

Instructions to Authors

Thank you for your interest in Annals of Cancer Epidemiology (ACE). Please consult the following instructions to help you prepare your manuscript, and feel free to contact us with any questions. To ensure fast peer review and publication, manuscripts that do not adhere to the following instructions will be returned to the corresponding author for technical revision before undergoing peer review. We are looking forward to your submission.

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1. ABOUT THE JOURNAL

Annals of Cancer Epidemiology (ACE) is an international, peer-reviewed, open-access journal with the goal to help human apply epidemiological studies to fight against cancers by providing an open-access platform and user-friendly facilities to worldwide cancer researchers, clinicians, scientists and policy makers, and shape an effective communication and collaboration amongst them. All submissions and review processes of ACE are conducted

electronically to expedite the reviews and publication process. ACE will spare no effort to minimize the duration of review and publication processes whilst maintaining the highest standards of each, and will maintain innovative efforts to meet readers' need.

The editors and an international advisory group of scientists and clinician-scientists as well as other experts will hold ACE articles to the high-quality standards. ACE welcomes submissions of Original Research (full length and short reports), Review Articles and Editorials to published research.

Journal Abbreviation: Ann Cancer Epidemiol

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2. MANUSCRIPT CATEGORIES

(1) ORIGINAL ARTICLE

Word limit: 5,000 words maximum including abstract but excluding references, tables and figures.

Abstract: 450 words maximum, with sub-headers (background, methods, results and conclusions).

References: no maximum.

Figures/ tables: no maximum, but 8 figures should be sufficient.

Description: Originality and clinical impact are essential for acceptance of Original Articles. Such an article is to present original basic science or clinical research findings by the authors in the field of cancer epidemiology. Systematic review with meta-analysis in ACE is addressed as original article. The abstract should contain the following subheadings: Background, Methods, Results and Conclusions. Original articles should entail a section describing the contribution of each author to the manuscript as well as Statement of Ethics Approval. See

section “AUTHORS’ CONTRIBUTION” and section “STATEMENT OF ETHICS APPROVAL” for details.

(2) REVIEW ARTICLE

Word limit: 6,000 words maximum including abstract but excluding references, tables and figures.

Abstract: 450 words maximum, unstructured (no use of sub-headers).

References: no maximum.

Figures/tables: minimum 1 figure or table.

Description: Reviews are comprehensive analyses of specific topics. ACE emphasizes that an acceptable Review Article should not be a ‘book chapter’ generally covering a topic, but should be a focused application of literature to address a relevant clinical issue. The Editors submit them upon invitation. Proposals for reviews may be submitted; however, in this case authors should only send an outline of the proposed paper for initial consideration. Both solicited and unsolicited review articles will undergo peer review prior to acceptance. Systematic review with no meta-analytics in ACE is addressed as Review Article. Review Articles should entail a section describing the contribution of each author to the manuscript. See section “AUTHORS’ CONTRIBUTION’ for details.

(3) MINI REVIEW

Word limit: 4,000 words maximum including abstract but excluding references, tables and figures.

Abstract: 450 words maximum, unstructured (no sub-headers).

References: no maximum.

Figures/tables: maximum 6 figures or tables.

Description: Mini Reviews are shorter reviews of topics that may be controversial or unresolved. They are submitted upon invitation by the Editors. Proposals for reviews may be submitted; however, in this case authors should only send an outline of the proposed paper for initial consideration. Both solicited and unsolicited review articles will undergo peer review prior to acceptance. Mini review should entail a section describing the contribution of each author made to the manuscript. See section “AUTHORS’ CONTRIBUTION “ for details.

(4) BRIEF REPORT

Word limit: 2,500 words including abstract but excluding references, tables and figures.

Abstract: 250 words, unstructured (no sub-headers).

References: 35 maximum.

Figures/tables: 8 maximum.

Description: Manuscripts containing pertinent and

interesting observations concerning cancer epidemiology and reports on new observations or studies that do not warrant publication as a full research article will be considered for the Brief Reports. These submissions will undergo full peer review.

(5) DESCRIPTIVE BRIEF REPORT FROM REGIONAL AND LOCAL REGISTRIES

Descriptive Brief Reports from Regional and Local Registries provides a forum for authors to report patterns of cancer obtained from regional or local cancer registries with limited resources. This section of ACE provides authors with the opportunity to carefully describe their data collection methods and cancer patterns observed.

Word limit: 2,500 words including abstract but excluding references, tables and figures.

Abstract: 250 words, unstructured (no sub-headers).

References: 35 maximum.

Figures/tables: 8 maximum.

(6) PERSPECTIVE

Word limit: 3000 words maximum including abstract but excluding references, tables and figures.

Abstract: 300 words maximum, unstructured (no use of sub-headers)

References: no maximum.

Figures/tables: minimum 1 figure or table

Description: Perspective articles can be more subjective, forward-looking or speculative. A paper presenting controversial positions or papers of the same topic advocating opposite opinions will be published as Perspectives. Most perspective articles will be solicited by editors. However, we also welcome timely, unsolicited perspective articles.

(7) EDITORIAL

Word Limit: 2,500 words maximum excluding references, tables and figures.

Abstract: not required for this manuscript type.

References: 25 maximum.

Figures/Tables: 2 maximum.

Description: Editorial is written by recognized leader(s) in the field. It is generally solicited by the (Deputy) Editor(s)-in-Chief.

(8) COMMENTARY

Word limit: 1,500 words maximum excluding references, tables and figures.

Abstract: not required for this manuscript type.

References: 20 maximum, including the article discussed.

Figures/tables: 2 maximum.

Description: Commentary, upon Editor's invitation, discusses a paper or report or event within the past few months or so, or in the near future. It should set the problems addressed by the paper/report/event in the wider context of the field. Proposals for Commentary may be submitted; however, in this case authors should only send an outline of the proposed paper for initial consideration.

(9) CORRESPONDENCE

Word limit: 1000 words maximum excluding references, tables and figures.

Abstract: not required for this manuscript type.

References: 10 maximum.

Figures/tables: Only 1 table or figure.

Description: Correspondence on content published in the Journal or on other topics of interest to our readers is welcomed. The journal might invite replies from the authors of the original publication, or pass on letters to these authors.

(10) VIEWPOINT

Word limit: 1200 words maximum excluding references, tables and figures.

Abstract: not required for this manuscript type.

References: 10 maximum.

Figures/tables: Only 1 table or figure.

Description: Viewpoints may address virtually any important topic in medicine, public health, research, ethics, health policy, or health law and generally are not linked to a specific article. Viewpoints should be well focused, scholarly, and clearly presented and must have no more than 3 authors.

(11) MEETING REPORT

Word limit: 4,000 words maximum including abstract but excluding references, tables and figures.

Abstract: 350 words maximum, unstructured (no use of sub-headers).

References: no maximum.

Figures/tables: no maximum, but 8 figures should be sufficient.

Description: Brief reports of symposia and conferences in related to cancer epidemiology. Reports must be submitted within 2 months of the meeting date in order to maintain their timeliness. Only those Meeting Reports dealing with topics of interest to the readership and that contain novel information and insights from the meeting are accepted for publication. A Meeting Report should be a thoughtful, critical commentary which shows an appreciation of the

connections among the various presentations and reveals the consensus, if any, which emerged at the meeting. Before submitting a full Meeting Report, authors should only send an outline of the proposed paper for initial consideration.

(12) CLINICAL GUIDELINE

Word limit: 6,000 words maximum including abstract but excluding references, tables and figures.

Abstract: 450 words maximum, unstructured (no sub-headers).

References: no maximum.

Figures/tables: minimum 1 figure or table.

Description: Guidelines need to be the product of a large group of individuals who are recognized authorities in their field. Guidelines will be written by a working party to include a steering committee (usually at least 4 members) and other authors representing a wide range of those with special relevant expertise as well as those whose everyday practice will be influenced by the guidelines.

3. STRUCTURE OF THE MANUSCRIPT

The length of manuscripts must adhere to the specifications under the section "MANUSCRIPT CATEGORIES". Manuscripts should be presented in the following order: (i) title page, (ii) abstract and key words, (iii) text, (iv) acknowledgments, (v) footnote, (vi) references, (vii) supplementary material, (viii) figure legends, (ix) tables (each table complete with title and footnotes) and (x) figures (it is recommended that figures, tables and videos are provided in separate files).

TITLE PAGE

The title page should include

- The title of the paper. Concise titles are easier to read than long, convoluted ones. Titles that are too short may, however, lack important information, such as study design (which is particularly important in identifying randomized controlled trials). Authors should include all information in the title that will make electronic retrieval of the article both sensitive and specific. (abbreviations is not allowed)
- The full names of the authors and the addresses of the institutions at which the work was carried out (in English).
- The full postal and email address, plus facsimile and telephone numbers, of the corresponding author.
- A short running title (less than 60 characters) should also be provided.
- Author's Contribution. In keeping with the latest guidelines of the International Committee of Medical

Journal Editors, for the original article, review article and systematic review/meta-analysis, the information of author contribution is needed (See section “Author’s Contribution” for details).

ABSTRACT AND KEYWORDS

The abstracts must adhere to the specifications under the section Manuscript Categories. The abstract of an original article, review article, systematic review and meta-analysis, should be structured into four paragraphs with sub-headers of background, methods, results and conclusions. The abstracts for all the other manuscript types should be unstructured. The abstract should not contain any abbreviations or acronyms, as well as citations of reference, figures or tables. And general statements (e.g. “the significance of the results is discussed”) should be avoided. Following the Abstract, 3-5 keywords should be given.

TEXT

The text part should be arranged into short/sharp paragraphs, which are best suited for reading on-screen. Authors must use the following sub-headers to divide the sections of their Original Article manuscript: Introduction, Methods, Results, Discussion, Acknowledgment, Footnote, References, and when relevant, Supplementary Material. Plus, authors should follow the same structures in systematic review and meta-analysis. However, review, perspective, viewpoint, commentary and others do not have those clear sections, they can be written in several sections with their own headers according to the topic (see detailed requirements in the previous section “MANUSCRIPT CATEGORIES”).

If an article describes any procedure, technology or apparatus that is new, has not been used in the indication described, or is being used for a purpose for which it was not originally intended, it is the responsibility of the authors to ensure that all ethical committee, institutional review board, and/or governing body approval has been properly obtained. Such approval must be explicitly stated in the main text.

The text should be keyed double-spaced throughout. A clearly readable font should be used (e.g. Arial, Calibri, Times New Roman, Verdana). Font size should be 10 or 12. Pages should be numbered. Language should be English. Spelling can be British or American, but consistent throughout. Any abbreviations should be defined on first usage in the text. Terms that are mentioned less than 3 or 4 times in the text should not be abbreviated.

AUTHORS’ CONTRIBUTION

This section is only required for original article, review

article and systematic review/meta-analysis. It describes the contribution of each author made to be manuscript. Authorship credit should be based on:

- 1) substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data;
- 2) drafting the article or revising it critically for important intellectual content; and
- 3) final approval of the version to be published.
- 4) Agreement to be accountable for all aspects of the work in ensuring that questions that related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Author should meet conditions 1, 2, 3, and 4, and all who meet the four criteria should be identified as authors. Those who do not meet all four criteria should be acknowledged (see section “ACKNOWLEDGMENTS” for details). Please note that acquisition of funding, collection of data, language editing or general supervision of the research group alone does not constitute authorship.

The “AUTHORS’ CONTRIBUTION” section should be completed as follow:

- (1) Conception and design:
- (2) Administrative support:
- (3) Provision of study material or patients:
- (4) Collection and assembly of data:
- (5) Data analysis and interpretation:
- (6) Manuscript writing: All authors.
- (7) Final approval of manuscript: All authors.

Note: 1. Manuscript writing part and Final approval of manuscript part are required to be included while other parts are based on actual applicability; 2. Contribution is not required when there is only one author.

ACKNOWLEDGMENTS

a. All contributors who do not meet the criteria for authorship should be listed in an acknowledgments section. Examples of those who might be acknowledged include a person who provided purely technical help, writing or language editing assistance, or a department chairman who provided only general support. Financial and material support should also be acknowledged.

b. Funding: Details of all funding sources for the work in question should be included in the Acknowledgment section.

The following rules should be followed:

The sentence should begin: “This work supported by...”; The full official funding agency name should be given, i.e. “National Institutes of Health”, not “NIH” (full RIN approved list of UK funding agencies).

Grant numbers should be given in brackets as follows: “[grant number XXX]”. Multiple grant numbers should be separated by a comma as follows: “[grant numbers XXX, YYY]”;

Agencies should be separated by a semi-colon (plus “and” before the last funding agency).

Where individuals need to be specified for certain sources of funding the following text should be added after the relevant agency or grant number “to [author initials]”;

An example is given here: “This work was supported by the National Institutes of Health [AA123456 to C.S.,BB765432 to M.H.]; and the Alcohol & Education Research Council [hfygr667789]”.

c. When there is nobody or funding to be acknowledged, please describe as “None”.

FOOTNOTE

a. Conflicts of Interest: See section “Conflicts of Interest” for details.

b. Financial Disclosure: Some variables, such as “measures of income inequality and degree of financial openness, are not included in our study because of the limited availability of good quality data across countries over the sample period”. When there is no financial disclosure, this section should be removed.

REFERENCES

A list of references to the literature should be arranged sequentially following appearance in the text. Personal communications, and unpublished data should not be included in the list of references, but can be mentioned in the text.

The Vancouver system of referencing should be used (examples are given below). In the text, references should be cited using Arabic numerals in round brackets in which they appear consecutively [e.g. “cancer-related mortality (19)”]; “heart failure (29,30)”. If cited in tables or figure legends, number according to the first identification of the table or figure in the text. In the reference list, cite the names of all authors when there are three or fewer; when three or more, list the first three followed by et al.

Do not use *ibid.* or *op cit.* Reference to unpublished data and personal communications should not appear in the list but should be cited in the text only (e.g. Smith A, 2000, unpublished data). All citations mentioned in the text, tables or figures must be listed in the reference list. Names of journals should be abbreviated in the style used in Pubmed. Authors are responsible for the accuracy of the references.

The format of reference sees as follow.

• Journal article

e.g., Gibas Z, Prout DF Jr, Pontes JR. Chromosome changes in germ cell tumours of the testis. *Cancer Genet Cytogenet* 1986; 19: 254-52.

• Online article not yet published in an issue

An online article that has not yet been published in an issue (therefore has no volume, issue or page numbers) can be cited by its Digital Object Identifier (DOI). The DOI will remain valid and allow an article to be tracked even after its allocation to an issue.

e.g., Furuya R, Takahashi R, Furuya S, et al. Is urethritis accompanied by seminal vesiculitis? *Int J Urol*. DOI:10.1111/j.1442-2042.2009.02314.x

• Book

e.g., Ernstoff M. *Urologic Cancer*. Blackwell Science, Boston, 1997.

• Chapter in a Book

e.g., Gilchrist RK. Further commentary: Continent stroma. In: King LR, Stone AR, Webster GD (eds). *Bladder Reconstruction and Continent Urinary Diversion*. Year Book Medical, Chicago, 1987; 204-5.

• Online publications

e.g., Hraska V, Photiadis J, Poruban R, Asfour B. Ross-Konno operation in children . *Multimed Man Cardiothorac Surg* doi: 10.1510/mmcts.2008.003160.

or

e.g., Thurber JS, Deb SJ, Collazo LR. Ascending-to-descending aortic bypass for coarctation of the aorta. *CTSNet* [published 12 May 2008, accessed 30 November 2011]. Available from: <http://www.ctsnet.org/sections/clinicalresources/adultcardiac/>

TABLES

Tables should be self-contained and complement, but not duplicate information contained in the text. All tables should be numbered consecutively in the order of reference in the text. Each Table should be on a separate page; tables must be typed and editable in tabular form that is convenient for copyediting and typesetting; and they should not be inserted as images.

Each column must carry an appropriate heading and, if measurements are given, the units should be given in the column heading. Column headings should be brief, with units of measurement in parentheses; all abbreviations must be defined in footnotes. Footnote symbols: †, ‡, §, ¶, should

be used (in that order) and *, **, *** should be reserved for P-values. Statistical measures such as SD or SEM should be identified in the headings. If tables have been reproduced from another source, a letter/permission from the copyright holder (usually the Publisher), stating authorization to reproduce the material, must be submitted as supplemental materials during paper submission.

FIGURES

All illustrations (line drawings and photographs) are classified as figures. Figures should be numbered consecutively in the order of reference in the text. Figures should be provided separately. Magnifications should be indicated using a scale bar on the illustration. If figures have been reproduced from another source, a letter/permission from the copyright holder (usually the Publisher), stating authorization to reproduce the material, must be submitted as supplemental materials.

Size: Figures should be sized to fit within the page column (82 mm), intermediate (118 mm) or the full text width (173mm).

Specifications:: Figures must be supplied as high resolution saved as .eps, .tif or .jpg; 300 dpi (dots per inch), figures containing text 400 dpi, Line figures 1,000 dpi. Pixel screen width: 1280, grayscale for black and white, RGB for color.

Line figures: Must be sharp, black and white graphs or diagrams, drawn professionally or with a computer graphics package.

Text sizing in figures: Lettering must be included and should be sized to be no larger than the journal text or 8 point (Should be readable after reduction – avoid large type or thick lines). Line width between 0.5 and 1 point.

Figure legends: Type figure legends on a separate page. Legends should be concise but comprehensive – the figure and its legend must be understandable without reference to the text. Include definitions of any symbols used and define/explain all abbreviations and units of measurement.

VIDEO

ACE will accept digital files in mp4, avi., mov., and wmv (keep the bit rate as high as possible), MPEG(MPEG video file), flash video (flv.), DVD video format, etc. Contributors are asked to be succinct, and the Editor-in-chief reserves the rights to require shorter video duration if necessary. Video files can be submitted with a manuscript online: <http://ace.amegroups.com/pages/view/submit-multimedia-files>.

Duration: Video files should be limited to 20 minutes.

Quality: Please set the video aspect ratio as 4:3 or 16:9(widescreen). The original video should be of high quality. The resolution is no less than 1280*720, the frame

rate no less than 24 frames per second and the bit rate no lower than 5Mbps.

Text in video: All the text notes, explanations or descriptions, etc. in the video must be in English. And the logo or watermark of hospital should not be stick on the screen. Plus, the information of patients should be erased from the video.

Video legends: Legends for the video files should be provided. The video files should be number consecutively in the order of reference in the text.

APPENDIX

The supplementary appendix should be paginated, with a table of contents, followed by the list of investigators (if there is one), text (such as methods), figures, tables, and then references. The supplementary appendix should not be included in the article's reference list.

The appendix must be submitted in a Word file. The appendix will not be edited for style. It will be presented online as additional information provided by the authors.

The published article will contain a statement that supplementary material exists online and will provide the reader with a URL and link. To reference the supplementary appendix in the text of the article, refer to it as in the following example:

“Many more regressions were run than can be included in the article. The interested reader can find them in a supplementary appendix online”.

EQUATIONS

Equations should be numbered sequentially with Arabic numerals; these should be ranged right in parentheses. All variables should appear in italics. Use the simplest possible form for all mathematical symbols.

4. STYLE OF THE MANUSCRIPT

Manuscripts must follow the style of the Vancouver agreement detailed in the International Committee of Medical Journal Editors' revised 'Uniform Requirements for Manuscripts Submitted to Biomedical Journals: Writing and Editing for Biomedical Publication', as presented at: <http://www.ICMJE.org/>.

Author name: Each author's given name should be followed by family name.

Capitalize each letter of the Family name. A hyphen could be used in Family name according to the rule in Author region

Capitalize the first letter of those words/syllables that they hope to be abbreviated in their given name, otherwise, DO NOT capitalize the first letter and use a hyphen to connect

it with its anterior word.

Spelling: The Journal uses US spelling and authors should therefore follow the latest edition of the Merriam–Webster’s Collegiate Dictionary.

Units: All measurements must be given in SI or SI-derived units. For more information about SI units, please go to the Bureau International des Poids et Mesures (BIPM) website at: <http://www.bipm.fr>.

Abbreviations: Must be used sparingly – only where they ease the reader’s task by reducing repetition of long, technical terms. Initially use the word in full, followed by the abbreviation in parentheses. Thereafter use the abbreviation only.

Trade names: Drugs should be referred to by their generic names. If proprietary drugs have been used in the study, refer to these by their generic name, mentioning the proprietary name, and the name and location of the manufacturer, in parentheses.

5. REVIEW PROCESS

Manuscripts are assigned sequentially to Science Editors. An Science Editor solicits reviewers (typically, two external reviews are sought). The reviewers’ evaluations and Science Editor’s comments are compiled by the Editor-in-Chief for disposition and transmittal to the authors. A decision is made usually within four weeks of the receipt of the manuscript.

The Editor-in-Chief will advise authors whether a manuscript is accepted, should be revised or is rejected. Minor revisions are expected to be returned within two weeks of decision; major revisions within three weeks. Manuscripts not revised within these time periods are subject to withdrawal from consideration for publication unless the authors can provide extenuating circumstances.

A number of manuscripts will have to be rejected on the grounds of priority and available space. A manuscript maybe returned to the authors without outside review if the Editor-in-Chief and Science Editor find it inappropriate for publication in the Journal. Similarly, the Editors may expedite the review process for manuscripts felt to be of high priority in order to reach a rapid decision. Such ‘fast track decisions’ will normally occur within one week of receipt of the manuscript.

Authors may provide the Editor-in-Chief with the names, addresses and email addresses of up to three suitably qualified individuals of international standing who would be competent to referee the work, although the Editor-in-Chief will not be bound by any such nomination. Likewise, authors may advise of any individual who for any reason, such as potential conflict of interest, might be inappropriate

to act as a referee, again without binding the Editor-in-Chief.

The Editor-in-Chief’s decision is final. If, however, authors dispute a decision and can document good reasons why a manuscript should be reconsidered, a rebuttal process exists. In the first place, authors should write to the Editor-in-Chief.

All journals Manuscripts should be written in a clear, concise, direct style so that they are intelligible to the professional reader who is not a specialist in the particular field. When contributions are judged as acceptable for publication, the Editor and the Publisher reserve the right to modify manuscripts to eliminate ambiguity and repetition and improve communication between authors and readers. If extensive alterations are required, the manuscript will be returned to the author for revision.

6. ETHICAL CONSIDERATIONS

Authors must state that the protocol for the research project has been approved by a suitably constituted Ethics Committee of the institution within which the work was undertaken and that it conforms to the provisions of in accordance with the Helsinki Declaration as revised in 2013, available at: <http://www.wma.net/en/30publications/10policies/b3/%20index.html>. The journal retains the right to reject any manuscript on the basis of unethical conduct of either human or animal studies. All investigations on human subjects must include a statement that the subject gave informed consent. Patient anonymity should be preserved. Photographs need to be cropped sufficiently to prevent human subjects being recognized (or an eye bar should be used).

❖ **For studies in the following categories:**

Randomized controlled trials or other intervention research: This category includes any study that carries out medical intervention(s) on patients or healthy individuals.

Case-control study: A case-control study is designed to retrospectively analyze the exposure to the risk factor of interest in subjects with known outcomes (with or without disease; dead or alive; or, with or without other predetermined endpoints).

Prospective cohort study: In a prospective cohort study, patients with known exposure to a risk factor are followed and then the outcomes (with or without disease; or, dead or alive) were identified.

Cross-sectional studies: Cross-sectional studies are performed to investigate the occurrence of a specific disease or the status quo of a clinical condition.

Basic or translational medical research using human

specimens:

- Authors must state whether their studies had been approved by an institutional review board (IRB) (if yes, please provide the number of approval document). For a multi-center study, IRB approval must be obtained from each center.
- The authors must state whether all the subjects had signed the informed consent forms. For subjects under 18 years of age or those with limited capacity for civil conduct, the authors must state whether their caregivers had signed the informed consent forms.
- Also, the authors should state whether the study outcomes will affect the future management of the patients.

❖ For other categories:

Retrospective and ambispective cohort studies: In these studies, the patients' exposure to risk factor(s) were retrospectively identified, followed by the retrospective follow-up of the patients to determine the relationship between the future or current endpoints (with or without disease; or, dead or alive) and the exposure.

- For studies in this category, authors must state whether their study had been approved by an institutional review board (IRB) (if yes, please provide the number of approval document). For a multi-center study, IRB approval must be obtained from each center.
- Also, the authors should state whether the study outcomes will affect the future management of the patients.
- The authors must state whether all the subjects had signed the informed consent forms before enrollment. For subjects under 18 years of age or those with limited capacity for civil conduct, the authors must state whether their caregivers had signed the informed consent forms. For deceased patients or those who had lost capacity for civil conduct, the informed consent forms could be signed by their family members or caregivers. For studies on patient data retrieved from hospital medical record system or social insurance systems, an informed consent form is not required; however, the authors still need to declare whether the patient's personal data have been secured.

Systematic review and meta-analysis, review, opinion, hypothesis, and editorial

- No statement on medical ethics is required.

Case report and visualized surgery:

- No statement on medical ethics is required. However,

in cases of involving new and controversial treatments, approval from IRC might be required.

- Informed consent must be obtained from the subjects or their caregivers.

Diagnostic accuracy tests: These studies are performed to evaluate the efficiency of a specific index test in disease diagnosis.

- For studies in this category, authors must state whether their study had been approved by an institutional review board (IRB) (if yes, please provide the number of approval document). For a multi-center study, IRB approval must be obtained from each center.
- Also, the authors should state whether the study outcomes will affect the future management of the patients.
- If the study has a prospective design: the authors must state whether all the subjects had signed the informed consent forms before enrollment. For subjects under 18 years of age or those with limited capacity for civil conduct, the authors must state whether their caregivers had signed the informed consent forms. However, for retrospective studies based on a hospital medical record system, no informed consent is required.

Nested case-control study: In a nested case-control study, the patients were followed up after the biological samples are obtained from the subjects, and then a subset of patients are chosen for the analysis.

If the study has a prospective design:

- Authors must state whether their study had been approved by an institutional review board (IRB) (if yes, please provide the number of approval document). For a multi-center study, IRB approval must be obtained from each center.
- Also, the authors should state whether the study outcomes will affect the future management of the patients.
- The authors must state whether all the subjects have signed the informed consent forms before they enter the study, no matter whether they enter the final analysis. For subjects under 18 years of age or those with limited capacity for civil conduct, the authors must state whether their caregivers had signed the informed consent forms.

If the study is based on a previously available specimen bank, the authors must:

- State whether the specimen bank had been approved by the IRB upon its establishment;
- State whether all the subjects had signed the informed

consent forms during the establishment of the bank(attached with the numbers of approval documents).

Post hoc analysis: In a post hoc analysis, the authors re-examines the currently available data from different perspectives.

- The authors need to state whether the previous studies had been approved by the local medical ethics committee(s)
- Also, it is important to state whether all the subjects had signed the informed consent forms in the previous studies.

For more information on statement of ethics, please feel free to consult our editorial staff.

7. STATEMENT OF ETHICS APPROVAL

Statement of Ethics Approval: We require every research article submitted to include a statement that the study obtained ethics approval (or a statement that it was not required and why), including the name of the ethics committee(s) or institutional review board(s), the number/ID of the approval(s), and a statement that participants gave informed consent before taking part. The statement should be described in the method section.

* When concerning experiments on human subjects, authors should indicate whether the procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national). Furthermore, authors also need to confirm that the patient has given their consent for the publication. The editorial office may request copies of the informed consent documentation at any time. We recommend the following wording used for the consent section as: "Written informed consent was obtained from the patient for publication of this article and any accompanying images. A copy of the written consent is available for review by the Editors-in-Chief of this journal."

* When concerning experiments on animals, authors should be asked to indicate whether the institutional and national guide for the care and use of laboratory animals was followed.

8. INFORMED CONSENT

Identifying information, including names, initials, or hospital numbers, should not be published in written descriptions, photographs, or pedigrees unless the information is essential for scientific purposes and the patient (or parent or guardian) gives written informed consent for publication. Informed consent is required for Case report, original/research articles and visualized

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