**How to submit your paper**

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Please ensure that all submissions to eBioMedicine follows the guidelines provided for each article type. For instruction on how to format the text of your paper, including tables, figures, panels, and references, please see our Formatting guidelines.

Articles

Reporting Standards

Interventional studies:
We require the registration of all interventional trials, whether early or late phase, in a primary register that participates in WHO’s International Clinical Trial Registry Platform (see Lancet 2007; 369: 1909–11) or in ClinicalTrials.gov, in accord with ICMJE recommendations. We also encourage full public disclosure of the minimum 21-item trial registration dataset at the time of registration and before recruitment of the first participant (see Lancet 2006; 367: 1631–35). The registry must be independent of for-profit interest.

Reports of trials must conform to CONSORT 2010 guidelines and should be submitted with their protocols.

All reports of randomised trials should include a section entitled Randomisation and masking, within the Methods section. Please refer to The Lancet’s formatting guidelines for randomised trials.

Cluster-randomised trials must be reported according to CONSORT extended guidelines.

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Please carefully follow the linked guidelines of reporting standards if your study falls within one the following categories, and fill in and return the checklist(s) where applicable:

Animal preclinical studies .......................................................... ARRIVE
Observational cohort and case-control studies* ....................... STROBE
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Systematic reviews and meta-analyses ..................................... PRISMA
Genetic association studies ...................................................... STREGA
Genetic risk prediction studies ................................................ GRIPS
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Case reports ............................................................................. CARE
Health economic evaluation ..................................................... CHEERS
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- Abstract
- Keywords (4–6)
- Research in Context
- Introduction
Information for Authors

- Methods (including Ethics, Statistics, and Role of Funders)
- Results
- Discussion
- Contributors
- Declaration of Interests
- Acknowledgments
- Data Sharing Statement
- References
- Figure Legends

Title page
Titles should be informative but not excessively detailed or heavy on jargon. Please avoid abbreviations in title. Please either define functionally (e.g., "the influenza viral HA protein") or spell out ("influenza viral hemagglutinin"). A brief title, author name(s), preferred degree (one only), affiliation(s), and full address(es) of the authors must be included. Full names for all authors must be included in the format Julie M. Moore, not Moore J.M. or J.M. Moore. The name and address of the corresponding author should be separately and clearly indicated with email and telephone details.

Abstract
Include an abstract (semi-structured summary), with five paragraphs (Background, Methods, Findings, Interpretation, and Funding), not exceeding 250 words. Our electronic submission system will ask you to copy and paste this section at the "Submit Abstract" stage. For randomised trials, the abstract should adhere to CONSORT extensions: abstracts (see Lancet 2008; 371: 281-83). For intervention studies, the abstract should include the primary outcome expressed as the difference between groups with a confidence interval on that difference (absolute differences are more useful than relative ones). Secondary outcomes can be included as long as they are clearly marked as secondary and all such outcomes are reported.

Keywords
Please provide a short list of keywords.

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All research papers (including systematic reviews/meta-analyses) submitted to eBioMedicine must include a panel putting their research into context with previous work in the format outlined below (see Lancet 2014; 384: 2176–77, for the original rationale). This panel should not contain references; key studies mentioned here should be referenced in the main text.

Evidence before this study
Authors should briefly state: the sources (databases, journal or book reference lists, etc) searched; the criteria used to include or exclude studies (including the exact start and end dates of the search), which should not be limited to English language publications; the search terms used; the quality (risk of bias) of that evidence; and the pooled estimate derived from meta-analysis of the evidence, if appropriate.

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Authors should state whether the results were substantiated by repetition under a range of conditions. Sufficient information about sample collection must be provided to distinguish between independent biological data points and technical replicates.

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Include a statement to indicate approval by appropriate ethics committee on animal and human experimentations.

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- Protein sequence: EMBL-EBI, Protein Data Bank
- Microarray and deep sequencing data: GEO, ArrayExpress
- SNPs and CNVs: dbSNP, DGVa, dbVAR
- Genotypes and phenotypes: dbGaP
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Tables should be provided in an editable Word or Excel format (so individual numbers/texts can be copied). Please ensure that each table fits within one A4-sized page.

Discussion
Please include discussion on limitations, generalisability, and interpretation of results. The Discussion should be no longer than 5 pages of A4 paper. Please do not include subheadings in the Discussion, and please do not repeat a description of the results.

Please conclude with a brief paragraph highlighting main points of study, including a statement regarding the translational value of the work. As with the Introduction and Abstract, please make sure the language is clear to the general audience, including non-specialists.

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From September 21, 2020, all submitted research Articles must contain a data sharing statement, to be included at the end of the manuscript. Data sharing statements must include:

- Whether data collected for the study, including individual participant data and a data dictionary defining each field in the set, will be made available to others (“undecided” is not an acceptable answer);
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Acknowledgements

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With a maximum of 150 words, briefly explain the necessary background and encapsulate the take-home message for a non-specialist reader. Please emphasise the recent developments or novel conclusions, concepts, or models that make your Review timely.

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Please provide 4-6 descriptive keywords.

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Example:

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Data for this Review were identified by searches of MEDLINE, Current Contents, PubMed, and references from relevant articles using the search terms “sentinel node”, “breast cancer”, and “axilla”. Abstracts and reports from meetings were included only when they related directly to previously published work. Only articles published in English between 1980 and 2006 were included.
Information for Authors

References (Reviews)
No more than 75 references, with particular emphasis on literature published in the past 5 years.

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**References**

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