Protocol for the development of a reporting guideline for causal and counterfactual prediction models in biomedicine

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ABSTRACT

Introduction While there are guidelines for reporting on observational studies (e.g., Strengthening the Reporting of Observational Studies in Epidemiology, Reporting of Studies Conducted Using Observational Routinely Collected Health Data Statement), estimation of causal effects from both observational data and randomised experiments (e.g., A Guideline for Reporting Mediation Analyses of Randomised Trials and Observational Studies, Consolidated Standards of Reporting Trials, PATH) and on prediction modelling (e.g., Transparent Reporting of a multivariable prediction model for Individual Prognosis or Diagnosis), none is purposely made for deriving and validating models from observational data to predict counterfactuals for individuals on one or more possible interventions, on the basis of given (or inferred) causal structures. This paper describes methods and processes that will be used to develop a Reporting Guideline for Causal and Counterfactual Prediction Models (PRECOG).

Methods and analysis PRECOG will be developed following published guidance from the Enhancing the Quality and Transparency of Health Research (EQUATOR) network and will comprise five stages. Stage 1 will be meetings of a working group every other week with rotating external advisors (active until stage 5). Stage 2 will comprise a systematic review of literature on counterfactual prediction modelling for biomedical sciences (registered in Prospective Register of Systematic Reviews). In stage 3, a computer-based, real-time Delphi survey will be performed to consolidate the PRECOG checklist, involving experts in causal inference, epidemiology, statistics, machine learning, informatics and protocols/standards. Stage 4 will involve the write-up of the PRECOG guideline based on the results from the prior stages. Stage 5 will seek the peer-reviewed publication of the guideline, the scoring/systematic review and dissemination.

Ethics and dissemination The study will follow the principles of the Declaration of Helsinki. The study has been registered in EQUATOR and approved by the University of Florida’s Institutional Review Board (202200495). Informed consent will be obtained from the working groups and the Delphi survey participants. The dissemination of PRECOG and its products will be done through journal publications, conferences, websites and social media.

Strengths and limitations of this study

⇒ There are no guidelines for the reporting of data-learnt prediction models that have the specific intent to calculate alternative scenarios (counterfactuals) and identify individualised effects of interventions.
⇒ Prediction of Counterfactuals Guideline (PRECOG) will fill a gap in reporting standards for counterfactual prediction modelling and will capitalise on the systematisation and quality of the Enhancing the Quality and Transparency of Health Research network.
⇒ PRECOG will be built on diverse (clinical researchers, computer scientists, epidemiologists, statisticians) expertise consensus across multiple development stages.
⇒ Even with rigorous study design, execution and reporting standard, causal claims made on observational data analyses might be still mistaken by wrong assumptions or unmeasured, hidden bias.

INTRODUCTION

The increasing availability of large electronic health record data has led to an explosion in the development of prediction models—both traditional statistics and machine learning—for diagnostic, prognostic and treatment optimisation purposes. Despite the availability of reporting guidelines, for example, ‘Transparent Reporting of a multivariable prediction model for Individual Prognosis or Diagnosis’ (TRIPOD),\textsuperscript{1} the quality of many studies is low, as well as adherence to reporting standards, and there is often a misinterpretation of the models’ operating capabilities, with possible misuse and harm at the individual and/or population level.\textsuperscript{2,3} One of the most common mistakes\textsuperscript{4,5} is to consider a prediction model readily usable for interventions on individuals, by changing certain variables with the intent to improve outcomes, that is, calculating alternative scenarios or so-called counterfactuals. Since prediction
models are often learnt from observational data, there
is no guarantee that the strongest predictors are causing
the outcome of interest and are not confounded, medi-
ated by others, or actually concomitant causes of it. While
such bias is not a problem for mere prediction in similar
populations—since variables are not being changed with
the intent to modify risk—it becomes problematic in new,
out-of-distribution populations (even when cross-
validation performance is high), and when trying to optimise outcomes.

Thus, formal causal assessment is needed when develop-
oping prediction models on observational data to be used
for alternative scenarios and interventions, that is, counterfactual prediction models. The approaches from traditional statistics, computational science and econometrics, including the potential outcomes framework, do-calculus and directed acyclic graphs, are often focused on estimating a population-level causal effect for
a single intervention query (treatment or exposure) but
can be used to calculate individualised treatment effects
and counterfactuals. Machine learning has also been
employed for counterfactual prediction. Several off-
the-shelf methodologies have been revisited, including
deep learning and random forests.

Given the rise in counterfactual prediction modelling
studies, there is a need for common grounds on model
reporting, to improve overall quality (although adhering
to a protocol might be necessary, yet not sufficient condi-
tion to study quality), and specifically on transparency
and reproducibility of results.

In the ‘Enhancing the Quality and Transparency of
Health Research’ (EQUATOR) network (https://www.
equator-network.org/), there are guidelines specifically
designed for reporting causal effects on randomised clinical
trials (RCTs), for example, ‘Consolidated Standards of
Reporting Trials’ and ‘A Guideline for Reporting Mediation Analyses of Randomised Trials and Observa-
tional Studies’. Reporting guidelines for observational
studies also mention causal effects inference, for example, ‘Strengthening the Reporting of Observational
Studies in Epidemiology Using Mendelian Randomisation’, ‘Reporting of Studies Conducted Using Observa-
tional Routinely Collected Health Data Statement for
Pharmacoepidemiology’, and the ‘Instrumental Variable Methods in Comparative Safety and Effectiveness Research’. Outside of EQUATOR, the Patient-Centered Outcomes Research Institute (PCORI) (https://www.
pcori.org/) provides ‘Standards for Causal Inference
Methods in Analyses of Data from Observational and
Experimental Studies in Patient-Centered Outcomes Research’. Also, there are guidelines for estimating
causal effects in pragmatic randomised trials. Worth
noting is the ‘predictive approaches to treatment effect heterogeneity’ (PATH) statement, which—although focused on RCTs—examines treatment effect heteroge-
enity by considering as effect modifier(s) either the risk
or the covariates, with both strategies aimed at guiding
treatment decisions. PATH provides guidance for specific
multivariable regression configurations and warns against
more ‘aggressive’ approaches (eg, machine learning
models with many df) that could bring overfitting. Overall, existing guidelines are not well fitted for causal
and counterfactual prediction modelling for observational biomedical data (or a mixture of RCTs and observ-
ational), although a number of them contain elements that are directly related.

Consequently, we aim to develop a new reporting
guideline, which we tentatively name as PRECOG—
acronym for ‘Prediction of Counterfactuals Guideline’. The primary focus of PRECOG is to provide guidance
on how to report causal assumptions as well as evaluate
derivation/validation of models—involving at least an
observational data source—that provide predictions of
individualised treatment/intervention effects in the form
of potential outcomes. On the one hand, the develop-
ment of these models can follow both risk-and-effect
modelling approaches as in PATH, but it is intended to
be more general, allowing any functional form and data
generation process. On the other hand, the validation
standard of these models falls within the TRIPOD scopes,
but it also evaluates how they are suitable for optimisation (eg, treatment decision, risk reduction) in addition to
diagnosis and prognosis, trusting on the counterfactuals
backed up by the causal claims. PRECOG is also expected
to provide guidance on software implementation and
interoperability. As a quality evaluation instrument,
PRECOG can help researchers (and general readers, peer reviewers, journal editors) as well as policy-makers
to carry out and critically appraise causal and counter-
factual prediction modelling studies. We anticipate
further expansion of the guideline for specific areas, for
example, pharmaceutical interventions. The primary use
cases of PRECOG are expected to fall within biomedical
sciences, but they could be applied to other fields such as
psychology or economics.

METHODS AND ANALYSIS
PRECOG will be developed following published guidance
from the EQUATOR network. We will develop the guide-
line in five stages, as shown in figure 1: (1) meeting of a
working group every other week; (2) scoping/systematic
review of causal and counterfactual prediction modelling
studies; (3) reporting checklist draft and real-time Delphi
exercise; (4) development of the final guideline and (5)
peer-review, publication and dissemination. These stages
are drawn from prior, successful development studies,
in primis the protocol used for the making of TRIPOD-
Artificial Intelligence (AI) and Prediction model Risk Of
Bias ASsessment Tool (PROBAST)-AI. The expected
timeline for stages 1–4 is 1 year, using 6–9 months for
stages 1–2, and 3–6 months for stages 3–4.

Stage 1: working group setup and meetings
The core working group is composed of the coauthors
of this protocol description, who met every other week

Xu J, et al. BMJ Open 2022;12:e059715. doi:10.1136/bmjopen-2021-059715
The purpose of the literature review is twofold: (1) to build a knowledge base on study design, methodological approaches, use cases and reporting commonalities among causal inference and counterfactual prediction studies in biomedical sciences; and (2) to help the development of reporting items for PRECOG. A subset of the working group members will concentrate on the review. Lin et al.34 published a scoping review on causal methods for predictions under hypothetical interventions, screening nearly 5000 papers and focusing on 13 key articles, including traditional statistical as well as machine learning modelling. Most works used marginal structural models and g-computation. The authors concluded that ‘techniques for validating causal prediction models’ are still in their infancy. Based on the results from the scoping review, and expanding the search strategy and the article sources, the team is going to move forward with a systematic review. The review will provide counts on methodology, review and applied papers, but then will focus on works that include at least one observational data source and an application use case, further deepening the validation strategies. The planned reporting statement of choice is the ‘Preferred Reporting Items for Systematic Reviews and Meta-Analyses’,35 and the working group will register the work in the ‘Prospective Register of Systematic Reviews’.36

As part of the review, we foresee discussing how to assess the potential risk of bias (which can lead to misuse and patients’ harm), and if current tools such as ‘PROBAST’ are appropriate.37

Stage 3: real-time Delphi exercise
We will conduct a real-time Delphi survey38 to review and refine the items of the PRECOG reporting checklist. Participants will be identified initially through the professional network of the core working group and of the external advisors, and further via literature search (including but not limited to the existing scoping review and the planned systematic review), social media screening and snowballing by the active participants. As for the expanded working group composition, participants will be invited from diverse and multicultural backgrounds and different countries. Invites will include academics at various career stages, researchers and investigators from non-profit and for-profit organisations, programme officers from national/federal funding agencies, entrepreneurs, healthcare professionals, journal editors, policy-makers, healthcare regulators and end-users of predictive models. The participant selection will be based on area expertise grouping (computer science, biostatistics, biomedical informatics, statistics, epidemiology, standards, causal inference, ethics), used to determine the sample size (discussed below). We choose a computer-based, real-time Delphi,38 since it offers some operational advantages with respect to conventional multi-round Delphi
techniques, for example, responder’s attrition. In brief, real-time Delphi is a ‘roundless’ exercise based on an online survey platform. Participants can access and modify their responses at any time during the survey time frame, and they can view the survey summaries calculated among all responders. In this way, participants can see if/how their opinion is unpopular and add further comments to support their cases.

The working group will develop an initial reporting checklist for PRECOG, based on the EQUATOR developing standard and existing related guidelines/statements. We anticipate that PRECOG will draw substantially from the reporting items of TRIPOD as well as the recommendations of PATH; however, we expect major differences rather than a simple merge. For instance, performance evaluation as recommended in TRIPOD should be modified to include specific metrics such as the Precision Estimation of Heterogeneous Effects, and emphasise out-of-distribution validation. Another important aspect is the causal assumptions. PATH relies on RCTs, where randomisation supports the strong ignorability of treatment assignments, while PRECOG models might be exclusively built on observational data (or a mixture of observational and RCT data) and a justification for causal claims will need to be provided.

An anonymous online survey will be created where each checklist item can be evaluated in relation to its importance and relevance for the guideline, using a five-point Likert scale, and a free text box for comments. Also, at the end of the survey, another text box will allow more generic comments and propositions, for example, new items to be added to the checklist. When a participant consents to participate and completes the survey for the first time, they can view the summary of all responses to date and can access the survey again within the next 6 weeks. The survey is closed after the required sample size is reached, or a maximum of 6 weeks are passed from the last recorded first response.

There is no consensus on the sample size of a Delphi panel but a minimum number of 10–18 panel members per area of expertise has been recommended. We will aim to reach a minimum sample size of 60 considering the aforementioned background expertise areas, compiling a list of 80–100 potential participants for the recruitment. At the end of the Delphi survey, the expanded working group will review the results and consolidate the checklist through a consensus meeting. The working group will also decide on the consensus rule. In general, for items ranked on a five-point Likert scale, the consensus rule is 80%. but there can be differences in how adjacent items are grouped or weighted toward consensus. For instance, Naughton et al. quantified the Likert points from 1 (most important) to 5 (least important), and defined consensus for items scoring a median of 2.5 or less overall, when at least 80% of responders gave 1–3 points. More recent works proposed entropy-based consensus.

Stage 4: development of the guideline and related products
On finalisation of the reporting checklist from the Delphi exercise, the extended working group will develop the full PRECOG guidelines. The manuscript will be posted to a public preprint website, for example, bioRxiv or medRxiv, before submission to a peer-review journal and possibly presented as an abstract/poster in international conferences, for example, the annual conference of the American Medical Informatics Association or the Society for Epidemiology Research. It is expected that the PRECOG initiative will produce at least the following papers:

- Guideline development protocol (this work).
- A systematic review of causal and counterfactual prediction models in biomedical sciences.
- PRECOG guideline.

Stage 5: publication and dissemination plan
After being posted on preprint servers, the aforementioned manuscripts will be submitted to peer-reviewed international journals for final publication. The authors’ list will be determined based on effective individual contributions, following the ‘contributor roles taxonomy’ (CRediT) (https://casrai.org/credit/), and might include additional contributors other than the working group members and external advisors. The dissemination strategy will be discussed during the workgroup meetings. In addition to conferences and publications, it is likely that social media platforms such as Twitter will be leveraged to inform on the PRECOG availability and utility.

Patient and public involvement
This study does not include patients. However, the participants of the working groups—by definition—will be involved in the design of the Delphi survey, in its evaluation, and in the finalisation of the PRECOG guideline (including authorship in papers). The participants of the Delphi survey can provide not only an evaluation of items but suggest new ones and re-evaluate the items during the time when the survey is open.

Acknowledgements
We thank the TRIPOD coauthors Dr G Collins (U Oxford, UK) and Dr KG Moons (UMC Utrecht, NL) and Dr N Peek (U Manchester, UK) for expressing their interest to join the PRECOG working groups.

Contributors
JX wrote and submitted the protocol description. YG performed an initial literature review on reporting standards. FW and HX performed an initial literature review on counterfactual prediction models. RL advised on protocol procedures and ethical review. JB and MP conceived the idea.

Funding
This work has been in part supported by National Institutes of Health (NIH)-National Institute of Allergy and Infectious Diseases (NIAID) grants no. R01AI145552 and R01AI141810 (MP), by National Institute on Aging (NIA) grants no. R33AG062884-03 (RL and MP) and 5R21AG068717-02 (JB and YG), by National Cancer Institute (NCI) grants no. 5R01CA246418-02, 3R01CA246418-02S1, 1R21CA245858-01A1, 3R21CA245858-01A1S1, and 1R21CA233394-01A1 (JB and YG), by the Centers for Disease Control and Prevention (CDC) grant no. U18DP006512 (JB, YG and MP), and by University of Florida Informatics Institute Seed grant.

Competing interests
None declared.

Patient and public involvement
Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.
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