What are the long-term symptoms and complications of COVID-19: a protocol for a living systematic review [version 2; peer review: 2 approved]

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Open Peer Review

Invited Reviewers
1
2
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report
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version 1
14 Dec 2020
report
report

1. Johannes Siegrist, University of Duesseldorf,

Abstract
Although the majority of people with Covid-19 will experience mild to moderate symptoms and will recover fully, there is now increasing evidence that a significant proportion will experience persistent symptoms for months after the acute phase of the illness. These symptoms include, among others, fatigue, problems breathing, lack of smell and taste, headaches, and depression and anxiety. It is also clear the virus has lasting fluctuating multiorgan sequelae, including affecting not only the respiratory system but also the heart, liver, and nervous system.

We present a protocol for a living systematic review that aims to synthesize the evidence on the prevalence and characteristics of post-acute COVID-19.

The living systematic review will be updated regularly, approximately
every 6 months, as new evidence emerges. We will include studies that follow up at least 100 people with Covid-19 at 12 or more weeks post Covid-19 onset, with no restrictions regarding country, setting, or language. We will use descriptive statistics and, for outcomes reported in two or more studies, we will use meta-analyses to estimate prevalence with 95% confidence intervals (CIs) using the exact method. Heterogeneity between estimates will be assessed using the I² statistic. Our findings will also be presented as infographics to facilitate transcription to lay audiences. Ultimately, we aim to support the work of policy makers, practitioners, and patients when planning rehabilitation for those recovering from Covid-19.

The protocol has been registered with PROSPERO (CRD42020211131, 25/09/2020).

**Keywords**

Living systematic review, COVID-19, long covid, lasting effects

This article is included in the Disease Outbreaks gateway.

This article is included in the Coronavirus collection.

This article is included in the Living Evidence collection.
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Author roles: Michelen M: Data Curation, Formal Analysis, Investigation, Methodology, Validation, Writing – Review & Editing; Cheng V: Data Curation, Formal Analysis, Investigation, Methodology, Validation, Writing – Review & Editing; Manoharan L: Data Curation, Formal Analysis, Methodology, Validation, Writing – Review & Editing; Elkheir N: Data Curation, Investigation, Methodology, Validation, Writing – Review & Editing; Dagens D: Data Curation, Investigation, Methodology, Writing – Review & Editing; Hastie C: Investigation, Validation, Writing – Review & Editing; O'Hara M: Investigation, Validation, Writing – Review & Editing; Suett JC: Investigation, Methodology, Validation, Writing – Review & Editing; Dahmash D: Data Curation, Formal Analysis, Validation, Writing – Review & Editing; Bugaeva P: Data Curation, Formal Analysis, Validation, Writing – Review & Editing; Rigby I: Data Curation, Formal Analysis, Validation, Writing – Review & Editing; Harriss E: Data Curation, Writing – Review & Editing; Munblit D: Formal Analysis, Validation, Writing – Review & Editing; Foote C: Investigation, Validation, Writing – Review & Editing; Scott J: Writing – Review & Editing; Carson G: Funding Acquisition, Validation, Writing – Review & Editing; Olliaro P: Funding Acquisition, Validation, Writing – Review & Editing; Sigfrid L: Funding Acquisition, Methodology, Validation, Writing – Review & Editing; Stavropoulou C: Conceptualization, Data Curation, Formal Analysis, Investigation, Methodology, Project Administration, Resources, Supervision, Validation, Writing – Original Draft Preparation, Writing – Review & Editing

Competing interests: No competing interests were disclosed.

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Background
The range of documented Covid-19 infections vary from asymptomatic to severe, but the vast majority of patients experience mild to moderate symptoms and do not require hospitalisation. We have previously conducted a rapid review of the literature to identify which symptoms and signs might differentiate mild and moderate from severe Covid-19. Since then, and as more data are being gathered, there is increasing evidence of a “long-tail” of Covid-19 illness, but limited information about the range and duration of symptoms experienced or longer term health complications. A community app developed at King’s College London, which tracks self-reported symptoms, has shown that about one in ten will be sick for three weeks or more (https://covid.joinzoe.com/post/covid-long-term). Some individuals with Covid-19 have reported “fatigue, headaches and tingling nerves” that lasted months after symptom onset. A recent longitudinal cohort of 143 patients followed after hospitalisation from Covid-19 in Italy reported that 87% had at least one ongoing symptom, most (55%) reporting three or more, at 60 day follow up. Fatigue (53%), dyspnoea (43%), joint pain (27%) and chest pain (22%) were the most common ongoing symptoms, but there is a variety of other symptoms and complications that have been reported including neurocognitive difficulties, muscle pains and weakness, gastrointestinal upset, rashes, metabolic disruption, thromboembolic conditions and mental health conditions. A prolonged course of illness has also been reported among people with mild Covid-19 who did not require hospitalisation.

The evidence to date remains fragmented as to the onset of symptoms and clinical features, how long symptoms may last, how this relates to the severity of the initial illness, and further lasting impacts to health. A better understanding of patients’ projected recovery from Covid-19 is helpful to patients, healthcare professionals, policymakers and commissioners. The clinical management of persisting symptoms of Covid-19 has started to be addressed in the clinical literature and NHS England has issued guidance for the multisystem needs of patients recovering from Covid-19. Our findings could help identify people requiring additional rehabilitation services and, where necessary, specialist referral to establish a secondary cause of their symptoms. Our findings will also be relevant to organisations such as NHS England, which have recently launched an online Covid-19 rehab service supporting patients suffering long-term effects of the disease (https://www.yourcovidrecovery.nhs.uk/) or the British Society of Immunologists, which recently released a briefing note recommending research into the long-term immunological health consequences of Covid-19.

The aim of this review is to synthesize and continually update the evidence on the characteristics, including prevalence and duration of symptoms and clinical features of post-acute COVID-19, as well as risk factors for developing Long Covid. This will inform clinical and public health management, prevention, and rehabilitation policies.

Methods
To address the aim of this study we will conduct a living systematic review (LSR). LSRs are used in areas where research evidence is emerging rapidly, current evidence is uncertain, and new research may influence policy or practice decisions. These are all features of Covid-19 research, where much about the long-term effects of the disease are still unknown and policy makers are calling for more evidence. The review will be updated approximately every six months, with update cycles under continuous review as the pace of new evidence generated develops through the pandemic. We aim to continue to update the review for up to two years. Our study methodology has been developed and strengthened through consultation with Long Covid Support (a patient support network).

Inclusion/exclusion criteria
We will include studies that meet the follow criteria:
- Studies following up with at least 100 people with suspected, laboratory confirmed, and/or clinically diagnosed Covid-19
- Studies assessing symptoms or outcomes at 12 or more weeks post Covid-19 onset
- Peer reviewed articles published since 1 January 2020
- No restriction regarding country, setting, or language

We will exclude:
- Studies that focus only on acute Covid-19
- Editorials and opinion papers

Search strategy
A search of the following databases will be conducted: Pubmed and CINAHL through the EBSCO database host for
general health peer-reviewed articles and Global Health for
global peer-reviewed articles through the Ovid database host.
In addition, we will search Google Scholar for grey literature.
We will also conduct complementary searches using the WHO
Global Research Database on Covid-19 and LitCOVID as
two databases that bring together evidence on Covid-19
from a worldwide dataset. A ‘backwards’ snowball search
will be conducted for the references of systematic reviews.
Finally, we will contact experts in the field and use social
media to identify relevant studies.

We will search using controlled subject headings and key-
words of the following concepts: Terms related to 1) COVID-19
OR COVID OR SARS-CoV-2; 2) symptoms OR clinical fea-
tures OR signs OR characteristics OR sequelae OR complica-
tions; 3) long-term OR post-acute OR long-tail OR persistent OR
chronic COVID OR long COVID OR post discharge OR pro-
longed symptoms OR long haul. The search terms were piloted
on Pubmed and CINAHL through the EBSCO database host
the week starting 14th September 2020 to ensure that high
profile research articles on long covid were included. No
important studies were missed.

An example is shown below:

| MEDLINE Search (16/3/2021) | 1952 |
|----------------------------|------|
| S1. COVID-19 OR OR covid OR SARS-CoV-2. ab |      |
| S2. symptom* OR "clinical features" OR signs OR characteristic* OR sequelae OR complication*.ab |      |
| S3. "long-term Covid" OR long-term N2 consequence* OR "long-term impact" OR "long-term effect" OR "post-
acute" OR long-tail OR persist* OR "chronic-COVID" OR "long-COVID" OR post-discharge OR postdischarge OR"prolonged symptom" OR "long-haul".ab |      |
| S4. S1 AND S2 AND S3 |      |

Screening
Search results will be managed and screened using a review
online platform, Rayyan12. Initial screening of titles and abstracts
as well as full text screening against the inclusion criteria
will be done by two reviewers independently. Non-English arti-
cles will be translated using Google Translate or reviewed by
a reviewer with good knowledge of the language. Disagree-
ments for inclusion will be resolved by consensus. Where disa-
grreements cannot be resolved, a third researcher will review
the papers to make the final decision.

Risk of bias
We will be using the Hoy et al. checklist13 to critically
appraise the studies included in the review, a validated tool for
assessing risk of bias in prevalence studies.

Data extraction
The following information will be extracted from each study
based on an extraction form informed by a previous review:

Outcomes
The primary outcome is to characterise the prevalence of symp-
toms and complications of long term Covid-19 in different
populations. Secondary outcomes include diagnostics and risk
factors for developing different sequelae.

Data analysis
We will use descriptive analysis, and present proportion of
symptoms and estimate their 95% confidence intervals (CIs)
using the exact method. When more than two studies provide
information on a symptom, we will perform a meta-analysis
using a random intercept logistic regression model. Heterogeneity
between estimates will be assessed using the I² statistic.

Where data is available, we will explore key factors that affect
prevalence estimates, e.g. hospitalisation, settings, location of
the study, sex and follow-up timing using subgroup analysis and
meta-regression analysis. We will identify these factors by
discussing with our clinical experts and patient advocates and
by reviewing the literature. The division for key factors used in
subgroup analysis will depend on the availability of reported
data across studies. We will align the division with the lit-
erature and expert opinions to form exploratory analysis and
to help the interpretation.

We will also conduct sensitivity analysis to examine the impact
of high risk of bias studies and conventional statistical meth-
ods on the prevalence estimates, e.g. Freeman-Tukey Double
arc sine transformation using inverse variance meta-analysis.
All analysis and data presentation will be performed using
meta and ggplot2 in R (version 4.0.5 or above) via RStudio
(version 1.3.1093 or above).

We will work with patient advocates to present the data to
facilitate transcription to lay audiences.

Protocol registration
This protocol report is structured according to the Preferred
Reporting Items for Systematic Reviews and Meta-Analyses
Protocols (PRISMA-P) statement guidelines14, was registered
with PROSPERO (CRD42020211131, 25 September 2020).
The protocol will be updated as we progress with the living
review as and if needed. CS is the guarantor for this study.

Data availability
Underlying data
No underlying data are associated with this article.

Reporting guidelines
Figshare: PRISMA-P checklist for “What are the long-term
symptoms and complications of COVID-19: a protocol for a
living systematic review”. https://doi.org/10.25383/city.13187456.
v115.
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15. Michelen M, Sigfrid L, Manoharan L, et al.: What are the long-term symptoms and complications of COVID-19: a protocol for a living systematic review. City, University of London. Journal contribution. 2020.
   http://www.doi.org/10.25383/city.13187456.v1
Open Peer Review

Current Peer Review Status: 

Version 2

Reviewer Report 22 September 2021
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✔ Madelon van Wely
Centre for Reproductive Medicine, Amsterdam University Medical Centre, University of Amsterdam, Amsterdam, The Netherlands

In my view all queries were answered and adequately addressed, the protocol quality has improved and I am content with the present version. I wish the authors good luck with their review.

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Epidemiology, meta-analyses, Obstetrics and Gynaecology

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Reviewer Report 31 August 2021
https://doi.org/10.5256/f1000research.58402.r92985

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✔ Johannes Siegrist
Senior Professorship on Work Stress Research, University of Duesseldorf, Duesseldorf, Germany

I have read the revision of the manuscript and approve the publication of this manuscript.

Competing Interests: No competing interests were disclosed.
Reviewer Expertise: Social epidemiology systematic reviews

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Version 1

Reviewer Report 22 March 2021

https://doi.org/10.5256/f1000research.30148.r80913

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Madelon van Wely
Centre for Reproductive Medicine, Amsterdam University Medical Centre, University of Amsterdam, Amsterdam, The Netherlands

No doubt this is an important LSR that needs to be performed. I am in favor of getting this protocol published, however, at present many details are lacking. In my view in LSRs it is crucial to decide beforehand what the authors plan to do. Though it may seem annoying now, it will be helpful in the end.

1. The outcomes. You performed a rapid review so you will know what main outcomes you are looking for. The outcomes are not stated in the Prospero protocol either.

   2. It would also be advisable to have list of inclusion criteria and data you want to extract like risk factors and demographics. You could combine these with outcomes and present them in one table.

3. Data analysis: descriptive analysis sub grouped by age, sex, comorbidities? As the idea is to provide proportions from different studies I think you will be able to pool these, stratified by age etc. You could for instance summarise proportions (maybe use Freeman-Tukey transformation to stabilise variances in case of studies with zero events?), provide 95% CIs and predictive intervals to report on the precision of estimates. (meta-analysis of proportions)

4. About the subgrouping. How are you planning to do so? Are the age groups pre-specified or will they be dichotomised of subgrouped on basis of distribution of the data?

5. The duration is very important, how do you intend to present time? As grouped or continuous variable? When you have adequate data you could present the prevalence estimates over time.

6. Besides prevalence studies you might also include cohorts in which case you may want
to assess the risk of bias using the Newcastle Ottawa Scale for comparative cohorts or another such checklist.

7. Search: I would also search specifically for the expected outcomes

8. I do advise to clarify when you consider a symptom to be persistent? You write: persistent mild, moderate or severe symptoms as defined by the article authors. I suppose post-discharge is T0.

9. How often do you at this moment intend to update and publish the results?

Is the rationale for, and objectives of, the study clearly described?
Partly

Is the study design appropriate for the research question?
Yes

Are sufficient details of the methods provided to allow replication by others?
No

Are the datasets clearly presented in a useable and accessible format?
Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Epidemiology, meta-analyses, Obstetrics and Gynaecology

I confirm that I have read this submission and believe that I have an appropriate level of expertise to state that I do not consider it to be of an acceptable scientific standard, for reasons outlined above.

Author Response 03 Jul 2021
Charitini Stavropoulou, City, University of London, London, UK

Thank you for your thorough review, please find an outline of the response to the comments raised.

1. The outcomes. You performed a rapid review so you will know what main outcomes you are looking for. The outcomes are not stated in the Prospero protocol either.

Response: Thanks for highlighting, we have updated the text in the methodology section to highlight that the primary outcome is to characterise the prevalence of symptoms and complications of long term Covid-19 in different populations. Secondary outcomes include diagnostics and risk factors for developing different sequelae. This is now stated on page 7 of the revised protocol.

1. It would also be advisable to have list of inclusion criteria and data you want to extract like risk factors and demographics. You could combine these with outcomes and present them in one table.
Response: We have provided further details in the methods, regarding the inclusion/exclusion criteria. We have also expanded on the data extraction section to show the clarify the data we will be extracting. Please see page 7 of the revised protocol.

1. Data analysis: descriptive analysis sub grouped by age, sex, comorbidities? As the idea is to provide proportions from different studies I think you will be able to pool these, stratified by age etc. You could for instance summarise proportions (maybe use Freeman-Tukey transformation to stabilise variances in case of studies with zero events?), provide 95% CIs and predictive intervals to report on the precision of estimates. (meta-analysis of proportions)

Response: We have provided further information in our analysis section to clarify that the analysis will go beyond descriptive statistics and that we will perform meta-analysis wherever this is possible, i.e. when there are more than two studies providing information on a symptom. Please see pages 7 and 8 of the revised protocol.

1. About the subgrouping. How are you planning to do so? Are the age groups pre-specified or will they be dichotomised of subgrouped on basis of distribution of the data?

Response: We plan to conduct subgroup analysis to explore the influence of key factors, e.g. age, on the estimates of prevalence. We will identify these key factors by discussing with our clinical experts, patient advocates and by reviewing the literature. We will try to align the division for subgroups with the literature to help inform the analysis and results. However, the methods will heavily depend on the availability and distribution of data. We have updated the methodology section accordingly.

1. The duration is very important, how do you intend to present time? As grouped or continuous variable? When you have adequate data you could present the prevalence estimates over time.

Response: We now clarify in the methods section that a subgroup analysis will be performed on follow up timing as indeed this is an important parameter as a group variable.

1. Besides prevalence studies you might also include cohorts in which case you may want to assess the risk of bias using the Newcastle Ottawa Scale for comparative cohorts or another such checklist.

Response: We will include any study design including cohort studies with symptom prevalence. We have chosen to use the Hoy et al risk of bias assessment checklist, a validated tool for assessing risk of bias in prevalence studies, which does not prevent us from using it with cohort studies.

1. Search: I would also search specifically for the expected outcomes

Thank you for your suggestion. We have conducted pilot searches and noticed considerable diversity in the reported outcomes. The inclusion of key terms for the expected outcomes in search strings generated excessive hits and contributed to low precision, aka a low percentage of useful publications found in the search results. It also limits searches by “expected” outcomes and loses the opportunities finding new types of long covid symptoms and risk factors. Therefore, we used a search strategy using simple search strings focusing on describing long covid and complement the searches by backwards snowball searches in the references of relevant publications and consulting experts.

1. I do advise to clarify when you consider a symptom to be persistent? You write: persistent mild, moderate or severe symptoms as defined by the article authors. I
suppose post-discharge is T0.

**Response.** Thanks for highlighting. We have provided further clarification in the inclusion criteria (page 5). In light of the new definitions issued by the Office for National Statistics (ONS), we clarify we will be including studies assessing symptoms or outcomes at 12 or more weeks post Covid-19 onset.

1. How often do you at this moment intend to update and publish the results?

**Response:** We will update the review every 6 months, as now clarified on page 5 of the methods section.

**Competing Interests:** No competing interests were disclosed.

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**Reviewer Report 12 February 2021**

https://doi.org/10.5256/f1000research.30148.r78755

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**Johannes Siegrist**
Senior Professorship on Work Stress Research, University of Duesseldorf, Duesseldorf, Germany

This protocol paper describes the aims and methods of conducting a living systematic review (LSR) on findings from studies following COVID-19 patients with long-term symptoms. The LSR is planned to be updated at 2-month intervals up to 2 years. The paper offers information on search strategy with key terms, data management procedure, screening with critical appraisal, data extraction, and strategies of data analysis. Overall, the approach is in line with the PRISMA guidelines of performing systematic reviews and meta-analyses. Moreover, the protocol was registered in PROSPERO. Critical appraisal was proposed according to Hoy *et al*. 2012, and authors are expected to ensure that relevant criteria of risk of bias can be assessed by this tool.

In summary, while the indexing of the paper is endorsed, several minor queries still need to be addressed, as detailed below.

**Search strategy:**
Authors claim that their search will not be restricted by country and language. However, it is unrealistic to identify and analyse written materials in more than 3 or 4 main languages, and these languages (English plus...) should be explicitly mentioned in the study protocol.

**Data extraction:**
I wonder why no data on treatment (therapy of COVID-19 symptoms) are included in the data extraction matrix. This information is crucial if aspects such as symptom severity and duration are being evaluated.

**Data analysis:**
This section is not well elaborated. Basic descriptive statistics only are mentioned. However, systematized data from respective publications will probably allow additional, more informative ways of synthesizing the data.

Is the rationale for, and objectives of, the study clearly described?
Yes

Is the study design appropriate for the research question?
Yes

Are sufficient details of the methods provided to allow replication by others?
Partly

Are the datasets clearly presented in a useable and accessible format?
Not applicable

**Competing Interests:** No competing interests were disclosed.

**Reviewer Expertise:** Social epidemiology systematic reviews

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

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**Author Response 03 Jul 2021**

Charitini Stavropoulou, City, University of London, London, UK

Thank you for your thorough review of our protocol, please find the response to the comments made outlined below.

**Search strategy:**
Authors claim that their search will not be restricted by country and language. However, it is unrealistic to identify and analyse written materials in more than 3 or 4 main languages, and these languages (English plus...) should be explicitly mentioned in the study protocol.

**Response:** The main search will be performed in English, articles identified in non-English languages will be translated using Google translate and assessed by a reviewer with good knowledge of the language. This is now stated on page 7 of the revised protocol.

**Data extraction:**
I wonder why no data on treatment (therapy of COVID-19 symptoms) are included in the data extraction matrix. This information is crucial if aspects such as symptom severity and duration are being evaluated.

**Response:** Thank you for the comment. This is part of the information included in the characteristics of population and is now explicitly mentioned in the protocol. Where data is
available and synthesizable, we plan to assess the association between acute treatment and symptom severity and duration.

**Data analysis:**
This section is not well elaborated. Basic descriptive statistics only are mentioned. However, systematized data from respective publications will probably allow additional, more informative ways of synthesizing the data.

**Response:** Thank you for your comments. We have amended the data analysis section to clarify that the analysis will go beyond descriptive statistics and, when possible, we will perform a meta-analysis. Please see pages 7 and 8 of the revised protocol.

**Competing Interests:** No competing interests were disclosed.