in a publicly available website. The most common websites used by authors are ClinicalTrials.gov and the International Clinical Trial Registry Platform hosted by the World Health Organization. These registries improve the reliability and transparency of published findings by preventing selective reporting of results, preventing unnecessary duplication of studies, and providing relevant material to patients that may enroll in such trials.²³ The Food and Drug Administration (FDA) Amendments Act and the International Committee of Medical Journal Editors (ICMJE) have both required registration of clinical trials prior to initiation of a study.^{24,25} Selective reporting bias, which includes demoting primary endpoints, omitting endpoints, or upgrading secondary endpoints in favor of statistical significance, may be especially pervasive and problematic. Numerous studies across several fields of medicine have evaluated the extent and magnitude of the problem.^{26–28} The consequences of selective reporting bias and manipulation of endpoints may compromise clinical decision making. Another issue—*p*-hacking—occurs when researchers repeatedly analyze study data until they achieve statistically significant results. Pre-registration of protocols and statistical analysis plans can be used to fact check published papers to ensure that any alterations made in the interim were made for good reason.

Third, transparency related to study funding and financial conflicts of interest should be emphasized. In a previous study, we found that one-third of the authors of pivotal oncology trials underlying FDA drug approvals failed to adequately disclose personal payments from the drug sponsor.²⁹ Recent news accounts of a prominent breast cancer researcher who failed to disclose financial relationships with pharmaceutical companies in dozens of publications has heightened awareness of the pervasiveness of this issue.³⁰ The ICMJE considers willful nondisclosure of financial interests to be a form of research misconduct.³¹ It is critical that the public be able to adequately evaluate financial relationships of the authors of the published studies in order to evaluate the likelihood of biased results and conclusions.

Several changes are needed to establish a culture of reproducibility and transparency. First, increased awareness of and training about these issues are needed. The National Institutes of Health has funded researchers to produce training and materials, which are available on the

Rigor and Reproducibility Initiative website,³² but more remains to be done. Strong mentorship is necessary to encourage trainees to adopt and incorporate reproducible research practices. Research on mentorship programs has found that trainees who have mentors report greater satisfaction with time allocation at work and increased academic self-efficacy compared with trainees without a mentor.³³ Conversely, poor mentorship can reinforce poor research practices among junior researchers, such as altering data to produce positive results or changing how results are reported.³⁴ Other research stakeholders must be involved as well. Although many journals recommend the use of reporting guidelines for various study designs, such as CONSORT and PRISMA, evidence suggests that these guidelines are not followed by authors or enforced by journals.³⁵ When journals enforce adherence to reporting guidelines, the completeness of reporting is improved.³⁶ Detractors of reporting guidelines are concerned that certain checklists (CONSORT, STROBE, STARD) will be used to judge research quality rather than improve writing clarity, that editors and peer reviewers will fail to enforce these guidelines, and that insufficient research exists to evaluate the outcomes from applying these guidelines.³⁷ We analyzed *COPD*, *NPJ Primary Care Respiratory Medicine*, and *Journal of Thoracic Oncology* from our sample as the top three journals for containing reproducibility indicators in their publications. These journals have explicit instructions for authors to provide things such as materials/protocols such that independent researchers may recreate the study or raw data to confirm calculations.^{38–40} Although reproducibility may be an emerging topic, these recommendations appear to be encouraging authors to include more thorough and complete research.

Our study has both strengths and limitations. We randomly sampled a large number of pulmonology journals containing various types of publications to generalize our findings across the specialty. Our study design also used rigorous training sessions and a standardized protocol to increase the reliability of our results. In particular, our data extraction process, which involved blinded and duplicate extraction by two investigators, is the gold standard systematic review methodology and is recommended by the Cochrane Collaboration.⁴¹ We have made all study materials available for public review to enhance the reproducibility of this study. Regarding limitations, our inclusion criteria for journals (i.e., published in English and MEDLINE indexed) potentially removed journals that contained more lax recommendations regarding indicators of reproducibility and transparency. Furthermore, although we obtained a random sample of publications for analysis, our sample may not have been representative of all pulmonology publications. Our

results should be interpreted in light of these strengths and limitations.

In conclusion, our study of the pulmonology literature found that reproducible and transparent research practices are not being incorporated into research. Sharing of study artifacts, in particular, needs improvement. The pulmonology research community should seek to establish norms of reproducible and transparent research practices.

Author contributions

DJT, MV: Substantial contributions to the conception and design of the work. CAS, JN, DJT: Acquisition, analysis, and interpretation of data for the work. CAS, JN, DJT, TEH, JP, KC, MV: Drafted the work and revised it critically for important intellectual content. MV: Final approval of the version submitted for publication. CAS: Accountability for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Conflicts of interest

The authors have no conflicts of interest to declare.

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Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.pulmoe.2020.07.001>.

References

- Goodman SN, Fanelli D, Ioannidis JPA. What does research reproducibility mean? *Sci Transl Med.* 2016;8(June (341)):341ps12, <http://dx.doi.org/10.1126/scitranslmed.aaf5027>.
- Herrera-Perez D, Haslam A, Crain T, Gill J, Livingston C, Kaestner V, et al. Meta-Research: a comprehensive review of randomized clinical trials in three medical journals reveals 396 medical reversals. *Elife.* 2019;8:e45183, <http://dx.doi.org/10.7554/eLife.45183>.
- Triple W, Center for Open Science. <https://cos.io/about/mission/>, 2006 [Accessed July 2019].
- Anderson CJ, Anderson J, van Assen MALM, Attridge PR, Attwood A, Axt J, et al. Reproducibility Project: Psychology. *Open Science Framework*; 2018, <http://dx.doi.org/10.17605/OSF.IO/EZCUJ>.
- Center for Open Science. <https://osf.io/e81xl/wiki/home/>, 2013 [Accessed 17 June 2019].
- Prinz F, Schlange T, Asadullah K. Believe it or not: how much can we rely on published data on potential drug targets? *Nat Rev Drug Discov.* 2011;10(August (9)):712, <http://dx.doi.org/10.1038/nrd3439-c1>.
- Baker M. 1,500 scientists lift the lid on reproducibility. *Nature.* 2016;533(May (7604)):452–4, <http://dx.doi.org/10.1038/533452a>.
- Ioannidis JPA. Contradicted and initially stronger effects in highly cited clinical research. *JAMA.* 2005;294(July (2)):218–28, <http://dx.doi.org/10.1001/jama.294.2.218>.
- Murad MH, Wang Z. Guidelines for reporting meta-epidemiological methodology research. *Evid Based Med.* 2017;22(August (4)):139–42, <http://dx.doi.org/10.1136/ebmed-2017-110713>.
- Liberati A, Altman DG, Tetzlaff J, Mulrow C, Gøtzsche PC, Ioannidis JPA, et al. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. *J Clin Epidemiol.* 2009;62(October (10)):e1–34, <http://dx.doi.org/10.1136/bmj.b2700>.
- Hardwicke TE, Wallach JD, Kidwell MC, Bendixen T, Crüwell S, Ioannidis JPA. An empirical assessment of transparency and reproducibility-related research practices in the social sciences (2014–2017). *R Soc Open Sci.* 2020;7(February (2)):190806, <http://dx.doi.org/10.1098/rsos.190806>.
- Dunnigan K. Confidence interval calculation for binomial proportions. In: MWSUG Conference. 2008. <http://www.mwsug.org/proceedings/2008/pharma/MWSUG-2008-P08.pdf> [Accessed 7 June 2020].
- Nosek BA, Spies JR, Motyl M. Scientific Utopia: II. restructuring incentives and practices to promote truth over publishability. *Perspect Psychol Sci.* 2012;7(November (6)):615–31, <http://dx.doi.org/10.1177/1745691612459058>.
- Munafò MR. Reliability and replicability of genetic association studies. *Addiction.* 2009;104(September (9)):1439–40, <http://dx.doi.org/10.1111/j.1360-0443.2009.02662.x>.
- Button KS, Ioannidis JPA, Mokrysz C, Nosek BA, Flint J, Robinson ESJ, et al. Power failure: why small sample size undermines the reliability of neuroscience. *Nat Rev Neurosci.* 2013;14(May (5)):365–76, <http://dx.doi.org/10.1038/nrn3475>.
- Open Science Collaboration, PSYCHOLOGY. Estimating the reproducibility of psychological science. *Science.* 2015;349(August (6251)):aac4716, <http://dx.doi.org/10.1126/science.aac4716>.
- Niven DJ, McCormick TJ, Straus SE, Hemmelgarn BR, Jeffs L, Barnes TRM, et al. Reproducibility of clinical research in critical care: a scoping review. *BMC Med.* 2018;16(February (1)):26, <http://dx.doi.org/10.1186/s12916-018-1018-6>.
- Carp J. The secret lives of experiments: methods reporting in the fMRI literature. *Neuroimage.* 2012;63(October (1)):289–300, <http://dx.doi.org/10.1016/j.neuroimage.2012.07.004>.
- Marcus E, Whole Cell team. A STAR is born. *Cell.* 2016;166(August (5)):1059–60, <http://dx.doi.org/10.1016/j.cell.2016.08.021>.
- Wilkinson MD, Dumontier M, Aalbersberg IJJ, Appleton G, Axton M, Baak A, et al. The FAIR Guiding Principles for scientific data management and stewardship. *Sci Data.* 2016;3(March):160018, <https://doi.org/10.1038/sdata.2016.18>.
- Mons B, Neylon C, Velterop J, Dumontier M, da Silva Santos L, Wilkinson MD. Cloudy, Increasingly FAIR; Revisiting the FAIR Data Guiding Principles for the European Open Science Cloud. *Inf Serv Use.* 2017;37(January 1):49–56, <http://dx.doi.org/10.3233/ISU-170824>.
- Longo DL, Drazen JM. More on data sharing. *N Engl J Med.* 2016;374(May (19)):1896–7, <http://dx.doi.org/10.1056/nejmc1602586>.
- ICMJE. <http://www.icmje.org/icmje-recommendations.pdf>, 2016 [Accessed July 2019].
- Food and Drug Administration Amendments Act of 2007, Pub. L. No. 110-185, 121 Stat. 823 (Sep 27, 2007).

25. Laine C, Horton R, DeAngelis CD, Drazen JM, Frizelle FA, Godlee F, et al. Clinical trial registration — looking back and moving ahead. *N Engl J Med*. 2007;356(June):2734–6, <http://dx.doi.org/10.1056/nejme078110>.
26. You B, Gan HK, Pond G, Chen EX. Consistency in the analysis and reporting of primary end points in oncology randomized controlled trials from registration to publication: a systematic review. *J Clin Oncol*. 2012;30(January (2)):210–6, <http://dx.doi.org/10.1200/jco.2011.37.0890>.
27. Mathieu S, Boutron I, Moher D, Altman DG, Ravaud P. Comparison of registered and published primary outcomes in randomized controlled trials. *JAMA*. 2009;302(September (9)):977–84, <http://dx.doi.org/10.1001/jama.2009.1242>.
28. Rankin J, Ross A, Baker J, O'Brien M, Scheckel C, Vassar M. Selective outcome reporting in obesity clinical trials: a cross-sectional review. *Clin Obes*. 2017;7(4):245–54, <http://dx.doi.org/10.1111/cob.12199>.
29. Wayant C, Turner E, Meyer C, Sinnett P, Vassar M. Financial conflicts of interest among oncologist authors of reports of clinical drug trials. *JAMA Oncol*. 2018;4(October(10)):1426–8, <http://dx.doi.org/10.1001/jamaoncol.2018.3738>.
30. Ornstein C, Thomas K, The New York Times. <https://www.nytimes.com/2018/09/08/health/jose-baselga-cancer-memorial-sloan-kettering.html>, 2018. September 8 [Accessed 26 June 2019].
31. ICMJE. Disclosure of Financial and Non-Financial Relationships and Activities, and Conflicts of Interest, <http://icmje.org/recommendations/browse/roles-and-responsibilities/author-responsibilities-conflicts-of-interest.html>. [Accessed 26 June 2019].
32. National Institutes of Health (NIH). <https://www.nih.gov/research-training/rigor-reproducibility/training>, 2015 [Accessed 26 June 2019].
33. Feldman MD, Arean PA, Marshall SJ, Lovett M, O'Sullivan P. Does mentoring matter: results from a survey of faculty mentees at a large health sciences university. *Med Educ Online*. 2010;15(April 1), <http://dx.doi.org/10.3402/meo.v15i0.5063>.
34. Boulbes DR, Costello T, Baggerly K, Fan F, Wang R, Bhat-tacharya R, et al. A survey on data reproducibility and the effect of publication process on the ethical reporting of laboratory research. *Clin Cancer Res*. 2018;24(July (14)):3447–55, <http://dx.doi.org/10.1158/1078-0432.ccr-18-0227>.
35. Sims MT, Checketts JK, Wayant C, Vassar M. Requirements for trial registration and adherence to reporting guidelines in critical care journals: a meta-epidemiological study of journals' instructions for authors. *Int J Evid Based Healthc*. 2018;16(March (1)):55–65, <http://dx.doi.org/10.1097/xeb.000000000000120>.
36. Agha RA, Fowler AJ, Limb C, Whitehurst K, Coe R, Sagoo H, et al. Impact of the mandatory implementation of reporting guidelines on reporting quality in a surgical journal: a before and after study. *Int J Surg*. 2016;30(June):169–72, <http://dx.doi.org/10.1016/j.ijvsu.2016.04.032>.
37. Vandembroucke JP. STREGA, STROBE, STARD, SQUIRE, MOOSE, PRISMA, GNOSIS, TREND, ORION, COREQ, QUOROM, REMARK... and CONSORT: for whom does the guideline toll? *J Clin Epidemiol*. 2009;62(June (6)):594–6, <http://dx.doi.org/10.1016/j.jclinepi.2008.12.003>.
38. Journal of Chronic Obstructive Pulmonary Disease. Submit to COPD, <https://www.tandfonline.com/action/authorSubmission?show=instructions&journalCode=icop20>. [Accessed 7 June 2020].
39. Npj Primary Care Respiratory Medicine. For Authors & Referees, <https://www.nature.com/npjpcrm/authors-and-referees>. [Accessed 7 June 2020].
40. Information for Authors: Journal of Thoracic Oncology. Author Information, <https://www.jto.org/content/authorinfo>. [Accessed 7 June 2020].
41. Higgins JPT, Deeks JJ. Selecting studies and collecting data. In: Higgins JPT, Green S, editors. *Cochrane handbook for systematic reviews of interventions*. Chichester (UK): John Wiley & Sons; 2008. p. 151–85.