



Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.

ORIGINAL ARTICLE

# Abstracts for reports of randomized trials of COVID-19 interventions had low quality and high spin

Dongguang Wang<sup>a,1</sup>, Lingmin Chen<sup>b,1</sup>, Lian Wang<sup>a</sup>, Fang Hua<sup>c,d</sup>, Juan Li<sup>e</sup>, Yuxi Li<sup>e</sup>,  
Yonggang Zhang<sup>f</sup>, Hong Fan<sup>a,\*</sup>, Weimin Li<sup>a,\*\*</sup>, Mike Clarke<sup>g,\*\*\*</sup>

<sup>a</sup>Department of Respiratory and Critical Care Medicine, West China Hospital/West China School of Medicine, Sichuan University, Chengdu, China

<sup>b</sup>Department of Anesthesiology and National Clinical Research Center for Geriatrics, West China Hospital, Sichuan University & The Research Units of West China (2018RU012, Chinese Academy of Medical Sciences), Chengdu, China

<sup>c</sup>Center for Evidenced-Based Stomatology, School & Hospital of Stomatology, Wuhan University, Wuhan, China

<sup>d</sup>Cochrane Oral Health, Division of Dentistry, School of Medical Sciences, Faculty of Biology, Medicine and Health, University of Manchester, Manchester Academic Health Science Center, Manchester, UK

<sup>e</sup>School of Health Preservation and Rehabilitation, Chengdu University of Traditional Chinese Medicine, Chengdu, China

<sup>f</sup>Department of Periodical Press, West China Hospital, Sichuan University, Chengdu, China

<sup>g</sup>Northern Ireland Clinical Trials Unit and Methodology Hub, Centre for Public Health, Queen's University Belfast, Belfast, UK

Accepted 29 June 2021; Available online 3 July 2021

## Abstract

**Objectives:** To assess the reporting quality of abstracts for published randomized controlled trials (RCTs) of interventions for coronavirus disease 2019 (COVID-19), including the use of spin strategies and the level of spin for RCTs with statistically nonsignificant primary outcomes, and to explore potential predictors for reporting quality and the severity of spin.

**Study Design and Setting:** PubMed was searched to find RCTs that tested interventions for COVID-19, and the reporting quality and spin in the abstracts were assessed. Linear regression analyses were used to identify potential predictors.

**Results:** Forty RCT abstracts were included in our assessment of reporting quality, and a higher word count in the abstract was significantly correlated with higher reporting scores (95% CI 0.044–0.658,  $P = 0.026$ ). Multiple spin strategies were identified. Our multivariate analyses showed that geographical origin was associated with severity of spin, with research from non-Asian regions containing fewer spin strategies (95% CI -0.756 to -0.096,  $P = 0.014$ ).

**Conclusions:** The reporting quality of abstracts of RCTs of interventions for COVID-19 is far from satisfactory. A relatively high proportion of the abstracts contained spin, and the findings reported in the results and conclusion sections of these abstracts need to be interpreted with caution. © 2021 Elsevier Inc. All rights reserved.

**Keywords:** Abstract; COVID-19; Randomized controlled trial; Primary outcome; Reporting quality; Spin

**Abbreviations:** RCT, randomized controlled trial; COVID-19, coronavirus disease 2019; CONSORT, consolidated standards of reporting trials; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2; IQR, interquartile range.

Declaration of competing interest: The authors have stated explicitly that they have no conflicts of interest in connection with this article.

<sup>1</sup> These authors contributed equally to this work.

\* Corresponding author. Tel.: +86 28 8542 3331; fax: +86 28 8542 3331.

\*\* Corresponding author. Tel.: +86 28 8542 3001; fax: +86 28 8542 3001.

\*\*\* Corresponding author. Tel.: +44-28-90978935; fax: +44-28-90235900.

E-mail addresses: fanhongfan@qq.com (H. Fan), weimi003@yahoo.com (W. Li), m.clarke@qub.ac.uk (M. Clarke).

## What is new?

### Key findings

- The median reporting score of 40 abstracts presenting the results of RCTs of interventions for COVID-19 was 8 (6, 10) of the 16 items in the CONSORT statement for abstracts.
- Fourteen (52%) of 27 abstracts with statistically nonsignificant primary outcomes had spin in the results section and 12 (44%) abstracts had spin in the conclusion section.
- Multivariate analyses showed that word count in the abstract was significantly correlated with reporting scores ( $P = 0.026$ ) and geographical origin was associated with severity of spin ( $P = 0.014$ ).

### What this adds to what is known?

- The reporting quality of abstracts of RCTs of 19 interventions for COVID-19 is low, and a relatively high proportion of the abstracts contains spin.
- This study explores potential predictors for reporting quality and the severity of spin.

### What is the implication?

- We recommend that authors, peer reviewers and editors make better use of reporting guidelines such as CONSORT and its extensions when preparing, appraising and editing research articles. Allowing authors more words in their abstracts might be a simple method to improve the reporting quality of abstracts.
- The findings reported in the results and conclusion sections of these abstracts need to be interpreted with caution, especially for those studies from Asian regions.

## 1. Introduction

In December 2019, a novel RNA coronavirus, which was subsequently named severe acute respiratory syndrome coronavirus (SARS-CoV-2) caused an outbreak of pneumonia in Hubei province of China, and quickly transmitted to other countries, resulting in millions of infections and deaths globally in just a few months. Unfortunately, there were no efficient methods to kill this virus or to treat the coronavirus disease 2019 (COVID-19) it caused. To address this, a large number of randomized controlled trials (RCTs) emerged very quickly to test interventions to prevent or treat COVID-19, along with a vast number of other pieces of research. This rapid, unprecedented outpouring of research on a specific condition means that policy makers, patients and clinicians are under great pressure to identify the most useful and reliable information from amidst

an overwhelming number of articles. They need relevant and valid material which can be accessed quickly and with minimal efforts [1], and as a result, the abstracts of the full reports become key to supporting clinical decision-making [2]. Clear reporting of a study's results in its abstract is likely to aid rational decision-making, but selective outcome reporting and other biases in the abstracts may make readers disoriented. It will be also important for users of these RCTs to consider whether the studies were justified, given that this has been identified as a problem for RCTs in the past [3].

To help readers, the consolidated standards of reporting trials (CONSORT) statement, published in 2010, lists 16 essential items for inclusion in the abstracts for reports of RCTs [4]. This provides the requirements for the content of a well reported abstract, but selectivity in how the authors present their results remains a problem, especially in studies without statistically significant outcomes where the authors might use spin to try to highlight results inappropriately.

The concept of spin was originally used in fields such as culture, politic and commerce, and is defined as a form of propaganda that could affect public views on an organization or public figure [5]. In health care, it was defined by Boutron et al. in 2010 as “specific reporting strategies from whatever motive, highlighting the interventions are beneficial despite the statistically nonsignificant differences for the primary outcomes” [6]. Since then, it has been widely evaluated in several medical specialties [7], including stomatology [5,8], otolaryngology [9], surgery [10,11], and cardiology [12]. Considering that such a large number of COVID-19 articles are being published with shorter periods for peer review and editorial oversight, this raises the possibility that more spin is making its way into the literature and that readers are being misled. Therefore, we have undertaken what we believe to be the first study of spin in reports of RCTs of interventions for COVID-19. This study investigates (1) the reporting quality of the abstracts for published RCTs of interventions for COVID-19 and their use of spin strategies, and the extent and level of spin in abstracts with statistically nonsignificant primary outcomes, and (2) potential predictors for reporting quality and the severity of spin.

## 2. Materials and methods

### 2.1. Search, eligibility, and selection of articles

We searched PubMed to identify reports of RCTs that had tested interventions for COVID-19 and were published up to 31 October 2020, using the following search strategy: (“COVID 19” [MeSH Terms] OR “2019 novel coronavirus” [All Fields] OR “2019 nCoV” [All Fields] OR “SARS CoV 2” [All Fields] OR “severe acute respiratory syndrome coronavirus 2” [All Fields] OR “coronavirus” [All Fields]) AND (“RCT” [All Fields] OR “randomized

controlled trial" [All Fields] OR "prospective cohort study" [All Fields] OR "longitudinal study" [All Fields] OR "cohort study" [All Fields]) AND ("2019/11/1" [Date - Publication]:"2020/10/31" [Date - Publication]). We included only reports published in English that presented the results of a RCT (defined as a prospective comparative study in which participants are allocated at random to one of the intervention arms). We excluded brief commentaries, research letters, observational studies (eg, cohort, case-control and cross-sectional studies), protocols, meta-analyses, and systematic reviews. We also excluded any reports that exclusively focused on cost-effective evaluations [13] or diagnostic test accuracy. Two reviewers (D.G.W. and Y.G.Z.) independently determined the eligibility of each abstract and a third reviewer (L.W.) was consulted in the event of any disagreements. The full text of the article linked to the abstract was retrieved when necessary, to determine eligibility or obtain additional information on the RCT (eg, source of funding).

## 2.2. Data extraction and reporting quality assessment of selected articles

For each selected RCT report, we extracted key information (journal type, average journal impact factor in recent 5 years, geographical location of first author, number of authors, type of institution, number of study centers, word count in the abstract, objective, structured format in the abstract, sample size, experimental interventions, number of intervention arms, number of primary outcomes, exact reporting of  $P$  values, and funding source) into an Excel document for further analyses.

Using the CONSORT explanation and elaboration statement for abstracts of parallel group randomized trials by Moher et al. [4] and previous reports [5,14], two reviewers worked independently (D.G.W. and Y.G.Z.) to make a judgment on whether an item in the RCT abstract was adequately reported. These two reviewers were calibrated through rounds of 10 randomly selected abstracts until strong agreement (Cohen's  $\kappa$  statistic  $\geq 0.90$ ) was reached. The reporting quality of all abstracts was then evaluated by these two assessors independently, with the overall reporting score being calculated for each abstract by giving a score of 1 for each of the 16 CONSORT essential items that were adequately described in the abstract and a score of 0 if the explanation was inadequate or unclear. These scores were summed to obtain a comprehensive reporting score for each selected abstract, which could range from 0 to 16. Any discrepancies were resolved in discussion with a third reviewer (L.W.).

## 2.3. Definition of primary outcomes of the RCTs

For this study, we defined an RCT's primary outcomes using the method outlined by Boutron et al. [6]:

- Explicit reporting of the primary outcomes in the original studies or clinical trial registrations; or
- Outcomes reported in the calculation of the required sample size for the RCT if no primary outcomes were explicitly reported; or
- Outcomes consistent with the primary objectives for the RCT if no primary outcomes were explicitly reported or mentioned in the sample size calculation.

RCTs with statistically nonsignificant primary outcomes ( $P \geq 0.05$  or a confidence interval for the effect estimate that included no difference) were included in the spin assessment.

## 2.4. Definition of spin and spin strategies

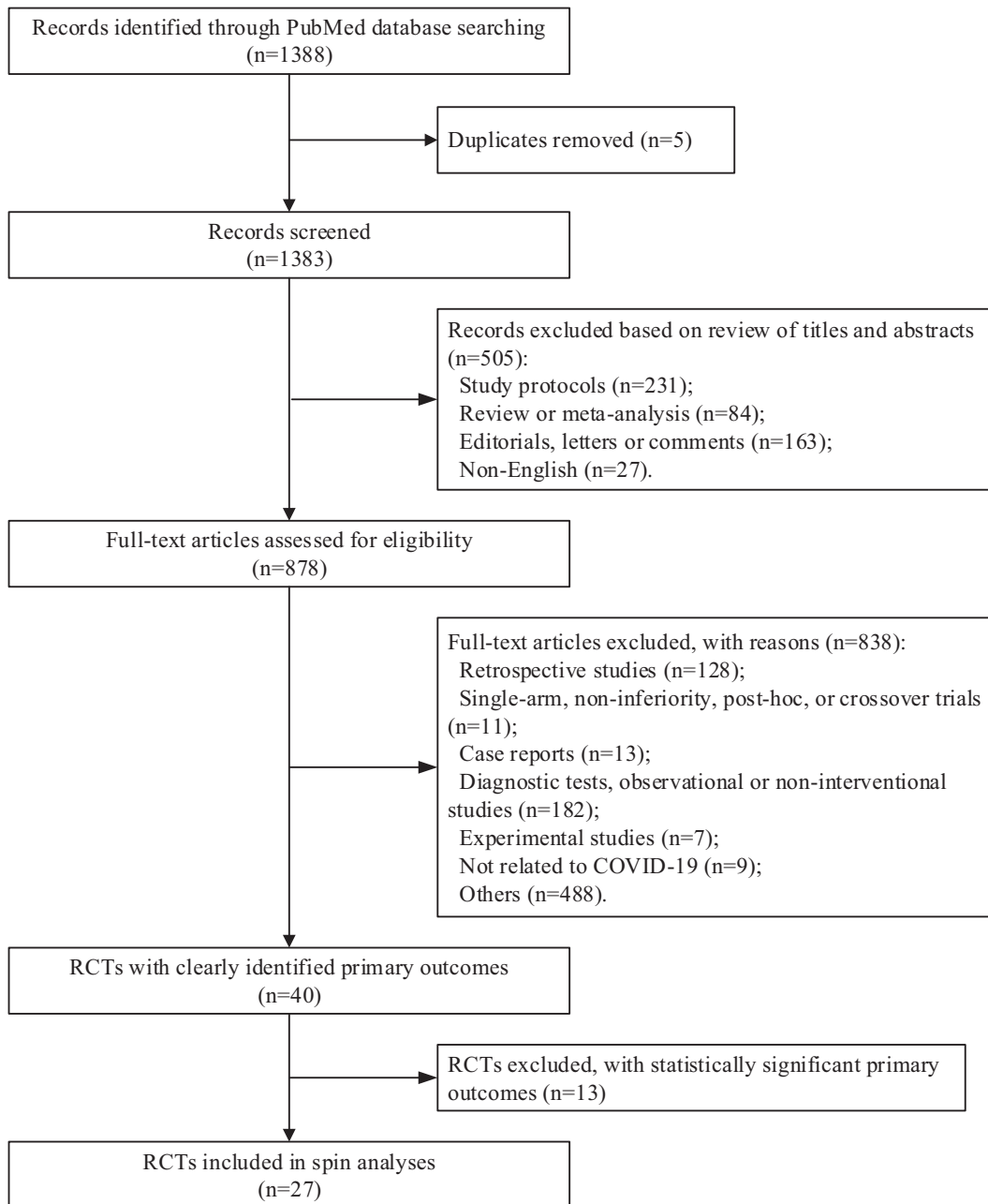
We used the Boutron et al. definition of spin: "specific reporting strategies from whatever motive, highlighting the interventions are beneficial despite the statistically nonsignificant differences for the primary outcomes, distorting the interpretation of study results and misleading readers" [6].

We classified spin strategies as follows, based on previous research [6,12]:

- Focusing on statistically significant results (eg, secondary outcomes, within-group comparisons and subgroup analyses).
- Claiming equivalent/noninferior/comparable/similar effects for statistically nonsignificant primary outcomes.
- Focusing only on primary outcomes or time-points with a statistically significant difference when there are several primary outcomes or multiple time-points for the primary outcomes.
- Claiming benefit of interventions with no consideration of the statistically nonsignificant results for primary outcomes or making recommendation for use of experimental interventions.

## 2.5. Spin assessment

Two reviewers (D.G.W. and Y.G.Z.) independently determined the presence of spin and spin strategies in the results and conclusion section of each abstract. As for the assessment of quality, an internal pilot study was conducted to calibrate these two assessors using rounds of 10 randomly selected abstracts until the agreement between them became strong (Cohen's  $\kappa$  statistic  $\geq 0.90$ ). A third reviewer (L.W.) was consulted to resolve any disagreements. We recorded the number of spin strategies in the results and conclusion section of each abstract, and assessed the level of spin in the conclusion section of the abstracts with a method from other research [6,12]: "high" was defined as no acknowledgment of statistically nonsignificant results for the primary outcome, no uncertainty and no recommendation for further trials; "moderate" was defined as no acknowledgment of statistically nonsignificant results for the primary outcome but a mention of uncertainty or



**Fig. 1.** Flow diagram for trial selections.

recommendations for further trials; “low” was defined as acknowledgment of statistically nonsignificant results for the primary outcome, or no acknowledgment of statistically nonsignificant results for the primary outcomes but a mention of uncertainty and recommendations for further trials, and “none” acted as a default category.

## 2.6. Statistical analyses

We calculated medians and interquartile ranges (IQR) for continuous variables and the number and proportion (%) of articles for categorical variables. We used linear

regression analyses to identify factors correlated with reporting quality, and the severity of spin. For factors associated with reporting quality, we used the overall reporting score as a dependent variable in regression analyses, and the univariate regression model was used first in the exploration of reporting quality predictors. Significant predictors were then tested by multivariate analyses. We used a similar process to explore factors relevant to the severity of spin (defined as the number of spin strategies in abstracts). All statistical analyses were performed with SPSS version 22.0 (IBM Corp, Armonk, NY, USA), and two-tailed *P* values less than 0.05 were considered statistically significant.

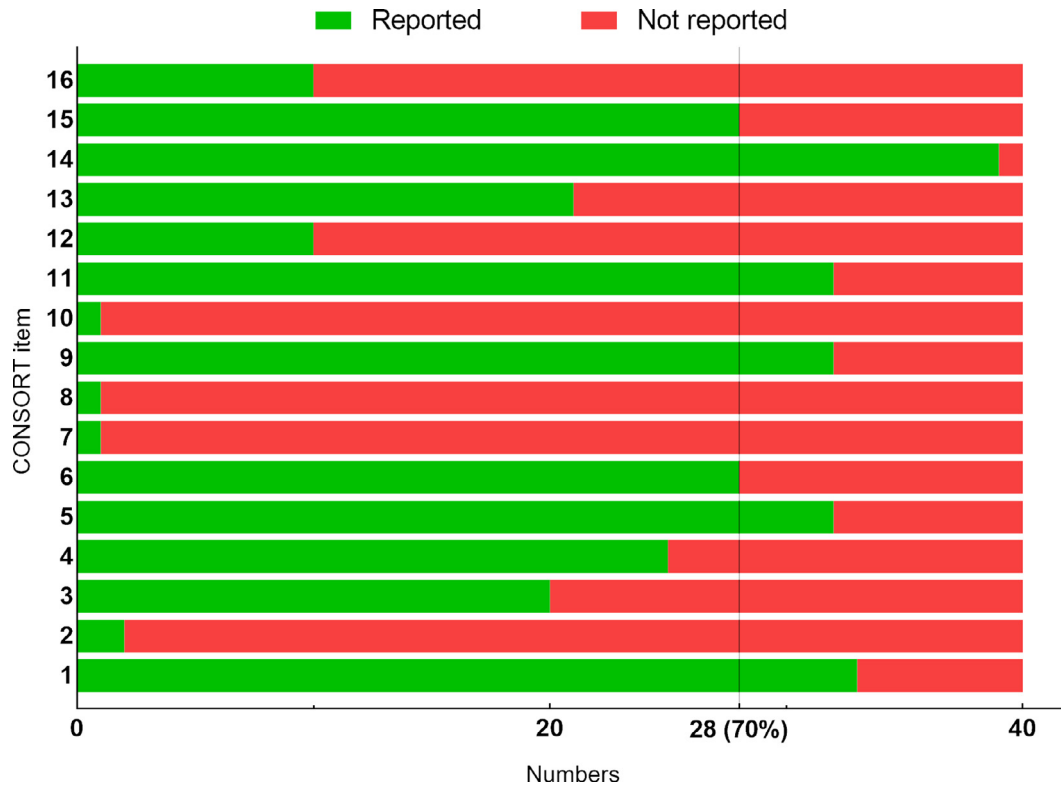


Fig. 2. Bar graph of the reporting quality of forty RCT abstracts based on the sixteen-item CONSORT statement for abstracts.

### 3. Results

#### 3.1. General characteristics of included studies

We retrieved a total of 1388 records from PubMed and identified 40 parallel-group RCTs among these, with 27 (67.5%) having a clearly identified, statistically nonsignificant primary outcome (Fig. 1).

Of the 40 eligible RCTs [15–54], 18 (45%) were single-center trials and most were conducted in the Asian region (26/40, 65%) and organized by universities (31/40, 78%). Thirty-three (83%) used a structured format for their abstract and same number of studies evaluated pharmacological interventions for the prevention or treatment of COVID-19. Ten (25%) abstracts were identified with more than 400 words and 17 (43%) studies reported a sample size more than 100 participants. The source of funding was not declared in the article for 6 (15%) RCTs. Detailed characteristics of the included studies are available in Table 1.

#### 3.2. Reporting quality assessment of the abstracts

A strong agreement was reached after two rounds of the pilot study for reporting quality evaluations ( $\kappa = 0.932$ ) before we assessed the reporting quality of all 40 abstracts (Fig. 2). The details for each item and sub-item are shown in Table 2 and Appendix 1 (Table 2a). Thirty-three (83%) studies could be identified as RCTs by the title of the ar-

ticle, 32 (80%) reported specific objectives or hypotheses, and 28 (70%) presented clearly defined primary outcomes in their abstract. Almost all abstracts mentioned random assignment but additional detail on sequence generation and allocation concealment was rare (37/40, 93%, 2/40, 5% and 1/40, 3%, respectively). The number of participants randomized to each group was found in 32 (80%) abstracts, but only one (3%) abstract reported the status of the trial (eg, closed or ongoing) in the results section, and only a quarter (10/40, 25%) provided full results for each intervention group (including primary outcome results, estimated effect size and its precision). Aside from their primary or secondary outcomes, 21 (53%) abstracts reported important adverse events or side effects in abstracts. Finally, 28 (70%) abstracts contained trial register information and 10 (25%) identified the source of funding.

#### 3.3. Predictors correlated to the reporting quality of abstracts in RCTs

The median overall reporting score for the 40 RCT abstracts was 8 (IQR: 6, 10). As shown in Figure 3, eight predictors were significantly correlated to a higher score in our univariate linear regression analyses: general journals ( $P = 0.009$ ), higher average IF ( $P < 0.001$ ), non-Asian locations ( $P = 0.015$ ), larger number of authors ( $P < 0.001$ ), multicenter studies ( $P = 0.008$ ), longer word count ( $P < 0.001$ ), structured format of abstracts ( $P = 0.006$ ), and

**Table 1.** General characteristics of the 40 RCTs related to COVID-19.

Characteristic	Category	N (%)
Journal type	Specialized medicine	23 (57.5)
	General medicine	17 (42.5)
Average journal impact factor in recent five years, median (IQR)	4.858 (2.735, 40.063)	
Geographical location	Asia	26 (65.0)
	Europe	7 (17.5)
	America	6 (15.0)
	Africa	1 (2.5)
No. of authors	≤7	6 (15.0)
	>7	34 (85.0)
Type of institution	University	31 (77.5)
	Others	9 (22.5)
Centers	Single center	18 (45.0)
	Multicenter	22 (55.0)
Word count in the abstract	<200	2 (5.0)
	200–400	28 (70.0)
	>400	10 (25.0)
Objective	Efficacy	20 (50.0)
	Safety	0 (0)
	Efficacy and safety	20 (50.0)
Structured format in the abstract	Yes	33 (82.5)
	No	7 (17.5)
Sample size	<50	10 (25.0)
	50–100	13 (32.5)
	>100	17 (42.5)
Intervention	Drugs	33 (82.5)
	Nonpharmaceutical interventions	7 (17.5)
Treatment arms	2	33 (82.5)
	>2	7 (17.5)
Primary outcomes	1	26 (65.0)
	>1	14 (35.0)
Exact P value	Yes	28 (70.0)
	No	12 (30.0)
Funding source	None	2 (5.0)
	Industry	5 (12.5)
	Nonindustry	21 (52.5)
	Industry and nonindustry	6 (15.0)
	Not reported	6 (15.0)
Total	40(100.0)	

IQR, interquartile range.

larger sample size ( $P = 0.002$ ). In multivariate models, only the predictor of longer word count ( $\beta = 0.351$ , 95% CI 0.044–0.658,  $P = 0.026$ ) remained as a statistically significant predictor (adjusted  $R^2 = 0.467$ ,  $P < 0.001$ ).

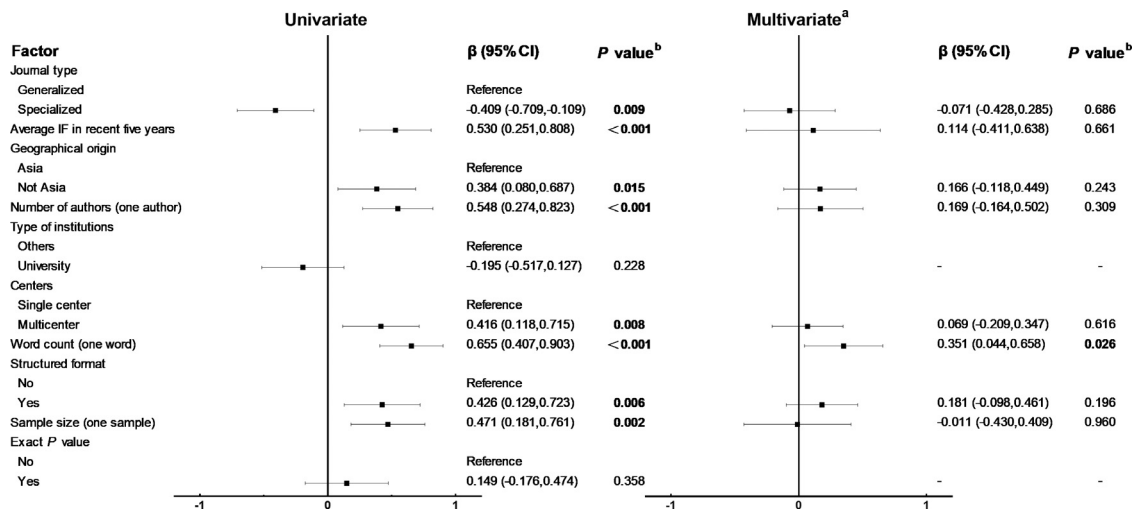
### 3.4. Spin strategies in abstracts

From the 40 RCT abstracts, the 27 (68%) with statistically nonsignificant primary outcomes were included in our

assessment of spin and 15 (56%) of these contained spin ( $\kappa = 0.977$ ), with multiple spin strategies present (Table 3 and Appendix 2 [Table 3a]). Fourteen (52%) abstracts had spin in the results section, with a focus on statistically significant secondary outcomes being the most frequent spin strategy (10/15, 67%). Other spin strategies found in the results section were focusing on statistically significant within-group comparison (2/15, 13%), focusing on statistically significant subgroup analyses (4/15, 27%), focusing

**Table 2.** Reporting of each item and subitem in forty RCT abstracts based on the CONSORT 2010 explanation and elaboration for abstracts.

Item	Criteria and subitems	N (%)
1. Title	Identification of the study as randomized in the title.	33 (82.5)
2. Trial design	Structured summary of the trial design (eg, parallel, cluster, crossover).	2 (5.0)
3. Participants	Eligibility criteria for participants and settings or locations where the data were collected.	20 (50.0)
	3a. Eligibility criteria for participants.	29 (72.5)
	3b. Settings or locations for data collections.	22 (55.0)
4. Interventions	Sufficient details of interventions intended for each group (eg, when, how).	25 (62.5)
5. Objectives	Specific objectives or hypotheses.	32 (80.0)
6. Primary outcomes	Clearly defined primary outcomes for this trial in methods.	28 (70.0)
7. Randomization	Scientific descriptions of how participants were allocated to interventions.	1 (2.5)
	7a. Random assignment (eg, random, randomized, randomization, random allocation).	37 (92.5)
	7b. Sequence generation (eg, random-number tables).	2(5.0)
	7c. Referring to allocation concealment.	1 (2.5)
8. Blinding(masking)	Whether or not participants, trial providers, and data collectors were blinded.	1 (2.5)
	8a. Brief descriptions only (eg, single-blind, double-blind, triple-blind).	9 (22.5)
9. Numbers randomized	Numbers of participants randomized to each group.	32 (80.0)
10. Recruitment and follow-up	Trial status (eg, on-going, closed to recruitment, closed to follow-up).	1 (2.5)
11. Numbers analyzed	Numbers of participants analyzed in each group.	32 (80.0)
	11a. Whether or not analyzed in accordance with the original grouping (eg, intention-to-treat analysis or pre-specified analysis).	1 (2.5)
12. Reports of primary outcomes	For the primary outcomes, a summary report of results for each group and the estimated effect size and its precision.	10 (25.0)
	12a. Primary outcome results for each group.	20 (50.0)
	12b. Estimated effect size.	17 (42.5)
	12c. Precision of the estimate (eg, 95%CI).	16 (40.0)
13. Harms	Important adverse events or side effects (seeing CONSORT for harms for specific guidance).	21 (52.5)
14. Conclusions	General interpretations corresponding to the results.	39 (97.5)
	14a. Benefits and harms balanced.	20 (50.0)
15. Trial registration	Trial registration number and the name of trial register.	28 (70.0)
16. Funding and supports	Supports of funding and supports.	10 (25.0)



**Fig. 3.** Factors associated with overall reporting score in forty RCT abstracts.

<sup>a</sup>For the multivariate linear regression analyses, adjusted  $R^2 = 0.467$ ,  $P < 0.001$ .

<sup>b</sup> $P < 0.05$  was considered statistically significant, shown in bold.

Abbreviations: IF, impact factor; CI, confidence interval.



**Table 3.** Spin strategies identified in 16 RCT abstracts with spin

	N (%)
Spin strategies in the result section	
Focusing on statistically significant within-group comparisons.	2 (12.5)
Focusing on statistically significant secondary outcomes.	12 (75.0)
Focusing on statistically significant subgroup analyses.	4 (25.0)
Focusing on statistically significant within- or between-group comparisons of secondary outcomes.	1 (6.3)
Focusing only on primary outcome of statistical significance when several primary outcomes exist.	3 (18.8)
Spin strategies in the conclusion section	
Claiming equivalent/noninferior/comparable/similar effects for statistically nonsignificant primary endpoints.	0 (0)
Focusing only on statistically significant results (eg, secondary outcomes, subgroup analyses, within-group analyses).	4 (25.0)
Claiming benefit with no consideration of the statistically nonsignificant primary outcomes.	12 (75.0)
Recommendation to use the experimental treatment.	5 (31.3)
Focusing only on outcomes with statistical significance when several primary outcomes exist.	3 (18.8)
Focusing only on time-points with statistical significance when multiple time-points for primary outcomes exist.	0 (0)

on statistically significant within- or between-group comparison of secondary outcomes (1/15, 7%), and focusing only on primary outcomes of statistical significance when there are several primary outcomes (3/15, 20%).

Spin was also identified in the conclusion section of 12 (44%) of 27 abstracts with statistically nonsignificant primary outcomes. Claiming benefit with no consideration of the statistically nonsignificant nature of the result for the primary outcomes was the most frequent spin strategy in the abstract conclusions (12/15, 80%). Five (33%) abstracts recommended use of experimental treatment, and 4 (27%) focused only on statistically significant results. Three (20%) abstracts focused only on a primary outcome of statistical significance in their conclusion section despite there being several primary outcomes in the RCT. None claimed equivalent/noninferior/comparable/similar effects for statistically nonsignificant results or focused only on a timepoint of statistical significance when there were multiple timepoints for the primary outcomes.

### 3.5. Level of spin evaluation in selected abstracts

Among the 15 abstracts with spin, 9 (60%) contained more than two spin strategies, with one (11%) having seven spin strategies. There were 1 or 2 spin strategies in the remaining 6 (40%) abstracts. Furthermore, among the 12 abstracts with spin in the conclusion section, we found a high level of spin in 6 (50%) abstracts. The level of spin was moderate in 3 (25%) and low in 3 (25%) conclusion sections (Table 4).

### 3.6. Potential predictors associated with the severity of spin

We explored whether the potential characteristic predictors were correlated with spin (Fig. 4). In summary, three factors were significantly associated with spin severity in our univariate analyses: less spin occurred in general and

**Table 4.** Assessment of level of spin in the conclusion section of RCT abstracts

Level of spin in the conclusion section	N (%)
Low <sup>a</sup>	3 (25.0)
Moderate <sup>b</sup>	3 (25.0)
High <sup>c</sup>	6 (50.0)

<sup>a</sup> Acknowledge statistically nonsignificant results for the primary outcome, or no acknowledgment of statistically nonsignificant results for the primary outcome but reported with uncertainty and recommendations for further trials.

<sup>b</sup> No acknowledgment of statistically nonsignificant results for the primary outcome but reported with uncertainty or recommendations for further trials.

<sup>c</sup> No acknowledgment of statistically nonsignificant results for the primary outcome, no uncertainty and no recommendations for further trials.

higher average IF journals and studies from non-Asian regions ( $P = 0.005$ , 0.023 and 0.004, respectively). Only the predictor of studies from non-Asian regions ( $\beta = -0.426$ , 95% CI -0.756 to -0.096,  $P = 0.014$ ) remained statistically significant in our multivariate analyses (adjusted  $R^2 = 0.388$ ,  $P = 0.002$ ).

## 4. Discussion

To the best of our knowledge, this is the first study to systematically assess the reporting quality and presence of spin in the abstracts of reports of RCTs testing interventions for COVID-19. The CONSORT statement for abstracts provides authors with guidance on the necessary details and clarity required for good reporting. It is intended to improve the reporting quality of abstracts, and recently, has been used as a tool to evaluate the reporting quality of abstracts [55–59]. Our results show that the overall reporting quality of the abstracts of these COVID-19 RCTs is far from satisfactory, with adherence of reports to the CONSORT items ranging from 2.5% to 97.5%.

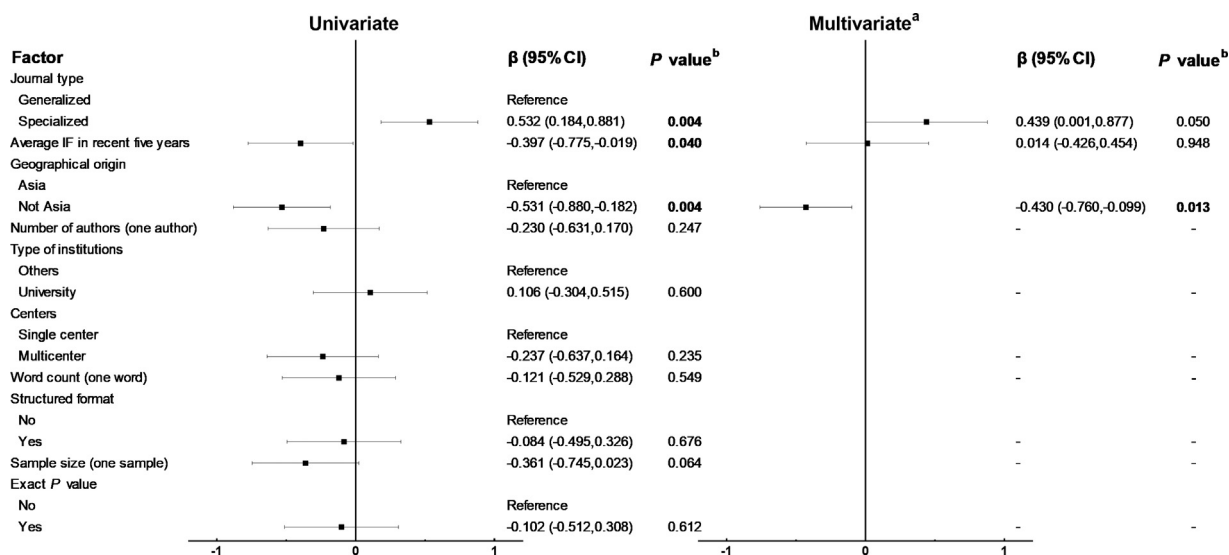


Fig. 4. Factors related to spin severity in 27 RCT abstracts with statistically nonsignificant primary outcomes.

<sup>a</sup>For the multivariate linear regression analyses, adjusted  $R^2=0.388$ ,  $P = 0.002$ .

<sup>b</sup> $P < 0.05$  was considered statistically significant, shown in bold.

Abbreviations: IF, impact factor; CI, confidence interval.

Notably, the CONSORT items most inadequately reported were those related to trial design, randomization, blinding, and trial status. Other studies in the medical literature have also shown a high prevalence of nonadherence to the CONSORT guidelines in RCT abstracts [60–65]. For example, Mozetic et al. found that the most underreported items were related to the methods items of CONSORT, such as trial design, allocation concealment, implementation of randomization sequence, and blinding [63]. Gallo et al. also found this, with limited adherence to the CONSORT for abstracts checklists among RCT abstracts published in the top plastic surgery journals [64]. They concluded that the most poorly reported items were trial registration (4%), method of randomization (2.4%), and source of trial funding (0%). Furthermore, Janackovic and Puljak evaluated 622 RCT abstracts in anesthesiology and observed the lowest consistency in trial design (18%), recruitment status (9%), number of participants analyzed (8%), randomization (3%), and funding (0.2%) [65]. Even in the top five highest-impact general medical journals, there was lack of adherence to the CONSORT statement for abstracts [66]. Overall, RCT abstracts in medical journals are poorly reported, providing readers with insufficient information, which means that readers might not get useful information in a short time and might even be misled by the incomplete results. One possible explanation for this failure to follow the guidelines is that some researchers are not familiar with the CONSORT statement, and that they may repeat the structure and content they have seen in other abstracts when they draft their report. In addition, the editorial office has certain responsibility, because abstracts are often structured in accordance with the au-

thor guidelines of academic journals. To remedy this, we recommend that authors, peer reviewers and editors make better use of reporting guidelines such as CONSORT and its extensions when preparing, appraising and editing research articles.

It should be noted that poor reporting quality of abstracts cannot be misinterpreted as poor study design. Limited length of the abstract means that it is often not possible to show all details of the research. Therefore, consistent with our results, items on RCT methods tend to have lower reporting rates in abstracts while, at least in journal papers with high impact, methodological details are well formulated in the full text.

The concept of spin, applying to clinical research, means selectively reporting significant findings while neglecting nonstatistically significant results. The role of spin is to make imperfect research results more meaningful, and thus stand out from similar studies. Up to now, there is no completely objective evaluation method for spin. In addition to the spin strategies mentioned by Boutron [6], more unrecognized strategies exist. We assessed the most common and widely used eleven spin strategies, and nearly 60% of the RCT abstracts with statistically nonsignificant primary outcomes in our study contained spin. High prevalence of spin has also been reported by other studies, ranging widely from 17% to 86% [5,8–12,67–70]. We speculate that high spin prevalence in abstracts of COVID-19 RCTs is due to the specific background at that time, that is, on the one hand, lower standards in medical journals and rapid peer review might lead to lax assessment of manuscripts, while on the other hand, high mortality and morbidity of COVID-19 needed prompt evidence on therapies, result-

ing in the emergence of large numbers of RCTs of interventions which were subsequently found to be ineffective. Consequently, the existence of spin in these early reports seems not surprising.

Among various forms of spin strategy, the most common one in the results section of the abstract was focusing on statistically significant secondary outcomes to claim benefit, while no consideration of the statistically nonsignificant primary outcomes was the most common strategy found in the conclusion sections. Similar findings were reported by Jellison et al. [68]. In another study, however, focusing on statistically significant within-group analysis was the most common spin strategy used in the result section of abstracts, and claiming equivalence or non-inferiority of results with nonsignificance was the most common in the conclusion section [5]. Moreover, Turrentine analyzed 83 scientific publications with spin in abstracts and concluded that the more common types of spin strategies in general obstetrics and gynecology were: emphasizing statistically significant secondary results (40%), interpreting nonstatistically significant primary outcomes as equivalent or similar effectiveness (37%) and claiming beneficial effects of treatment despite the statistical nonsignificance (15%) [70]. Lockyer et al. indicated that there was potential for spin in wound care trials emphasizing study results of significance rather than the importance of outcomes [67]. This is concerning because clinicians are prone to misunderstand the outcomes of a trial when spin is present and make inappropriate clinical decisions [6]. This makes it especially important to find ways to identify and mitigate spin [10]. Readers should keep the concept of spin in mind when reading abstracts and be aware of the diversity and heterogeneity of spin strategies, and researchers should properly report their results and conclusions within the limited word count for the abstract, instead of giving space to only those results that they wish to highlight to show the importance of their research. Journal reviewers and editors also need to be rigorous in their assessment of manuscripts. Our analyses of the predictors potentially associated with reporting quality show that a larger word count was significantly associated with better quality reporting in RCT abstracts. For this reason, allowing authors more words in their abstracts might be a simple method to improve the reporting quality of abstracts. In addition to larger word counts, reporting of trial registration and funding were also positively correlated with high-quality reporting [71]. In the multivariate analysis of relevant factors to spin severity, we found that research from non-Asian regions might be relevant to fewer spin strategies. Both Cooper and Reynolds-Vaughn demonstrated that a majority of abstracts with spin were funded by industry [9,72], while Jellison et al. found no relationship between industry funding and spin in abstracts [68], and use of statistician and article section were further confirmed to be unrelated to the presence of spin in another study [11]. Furthermore,

numbers of research centers were reported associated with presence of spin by Wu et al. [8], and Checketts et al. found that word count limit promoted the prevalence of spin [10]. Whatever, these findings raise concerns about the reporting specification of abstracts, and it is of vital importance for researchers to objectively and accurately report their findings.

There are some limitations to our study. Although we included all abstracts for RCTs of interventions for COVID-19 that we identified in PubMed up to the end of October 2020, the number of included RCTs is still relatively small, which means that our estimates might change if the study was expanded to include more abstracts. Second, although we evaluated spin strategies with a predesigned 11-item form used in other studies, some other potential spin strategies might have been omitted, leading to an underestimate of the presence of spin. Third, only RCTs testing interventions for COVID-19 were included when evaluating the reporting quality and spin of abstracts, that is to say, our analyses was focused on abstract sections, and our conclusions of poor reporting quality and high spin should not be extended to the full text. Fourth, evaluations of reporting quality and spin are subjective, and although we adopted an approach of double, independent and calibrated assessment to control the magnitude of subjectivity, if others repeated our assessments they might obtain different results. Despite these limitations, our study provides important new insights for the reporting quality of RCT abstracts which may have implications to research reporting more generally, as well as its specific relevance to the reporting of RCTs of interventions for COVID-19.

## 5. Conclusions

As of the end of October 2020, the reporting quality of the abstracts for reports of RCTs of interventions for COVID-19 is far from satisfactory. The frequency, extent and level of spin are relatively high in these abstracts, highlighting a need for the results and conclusion sections of such abstracts to be interpreted with caution. We hope that this assessment will raise readers' awareness of the need to carefully appraise abstracts, to be aware of the concept of spin, and to be especially cautious if the pressures of the COVID-19 pandemic increase their reliance on the results and conclusions reported in abstracts.

## Author contributions

Concept and design: YGZ, HF and WML.

Data extraction, statistical analysis and interpretation: DGW, LMC, YGZ, LW, FH, JL, YXL and MC.

Drafting the manuscript: DGW, LMC and YGZ.

Revising the manuscript: FH and MC.

Supervision: HF and WML.

## Funding

This work was supported by National Key R&D Program of China (2017YFC1309703) and 1.3.5 project for disciplines of excellence-Clinical Research Incubation Project, West China Hospital, Sichuan University (2019HXFH008).

## Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.jclinepi.2021.06.027.

## References

- [1] Smith R. What clinical information do doctors need? *BMJ* 1996;313:1062–8 PubMed PMID: 8898602; PubMed Central PMCID: PMCPMC2352351. doi:10.1136/bmj.313.7064.1062.
- [2] Johnson HL, Fontelo P, Olsen CH, Jones KD 2nd, Gimbel RW. Family nurse practitioner student perception of journal abstract usefulness in clinical decision making: a randomized controlled trial. *J Am Assoc Nurse Pract* 2013;25:597–603 PubMed PMID: 24170534. doi:10.1111/1745-7599.12013.
- [3] Walters C, Torgerson T, Fladie I, Clifton A, Meyer C, Vassar M. Are randomized controlled trials being conducted with the right justification? *J Evid Based Med* 2020;13:181–2 PubMed PMID: 32615030. doi:10.1111/jebm.12405.
- [4] Moher D, Hopewell S, Schulz KF, Montori V, Gotzsche PC, Devereaux PJ, et al. CONSORT 2010 explanation and elaboration: updated guidelines for reporting parallel group randomised trials. *J Clin Epidemiol* 2010;63:e1–37 PubMed PMID: 20346624. doi:10.1016/j.jclinepi.2010.03.004.
- [5] Fang X, Hua F, Riley P, Chen F, Zhang L, Walsh T, et al. Abstracts of published randomised controlled trials in endodontics: reporting quality and spin. *Int Endod J* 2020;1050–61 PubMed PMID: 32333794. doi:10.1111/iej.13310.
- [6] Boutron I, Dutton S, Ravaud P, Altman DG. Reporting and interpretation of randomized controlled trials with statistically nonsignificant results for primary outcomes. *JAMA* 2010;303:2058–64 PubMed PMID: 20501928. doi:10.1001/jama.2010.651.
- [7] Bero L, Chiu K, Grundy Q. The SSSPIN study-spin in studies of spin: meta-research analysis. *BMJ* 2019;367:l6202. PubMed PMID: 31852680; PubMed Central PMCID: PMCPMC7191944 at [www.icmje.org/coi\\_disclosure.pdf](http://www.icmje.org/coi_disclosure.pdf) and declare: no support from any organisation for the submitted work; no financial relationships with any organisations that might have an interest in the submitted work in the previous three years; no other relationships or activities that could appear to have influenced the submitted work. doi:10.1136/bmj.l6202.
- [8] Wu X, Yan Q, Fang X, Hua F, Shi B, Tu YK. Spin in the abstracts of randomized controlled trials in periodontology and oral implantology: a cross-sectional analysis. *J Clin Periodontol* 2020 PubMed PMID: 32618017. doi:10.1111/jcpe.13340.
- [9] Cooper CM, Gray HM, Ross AE, Hamilton TA, Bea Downs J, Wayant C. Evaluation of spin in the abstracts of otolaryngology randomized controlled trials. *Laryngoscope* 2018;2036–40 PubMed PMID: 30578543. doi:10.1002/lary.27750.
- [10] Checketts JX, Riddle J, Zaaza Z, Boose MA, Whitener JH, Vassar MB. An evaluation of spin in lower extremity joint trials. *J Arthroplasty* 2019;34:1008–12 PubMed PMID: 30733070. doi:10.1016/j.arth.2019.01.016.
- [11] Arunachalam L, Hunter IA, Killeen S. Reporting of randomized controlled trials with statistically nonsignificant primary outcomes published in high-impact surgical journals. *Ann Surg* 2017;265:1141–5 PubMed PMID: 27257737. doi:10.1097/SLA.0000000000001795.
- [12] Khan MS, Lateef N, Siddiqi TJ, Rehman KA, Alnaimat S, Khan SU, et al. Level and prevalence of spin in published cardiovascular randomized clinical trial reports with statistically nonsignificant primary outcomes: a systematic review. *JAMA Netw Open* 2019;2:e192622 PubMed PMID: 31050775; PubMed Central PMCID: PMCPMC6503494. doi:10.1001/jamanetworkopen.2019.2622.
- [13] Clarke L. An introduction to economic studies, health emergencies, and COVID-19. *J Evid Based Med* 2020;13:161–7 PubMed PMID: 32470229; PubMed Central PMCID: PMCPMC7283784. doi:10.1111/jebm.12395.
- [14] Liu K, Zeng J, Pei W, Chen S, Luo Z, Lu L. Assessing the reporting quality in randomized controlled trials of acupuncture for postherpetic neuralgia using the CONSORT statement and STRICTA guidelines. *J Pain Res* 2019;12:2359–70 PubMed PMID: 31534360; PubMed Central PMCID: PMCPMC6681161. doi:10.2147/JPR.S210471.
- [15] Zhu FC, Guan XH, Li YH, Huang JY, Jiang T, Hou LH, et al. Immunogenicity and safety of a recombinant adenovirus type-5-vectored COVID-19 vaccine in healthy adults aged 18 years or older: a randomised, double-blind, placebo-controlled, phase 2 trial. *Lancet* 2020;396:479–88 PubMed PMID: 32702299; PubMed Central PMCID: PMCPMC7836858. doi:10.1016/S0140-6736(20)31605-6.
- [16] Zheng F, Zhou Y, Zhou Z, Ye F, Huang B, Huang Y, et al. SARS-CoV-2 clearance in COVID-19 patients with Nofaferon treatment: a randomized, open-label, parallel-group trial. *Int J Infect Dis* 2020;99:84–91 PubMed PMID: 32758689; PubMed Central PMCID: PMCPMC7397938. doi:10.1016/j.ijid.2020.07.053.
- [17] Zhao J, Yang X, Wang C, Song S, Cao K, Wei T, et al. Yidu-toxicity blocking lung decoction ameliorates inflammation in severe pneumonia of SARS-COV-2 patients with Yidu-toxicity blocking lung syndrome by eliminating IL-6 and TNF- $\alpha$ . *Biomed Pharmacother* 2020;129:110436 PubMed PMID: 32768938; PubMed Central PMCID: PMCPMC7303599. doi:10.1016/j.biopha.2020.110436.
- [18] Ye YA, Group GCC. Guideline-based chinese herbal medicine treatment plus standard care for severe coronavirus disease 2019 (G-CHAMPS): evidence from China. *Front Med (Lausanne)* 2020;7:256 PubMed PMID: 32574340; PubMed Central PMCID: PMCPMC7267028. doi:10.3389/fmed.2020.00256.
- [19] Xiong WZ, Wang G, Du J, Ai W. Efficacy of herbal medicine (Xuanfei Baidu decoction) combined with conventional drug in treating COVID-19: a pilot randomized clinical trial. *Integr Med Res* 2020;9:100489 PubMed PMID: 32874913; PubMed Central PMCID: PMCPMC7452296. doi:10.1016/j.imr.2020.100489.
- [20] Xiao M, Tian J, Zhou Y, Xu X, Min X, Lv Y, et al. Efficacy of Huoxiang Zhengqi dropping pills and Lianhua Qingwen granules in treatment of COVID-19: a randomized controlled trial. *Pharmacol Res* 2020;161:105126 PubMed PMID: 32781283; PubMed Central PMCID: PMCPMC7414728. doi:10.1016/j.phrs.2020.105126.
- [21] Wu X, Yu K, Wang Y, Xu W, Ma H, Hou Y, et al. Efficacy and safety of triazavirin therapy for coronavirus disease 2019: a pilot randomized controlled trial. *Engineering (Beijing)* 2020;6:1185–91 PubMed PMID: 32923016; PubMed Central PMCID: PMCPMC7476906. doi:10.1016/j.eng.2020.08.011.
- [22] Wei N, Huang BC, Lu SJ, Hu JB, Zhou XY, Hu CC, et al. Efficacy of internet-based integrated intervention on depression and anxiety symptoms in patients with COVID-19. *J Zhejiang Univ Sci B* 2020;21:400–4 PubMed PMID: 32425006; PubMed Central PMCID: PMCPMC7203540. doi:10.1631/jzus.B2010013.
- [23] Wang Y, Zhang D, Du G, Du R, Zhao J, Jin Y, et al. Remdesivir in adults with severe COVID-19: a randomised, double-blind, placebo-controlled, multicentre trial. *Lancet* 2020;395:1569–78 PubMed PMID: 32423584; PubMed Central PMCID: PMCPMC7190303. doi:10.1016/S0140-6736(20)31022-9.
- [24] Wang JB, Wang ZX, Jing J, Zhao P, Dong JH, Zhou YF, et al. Exploring an integrative therapy for treating COVID-19: a randomized controlled trial. *Chin J Integr Med* 2020;26:648–55 PubMed PMID: 32676976; PubMed Central PMCID: PMCPMC7364292. doi:10.1007/s11655-020-3426-7.

- [25] Vlaar APJ, de Bruin S, Busch M, Timmermans S, van Zeggeren IE, Koning R, et al. Anti-C5a antibody IFX-1 (vilobelimab) treatment versus best supportive care for patients with severe COVID-19 (PANAMO): an exploratory, open-label, phase 2 randomised controlled trial. *Lancet Rheumatol* 2020;2:e764–ee73 PubMed PMID: 33015643; PubMed Central PMCID: PMC7521913. doi:10.1016/S2665-9913(20)30341-6.
- [26] Tang W, Cao Z, Han M, Wang Z, Chen J, Sun W, et al. Hydroxychloroquine in patients with mainly mild to moderate coronavirus disease 2019: open label, randomised controlled trial. *BMJ* 2020;369:m1849 PubMed PMID: 32409561; PubMed Central PMCID: PMC7221473 at [www.icmje.org/coi\\_disclosure.pdf](http://www.icmje.org/coi_disclosure.pdf) and declare: no support from any organisation for the submitted work other than those listed above; no financial relationships with any organisations that might have an interest in the submitted work in the previous three years; no other relationships or activities that could appear to have influenced the submitted work. doi:10.1136/bmj.m1849.
- [27] Shu L, Niu C, Li R, Huang T, Wang Y, Huang M, et al. Treatment of severe COVID-19 with human umbilical cord mesenchymal stem cells. *Stem Cell Res Ther* 2020;11:361 PubMed PMID: 32811531; PubMed Central PMCID: PMC7432540. doi:10.1186/s13287-020-01875-5.
- [28] Sekhavati E, Jafari F, SeyedAlinaghi S, Jamalimoghdamshahkhalil S, Sadr S, Tabarestani M, et al. Safety and effectiveness of azithromycin in patients with COVID-19: an open-label randomised trial. *Int J Antimicrob Agents* 2020;56:106143 PubMed PMID: 32853672; PubMed Central PMCID: PMC7445147. doi:10.1016/j.ijantimicag.2020.106143.
- [29] Sadeghi A, Ali Asgari A, Norouzi A, Kheiri Z, Anushirvani A, Montazeri M, Sofosbuvir and dactatasvir compared with standard of care in the treatment of patients admitted to hospital with moderate or severe coronavirus infection (COVID-19): a randomized controlled trial. *J Antimicrob Chemother* 2020;75:3379–85 PubMed PMID: 32812039; PubMed Central PMCID: PMC7454592. doi:10.1093/jac/dkaa334.
- [30] Beigel JH, Tomashek KM, Dodd LE, Mehta AK, Zingman BS, Kalil AC, et al. Remdesivir for the treatment of COVID-19—final report. *N Engl J Med* 2020;383:1813–26 PubMed PMID: 32445440; PubMed Central PMCID: PMC7262788. doi:10.1056/NEJMoa2007764.
- [31] Mitja O, Corbacho-Monne M, Ubals M, Tebe C, Penafiel J, Tobias A, et al. Hydroxychloroquine for early treatment of adults with mild COVID-19: a randomized-controlled trial. *Clin Infect Dis* 2020 PubMed PMID: 32674126; PubMed Central PMCID: PMC7454406. doi:10.1093/cid/ciaa1009.
- [32] Miller J, Bruen C, Schnaus M, Zhang J, Ali S, Lind A, et al. Auxora versus standard of care for the treatment of severe or critical COVID-19 pneumonia: results from a randomized controlled trial. *Crit Care* 2020;24:502 10.1186/s13054-020-03220-x. PubMed PMID: 32795330; PubMed Central PMCID: PMC7427272.
- [33] Lyngbakken MN, Berdal JE, Eskesen A, Kvale D, Olsen IC, Rueegg CS, et al. A pragmatic randomized controlled trial reports lack of efficacy of hydroxychloroquine on coronavirus disease 2019 viral kinetics. *Nat Commun* 2020;11:5284 PubMed PMID: 33082342; PubMed Central PMCID: PMC7576792. doi:10.1038/s41467-020-19056-6.
- [34] Lou Y, Liu L, Yao H, Hu X, Su J, Xu K, et al. Clinical outcomes and plasma concentrations of baloxavir marboxil and favipiravir in COVID-19 patients: an exploratory randomized, controlled trial. *Eur J Pharm Sci* 2021;157:105631 PubMed PMID: 33115675; PubMed Central PMCID: PMC7585719. doi:10.1016/j.ejps.2020.105631.
- [35] Liu K, Chen Y, Wu D, Lin R, Wang Z, Pan L. Effects of progressive muscle relaxation on anxiety and sleep quality in patients with COVID-19. *Complement Ther Clin Pract* 2020;39:101132 PubMed PMID: 32379667; PubMed Central PMCID: PMC7102525. doi:10.1016/j.ctcp.2020.101132.
- [36] Li Y, Xie Z, Lin W, Cai W, Wen C, Guan Y, et al. Efficacy and safety of lopinavir/ritonavir or arbidol in adult patients with mild/moderate COVID-19: an exploratory randomized controlled trial. *Med (N Y)* 2020;1 105-13 e4PubMed PMID: 32838353; PubMed Central PMCID: PMC7235585. doi:10.1016/j.medj.2020.04.001.
- [37] Li L, Zhang W, Hu Y, Tong X, Zheng S, Yang J, et al. Effect of convalescent plasma therapy on time to clinical improvement in patients with severe and life-threatening COVID-19: a randomized clinical trial. *JAMA* 2020;324:460–70 PubMed PMID: 32492084; PubMed Central PMCID: PMC7270883. doi:10.1001/jama.2020.10044.
- [38] Hung IF, Lung KC, Tso EY, Liu R, Chung TW, Chu MY, et al. Triple combination of interferon beta-1b, lopinavir-ritonavir, and ribavirin in the treatment of patients admitted to hospital with COVID-19: an open-label, randomised, phase 2 trial. *Lancet* 2020;395:1695–704 PubMed PMID: 32401715; PubMed Central PMCID: PMC7211500. doi:10.1016/S0140-6736(20)31042-4.
- [39] Hu K, Guan WJ, Bi Y, Zhang W, Li L, Zhang B, et al. Efficacy and safety of Lianhuaqingwen capsules, a repurposed Chinese herb, in patients with coronavirus disease 2019: A multicenter, prospective, randomized controlled trial. *Phytomedicine* 2020;153242 PubMed PMID: 32425361; PubMed Central PMCID: PMC7229744. doi:10.1016/j.phymed.2020.153242.
- [40] Furtado RHM, Berwanger O, Fonseca HA, Correa TD, Ferraz LR, Lapa MG, et al. Azithromycin in addition to standard of care versus standard of care alone in the treatment of patients admitted to the hospital with severe COVID-19 in Brazil (COALITION II): a randomised clinical trial. *Lancet* 2020;396 959-67PubMed PMID: 32896292; PubMed Central PMCID: PMC7836431. doi:10.1016/S0140-6736(20)31862-6.
- [41] Deftereos SG, Giannopoulos G, Vrachatis DA, Siasos GD, Giotaki SG, Gargalianos P, et al. Effect of colchicine vs standard care on cardiac and inflammatory biomarkers and clinical outcomes in patients hospitalized with coronavirus disease 2019: the GRECCO-19 randomized clinical trial. *JAMA Netw Open* 2020;3:e2013136 PubMed PMID: 32579195; PubMed Central PMCID: PMC7315286. doi:10.1001/jamanetworkopen.2020.13136.
- [42] Cao Y, Wei J, Zou L, Jiang T, Wang G, Chen L, et al. Ruxolitinib in treatment of severe coronavirus disease 2019 (COVID-19): a multicenter, single-blind, randomized controlled trial. *J Allergy Clin Immunol* 2020;146 137-46 e3PubMed PMID: 32470486; PubMed Central PMCID: PMC7250105. doi:10.1016/j.jaci.2020.05.019.
- [43] Cao B, Wang Y, Wen D, Liu W, Wang J, Fan G, et al. A trial of lopinavir-ritonavir in adults hospitalized with severe COVID-19. *N Engl J Med* 2020;382 1787-99PubMed PMID: 32187464; PubMed Central PMCID: PMC7121492. doi:10.1056/NEJMoa2001282.
- [44] Boulware DR, Pullen MF, Bangdiwala AS, Pastick KA, Lofgren SM, Okafor EC, et al. A randomized trial of hydroxychloroquine as postexposure prophylaxis for COVID-19. *N Engl J Med* 2020;383 517-25PubMed PMID: 32492293; PubMed Central PMCID: PMC7289276. doi:10.1056/NEJMoa2016638.
- [45] Borba MGS, Val FFA, Sampaio VS, Alexandre MAA, Melo GC, Brito M, et al. Effect of high vs low doses of chloroquine diphosphate as adjunctive therapy for patients hospitalized with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection: a randomized clinical trial. *JAMA Netw Open* 2020;3:e208857 PubMed PMID: 32330277. doi:10.1001/jamanetworkopen.2020.8857.
- [46] Abd-El salam S, Esmail ES, Khalaf M, Abdo EF, Medhat MA, Abd El Ghafar MS. Hydroxychloroquine in the treatment of COVID-19: a multicenter randomized controlled study. *Am J Trop Med Hyg* 2020;103 1635-9PubMed PMID: 32828135; PubMed Central PMCID: PMC7543820. doi:10.4269/ajtmh.20-0873.

- [47] Abbaspour Kasgari H, Moradi S, Shabani AM, Babamahmoodi F, Davoudi Badabi AR, Davoudi L. Evaluation of the efficacy of sofosbuvir plus daclatasvir in combination with ribavirin for hospitalized COVID-19 patients with moderate disease compared with standard care: a single-centre, randomized controlled trial. *J Antimicrob Chemother* 2020;75:3373–8. PubMed PMID: 32812025; PubMed Central PMCID: PMCPCMC7454669. doi:10.1093/jac/dkaa332.
- [48] Spinner CD, Gottlieb RL, Criner GJ, Arribas Lopez JR, Cattellan AM, Soriano Viladomiu A. Effect of remdesivir vs standard care on clinical status at 11 days in patients with moderate COVID-19: a randomized clinical trial. *JAMA* 2020;324:1048–57. PubMed PMID: 32821939; PubMed Central PMCID: PMCPCMC7442954. doi:10.1001/jama.2020.16349.
- [49] Skipper CP, Pastick KA, Engen NW, Bangdiwala AS, Abassi M, Lofgren SM, et al. Hydroxychloroquine in nonhospitalized adults with early COVID-19: a randomized trial. *Ann Intern Med* 2020;173:623–31. PubMed PMID: 32673060; PubMed Central PMCID: PMCPCMC7384270. doi:10.7326/M20-4207.
- [50] Folegatti PM, Ewer KJ, Aley PK, Angus B, Becker S, Belij-Rammerstorfer S, et al. Safety and immunogenicity of the ChAdOx1 nCoV-19 vaccine against SARS-CoV-2: a preliminary report of a phase 1/2, single-blind, randomised controlled trial. *Lancet* 2020;396:467–78. PubMed PMID: 32702298; PubMed Central PMCID: PMCPCMC7445431. doi:10.1016/S0140-6736(20)31604-4.
- [51] Entrenas Castillo M, Entrenas Costa LM, Vaquero Barrios JM, Alcalá Diaz JF, Lopez Miranda J, Bouillon R. Effect of calcifediol treatment and best available therapy versus best available therapy on intensive care unit admission and mortality among patients hospitalized for COVID-19: a pilot randomized clinical study. *J Steroid Biochem Mol Biol* 2020;203:105751. PubMed PMID: 32871238; PubMed Central PMCID: PMCPCMC7456194. doi:10.1016/j.jsmb.2020.105751.
- [52] Gharebaghi N, Nejadrahim R, Mousavi SJ, Sadat-Ebrahimi SR, Hajizadeh R. The use of intravenous immunoglobulin gamma for the treatment of severe coronavirus disease 2019: a randomized placebo-controlled double-blind clinical trial. *BMC Infect Dis* 2020;20:786. PubMed PMID: 33087047; PubMed Central PMCID: PMCPCMC7576972. doi:10.1186/s12879-020-05507-4.
- [53] Liu K, Zhang W, Yang Y, Zhang J, Li Y, Chen Y. Respiratory rehabilitation in elderly patients with COVID-19: a randomized controlled study. *Complement Ther Clin Pract* 2020;39:101166. PubMed PMID: 32379637; PubMed Central PMCID: PMCPCMC7118596. doi:10.1016/j.ctcp.2020.101166.
- [54] Davoudi-Monfared E, Rahmani H, Khalili H, Hajiabdolbaghi M, Salehi M, Abbasian L, et al. A randomized clinical trial of the efficacy and safety of interferon beta-1a in treatment of severe COVID-19. *Antimicrob Agents Chemother* 2020;64. PubMed PMID: 32661006; PubMed Central PMCID: PMCPCMC7449227. doi:10.1128/AAC.01061-20.
- [55] Song SY, Kim B, Kim I, Kim S, Kwon M, Han C, et al. Assessing reporting quality of randomized controlled trial abstracts in psychiatry: adherence to CONSORT for abstracts: a systematic review. *PLoS One* 2017;12:e0187807. PubMed PMID: 29117269; PubMed Central PMCID: PMCPCMC5678722. doi:10.1371/journal.pone.0187807.
- [56] Speich B, Mc Cord KA, Agarwal A, Gloy V, Gryaznov D, Moffa G, et al. Reporting quality of journal abstracts for surgical randomized controlled trials before and after the implementation of the CONSORT extension for abstracts. *World J Surg* 2019:2371–8. PubMed PMID: 31222645; PubMed Central PMCID: PMCPCMC6722149. doi:10.1007/s00268-019-05064-1.
- [57] Narayan VM, Cone EB, Smith D, Scales CD Jr, Dahm P. Improved reporting of randomized controlled trials in the urologic literature. *Eur Urol* 2016;70:1044–9. PubMed PMID: 27503840. doi:10.1016/j.eururo.2016.07.042.
- [58] Ghimire S, Kyung E, Lee H, Kim E. Oncology trial abstracts showed suboptimal improvement in reporting: a comparative before-and-after evaluation using CONSORT for abstract guidelines. *J Clin Epidemiol* 2014;67:658–66. PubMed PMID: 24439069. doi:10.1016/j.jclinepi.2013.10.012.
- [59] Chhapola V, Tiwari S, Brar R, Kanwal SK. Reporting quality of trial abstracts-improved yet suboptimal: a systematic review and meta-analysis. *J Evid Based Med* 2018;11:89–94. PubMed PMID: 29460397. doi:10.1111/jebm.12294.
- [60] Guo JW, Iribarren SJ. Reporting quality for abstracts of randomized controlled trials in cancer nursing research. *Cancer Nurs* 2014;37:436–44. PubMed PMID: 24406384. doi:10.1097/NCC.000000000000112.
- [61] Kumar S, Mohammad H, Vora H, Kar K. Reporting quality of randomized controlled trials of periodontal diseases in journal abstracts—a cross-sectional survey and bibliometric analysis. *J Evid Based Dent Pract* 2018;18:130–41. PubMed PMID: 29747793. doi:10.1016/j.jebdp.2017.08.005.
- [62] Khan MS, Shaikh A, Ochani RK, Akhtar T, Fatima K, Khan SU, et al. Assessing the quality of abstracts in randomized controlled trials published in high impact cardiovascular journals. *Circ Cardiovasc Qual Outcomes* 2019;12:e005260. PubMed PMID: 31030545; PubMed Central PMCID: PMCPCMC7457741. doi:10.1161/CIRCOUTCOMES.118.005260.
- [63] Mozetic V, Leonel L, Leite Pacheco R, de Oliveira Cruz Latorraca C, Guimaraes T, Logullo P. Reporting quality and adherence of randomized controlled trials about statins and/or fibrates for diabetic retinopathy to the CONSORT checklist. *Trials* 2019;20:729. PubMed PMID: 31842982; PubMed Central PMCID: PMCPCMC6916100. doi:10.1186/s13063-019-3868-4.
- [64] Gallo L, Wakeham S, Dunn E, Avram R, Thoma A, Voineskos S. The reporting quality of randomized controlled trial abstracts in plastic surgery. *Aesthet Surg J* 2020;40:335–41. PubMed PMID: 31353409. doi:10.1093/asj/sjz199.
- [65] Janackovic K, Puljak L. Reporting quality of randomized controlled trial abstracts in the seven highest-ranking anesthesiology journals. *Trials* 2018;19:591. doi:10.1186/s13063-018-2976-x. PubMed PMID: 30373644; PubMed Central PMCID: PMCPCMC6206658.
- [66] Hays M, Andrews M, Wilson R, Callender D, O'Malley PG, Douglas K. Reporting quality of randomized controlled trial abstracts among high-impact general medical journals: a review and analysis. *BMJ Open* 2016;6:e011082. PubMed PMID: 27470506; PubMed Central PMCID: PMCPCMC4985789. doi:10.1136/bmjopen-2016-011082.
- [67] Lockyer S, Hodgson R, Dumville JC, Cullum N. "Spin" in wound care research: the reporting and interpretation of randomized controlled trials with statistically non-significant primary outcome results or unspecified primary outcomes. *Trials* 2013;14:371. PubMed PMID: 24195770; PubMed Central PMCID: PMCPCMC3832286. doi:10.1186/1745-6215-14-371.
- [68] Jellison S, Roberts W, Bowers A, Combs T, Beaman J, Wayant C, et al. Evaluation of spin in abstracts of papers in psychiatry and psychology journals. *BMJ Evid Based Med* 2019:178–81. PubMed PMID: 31383725. doi:10.1136/bmjebm-2019-111176.
- [69] Latronico N, Metelli M, Turin M, Piva S, Rasulo FA, Minelli C. Quality of reporting of randomized controlled trials published in Intensive Care Medicine from 2001 to 2010. *Intensive Care Med* 2013;39:1386–95. PubMed PMID: 23743522. doi:10.1007/s00134-013-2947-3.
- [70] Turrentine M. It's all how you "spin" it: interpretive bias in research findings in the obstetrics and gynecology literature. *Obstet Gynecol* 2017;129:239–42. PubMed PMID: 28079777. doi:10.1097/AOG.0000000000001818.

- [71] Shaqman M, Al-Abedalla K, Wagner J, Swede H, Gunsolley JC, Ioannidou E. Reporting quality and spin in abstracts of randomized clinical trials of periodontal therapy and cardiovascular disease outcomes. *PLoS One* 2020;15:e0230843 PubMed PMID: 32302309; PubMed Central PMCID: PMC7164582. doi:[10.1371/journal.pone.0230843](https://doi.org/10.1371/journal.pone.0230843).
- [72] Reynolds-Vaughn V, Riddle J, Brown J, Schiesel M, Wayant C, Vas-sar M. Evaluation of spin in the abstracts of emergency medicine randomized controlled trials. *Ann Emerg Med* 2019;423–31 PubMed PMID: 31101371. doi:[10.1016/j.annemergmed.2019.03.011](https://doi.org/10.1016/j.annemergmed.2019.03.011).