Quantitative Imaging in Medicine and Surgery
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INSTRUCTION FOR AUTHORS

Thank you for your interest in *Quantitative Imaging in Medicine and Surgery* (*QIMS*). 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. Upon acceptance, we aim to publish the papers within 3 months, and we will make sure that all accepted papers will be published within 6 months. We are looking forward to your submission.

TABLE OF CONTENT

1. ABOUT THE JOURNAL

The *Quantitative Imaging in Medicine and Surgery* (Print ISSN 2223-4292; Online ISSN 2223-4306; *QIMS*) publishes peer-reviewed original reports and reviews in medical imaging, including X-ray, ultrasound, computed tomography, magnetic resonance imaging and spectroscopy, nuclear medicine and related modalities, and their application in medicine and surgery. While focus is on clinical investigations, papers on medical physics, image processing, or biological studies which have apparent clinical relevance are also published. This journal encourages authors to look at the medical images from a quantitative angle. Descriptive papers are of particular interests are also published. This journal is also interested in publishing papers on biomedical research policy, medical education and training, public health, and philological and historical thoughts related to biology and medicine.

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2. REVIEW PROCESS

Manuscripts are assigned sequentially to Associate Editors. An Associate Editor solicits reviewers (typically, two external reviews are sought). The reviewers' evaluations and Associate Editor's comments are compiled by the Editor-in-Chief for disposition and transmittal to the authors. A decision is made usually within six 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 four weeks of decision; major revisions within three months. 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 may be returned to the authors without outside review if the Editor-in-Chief and Associate 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 so that they are intelligible to the professional reader who is not a specialist in the particular field. They should be written in a clear, concise, direct style. Where 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 author and reader. If extensive alterations are required, the manuscript will be returned to the author for revision.

3. MANUSCRIPT CATEGORIES

(1) ORIGINAL ARTICLES

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

Abstract: 450 words maximum, structured with the following subheadings: Background, Methods, Results and Conclusions.

References: no limit.

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

Description: Full-length reports of current research in either basic or clinical science. Meta-analysis will be categorized into this type.

(2) INVITED REVIEWS

Word limit: 6,000 words maximum including abstract but excluding references, tables and figures.
Abstract: 450 words maximum.
References: no maximum.
Figures/tables: minimum 1 image or figure.
Description: Reviews are comprehensive analyses of specific topics. 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.

(3) MINI REVIEWS (research highlight)

Word limit: 4,000 words maximum including abstract but excluding references, tables and figures.
Abstract: 450 words maximum.
References: no maximum.
Figures/tables: maximum 6 images or figures.
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.

(4) IMAGING IN MEDICINE AND SURGERY

Word limit: 1000 words excluding references, tables and figures.
Abstract: A brief abstract, usually of 3-4 sentences, is required.
References: Up to 10.
Figures/Videos: 2 still images maximum for the print and PDF article, supplemented by 2 video maximum online. It is important to demonstrate how quantitative measure supported the diagnosis or management of the disease.
Description: Videos which are unique or highly illustrative of specific occurrences. They will be reviewed by the Editors prior to acceptance, but they do not have to go out for external peer review. They must be accompanied by a brief one paragraph description of relevant information. Please note our journal can publish limited such images per year.

(5) CLINICAL GUIDELINES

Word limit: 6,000 words maximum including abstract but excluding references, tables and figures.
Abstract: 450 words maximum.
References: no maximum.
Figures/tables: minimum 1 image or figure.
Description: Guidelines need to be the product of a large group of individuals who are recognised 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.

(6) LETTERS TO THE EDITOR

Word limit: 1000 words maximum excluding references, tables and figures.
Abstract: not required for this manuscript type.
References: 10 maximum.
Figures/ tables: 1 maximum.
Description: Letters usually offer perspective to content published in QIMS. In this case, a Letter must reference the original source, and a Response to a Letter must reference the Letter in the first few paragraphs. Letters can use an arbitrary title, but a Response must cite the title of the Letter: e.g. Response to [title of Letter]. This ensures that readers can track the line of discussion. Letters of any matter of interest to readers of the QIMS are also published.

(7) EDITORIALS

Word Limit: 2,500 words maximum excluding references, tables and figures.
Abstract: Not required.
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) COMMENTARIES

Word limit: 1,500 words maximum excluding references.
Title: 20 words maximum.
Abstract: no abstract required for this manuscript type.
References: 20 maximum, including the article discussed.
Figures/tables: 2 maximum.
Description: Commentaries, upon Editor's invitation, discuss a paper published in a specific issue and should set the problems addressed by the paper in the wider context of the field. Proposals for Commentaries may be submitted; however, in this case authors should only send an outline of the proposed paper for initial consideration

(9) PERSPECTIVES

Word limit: 3000 words maximum including abstract but excluding references, tables and figures.
Abstract: Unstructured. 300 words maximum.
References: no maximum.
Description: Perspectives can be more personal, forward-looking or speculative, compared with reviews of a scientific topic. A paper presenting controversial positions or papers of the same topic advocate opposite sides will be published as Perspectives. Most of Perspectives will be solicited by the editors; however, we also welcome timely, unsolicited Perspectives. Proposals for perspectives may be submitted; however, in this case authors should send an outline of the proposed article prior to submission.

(10) VIEWPOINT

Word limit: 1200 words maximum excluding references, tables and figures.
Abstract: Not required.
References: 10 maximum.
Figures/tables: Only one 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) CASE REPORTS

Word limit: 2,500 words maximum excluding references, tables and figures.
Abstract: A brief abstract, usually of 3-4 sentences, is required.
References: 20 maximum.
Figures/ tables: 8 maximum.
Description: New observations of diseases, clinical findings or novel/unique treatment outcomes relevant to practitioners in medicine and surgery. The text should be arranged as follows: Introduction, Case Report, Discussion. The authors should provide a statement at the end of the main text to confirm that the patient has given their consent for the Case reports to be published. The editorial office may request copies of the informed consent documentation at any time. We recommend the following wording is used for the consent section: "Written informed consent was obtained from the patient for publication of this Case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal."

If the patient has died, then consent for publication must be sought from the next of kin of the patient. If the patient is a minor, or unable to provide consent, then consent must be sought from the parents or legal guardians of the patient. In these cases, the statement in the 'Consent' section of the manuscript should be amended accordingly. Only cases of exceptional interest and novelty are considered. It is important to demonstrate how quantitative measure supported the diagnosis or management of the disease. This journal welcomes the initial cases of implementing or validation of novel imaging technique. Note the rarity of a disease is not the focus of this journal. For manuscripts that do not qualify, Editors may ask authors to shorten manuscripts and rewrite as Letters to the Editor. Please

note our journal can publish limited such cases per year.

(12) MEETING REPORTS

Word limit: 4,000 words maximum including abstract but excluding references, tables and figures.
Abstract: 350 words maximum, with sub-headers.
References: no limit.
Figures/ tables: no limit, but 8 figures should be sufficient.
Description: Brief reports of symposia and conferences in quantitative research. 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.

(13) TECHNICAL NOTES

Word limit: 2,500 words including abstract but excluding references, tables and figures.
Abstract: 250 words, unstructured (no use of sub-headers).
References: Up to 35.
Figures/tables: Up to 10 in total.
Description: Technical notes articles should present a new experimental or improved method, test or procedure. The method described may either be completely new, or may offer a better version of an existing method. The article must describe a demonstrable advance on what is currently available. The method needs to have been well tested and ideally, but not necessarily, used in a way that proves its value.

(14) BRIEF REPORTS

Word limit: 2,500 words including abstract but excluding references, tables and figures.
Abstract: 250 words, unstructured (no use of sub-headers).
References: Up to 35.
Figures/tables: Up to 8 in total.
Description: Manuscripts containing pertinent and interesting observations concerning quantitative imaging research in medicine and surgery 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.

4. DISCLOSURE

At the time of submission, the submitting author must include a disclosure statement in the body of the manuscript. The statement whether the authors have published or submitted the manuscript elsewhere. The statement will also describe all of the authors' relationships with companies that may have a financial interest in the information contained in the manuscript. This information should be provided under the heading titled 'Disclosure,' which should appear after the 'Acknowledgement' section and before the 'References' section. The absence of any interest to disclose must also be stated. In addition, any financial interests must be detailed in the Financial Disclosure form, which will be provided to the corresponding author upon acceptance for distribution to each author.

5. 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 pre-determined 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.

6. 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 surgery**. The statement should be included in the footnote.

It may be possible to publish without explicit consent if the report is important to public health (or is in some other way important); consent would be unusually burdensome to obtain;

and a reasonable individual would be unlikely to object to publication (all three conditions must be met).

7. POLICIES ON CONFLICT OF INTEREST

Our journal complies with the International Committee of Medical Journal Editors' uniform requirements on Conflict of Interest statement.

Conflict of Interest exists when an author (or the author's institution), reviewer, or editor has financial or personal relationships with other persons or organizations that inappropriately influence (bias) his or her actions. The existence of such relationships does not necessarily represent true conflict of interest. The potential for conflict of interest can exist whether or not an individual believes that the relationship affects their judgment. Financial relationships (such as employment, consultancies, stock ownership, honoraria, paid expert testimony, patents) are the most easily identifiable conflicts of interest and the most likely to undermine the credibility of the journal, the authors, and of science itself (http://www.icmje.org/index.html).

(1) PARTICIPANTS

All participants in the peer-review and publication process—not only authors but also peer reviewers, editors, and editorial board members of journals—must consider their conflicts of interest when fulfilling their roles in the process of article review and publication and must disclose all relationships that could be viewed as potential conflicts of interest.

a. Authors

When authors submit a manuscript of any type or format they are responsible for disclosing all financial and personal relationships that might bias or be seen to bias their work.

The ICMJE recommends that authorship be based on the following 4 criteria:

- Substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work; AND
- Drafting the work or revising it critically for important intellectual content; AND
- Final approval of the version to be published; AND
- Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

b. Peer Reviewers

Reviewers should be asked at the time they are asked to critique a manuscript if they have conflicts of interest that could complicate their review. Reviewers must disclose to editors any conflicts of interest that could bias their opinions of the manuscript, and should recuse themselves from reviewing specific manuscripts if the potential for bias exists. Reviewers must not use knowledge of the work they're reviewing before its publication to further their own interests.

c. Editors and Journal Staff

Editors who make final decisions about manuscripts should recuse themselves from editorial decisions if they have conflicts of interest or relationships that pose potential conflicts related to articles under consideration. Other editorial staff members who participate in editorial decisions must provide editors with a current description of their financial interests or other conflicts (as they might relate to editorial judgments) and recuse themselves from any decisions in which a conflict of interest exists. Editorial staff must not use information gained through working with manuscripts for private gain. Editors should publish regular disclosure statements about potential conflicts of interests related to the commitments of journal staff. Guest editors should follow these same procedures.

(2). REPORTING CONFLICTS OF INTEREST

Articles should be published with statements or supporting documents, declaring:

- Authors' of interest; and
- Sources of support for the work, including sponsor names along with explanations of the role of those sources if any in study design; collection, analysis, and interpretation of data; writing of the report; the decision to submit the report for publication; or a statement declaring that the supporting source had no such involvement; and
- Whether the authors had access to the study data, with an explanation of the nature and extent of access, including whether access is on-going.

To support the above statements, editors may request that authors of a study sponsored by a funder with a proprietary or financial interest in the outcome sign a statement, such as "I had full access to all of the data in this study and I take complete responsibility for the integrity of the data and the accuracy of the data analysis."

8. CLINICAL TRIALS REGISTRY

We require, as a condition of consideration for publication, registration in a public trials registry. Trials must register at or before the onset of patient enrollment. This policy applies to any clinical trial starting enrollment after January 1, 2006. For trials that began enrollment before this date, we require registration by April 1, 2006, before considering the trial for publication. We define a clinical trial as any research project that prospectively assigns human subjects to intervention or comparison groups to study the cause-and-effect relationship between a medical intervention and a health outcome. Studies designed for other purposes, such as to study pharmacokinetics or major toxicity (e.g., phase 1 trials) are exempt.

We do not advocate one particular registry, but registration

must be with a registry that meets the following minimum criteria: (1) accessible to the public at no charge; (2) searchable by standard, electronic (Internet-based) methods; (3) open to all prospective registrants free of charge or at minimal cost; (4) validates registered information; (5) identifies trials with a unique number; and (6) includes information on the investigator(s), research question or hypothesis, methodology, intervention and comparisons, eligibility criteria, primary and secondary outcomes measured, date of registration, anticipated or actual start date, anticipated or actual date of last follow-up, target number of subjects, status (anticipated, ongoing or closed) and funding source(s).

Registries that currently meet these criteria include: (1) the registry sponsored by the United States National Library of Medicine (www.clinicaltrials.gov); (2) the International Standard Randomized Controlled Trial Number Registry (http://www.controlled-trials.com); (3) the Australian Clinical Trials Registry (http://www.actr.org.au); (4) the Chinese Clinical Trials Register (http://www.chictr.org); and (5) the Clinical Trials Registry - India (http://www.ctri.in).

9. RANDOMIZED CONTOLLED TRIALS

Reporting of randomized controlled trials should follow the guide-lines of The CONSORT Statement:

http://www.consort-statement.org

10. COPYRIGHT

Papers accepted for publication in the journal become copyright of QIMS and authors will be asked to sign a transfer of copyright form. In signing the transfer of copyright, it is assumed that authors have obtained permission to use any copyrighted or previously published material. All authors must read and agree to the conditions outlined in the Copyright Assignment Form, and must sign the Form or agree that the corresponding author can sign on their behalf. Acceptance of a manuscript is contingent upon receipt of a signed Copyright Assignment Form.

11. 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 captilize 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.

12. 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) disclosure, (vi) references, (vii) supplementary material, (viii) figure legends, (ix) tables (each table complete with title and footnotes) and (x) figures. Footnotes to the text are not allowed and any such material should be incorporated into the text as parenthetical matter.

TITLE PAGE

The title page should contain (i) 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. (ii) the full names of the authors and (iii) the addresses of the institutions at which the work was carried out together with (iv) the full postal and email address, plus facsimile and telephone numbers, of the author to whom correspondence about the manuscript should be sent. The present address of any author, if different from that where the work was carried out, should be supplied in a footnote. The title should be short, informative and contain the major key words so that readers and in particular online users will discover the article easily in online search. Do not use abbreviations in the title. A short running title (less than 60 characters) should also be provided.

ABSTRACT AND KEYWORDS

The length of abstracts must adhere to the word count specifications under the section Manuscript Categories. Do not

use reference, table or figure in the abstract. The abstract of an original article should be structured into four paragraphs with headings of Background, Methods, Results and Conclusions. The abstracts for all other manuscript types should be non-structured. The use of abbreviations and acronyms should be limited and general statements (e.g. ''the significance of the results is discussed'') should be avoided.

Three to five key words should be supplied below the abstract, in alphabetical order, and should be taken from those recommended by the US National Library of Medicine's Medical Subject Headings (MeSH) browser list at: http://www.nlm.nih.gov/mesh/meshhome.html.

TEXT

Authors must use the following subheadings to divide the sections of their Original Article manuscript: Introduction, Materials and Methods, Results, Discussion, Acknowledgment, Disclosure, References, and when relevant, Supplementary Material.

ACKNOWLEDGMENTS

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 chairperson who provided only general support. Financial and material support should also be acknowledged.

When there is no one to be acknowledged, authors should also indicate 'Acknowledgements' section as 'None'.

AUTHOR CONTRIBUTION

This section is only required for original article, systematic review and meta-analysis article. It describes the contribution each author made to the 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. Authors should meet conditions 1, 2, and 3. Please note that acquisition of funding, collection of data, language editing or general supervision of the research group alone does not constitute authorship.

Example:
M.M.M. designed the overall study with contributions from S.F. S.F. designed and carried out experiments, collected and analyzed data, and cowrote the paper. Y.S. designed and carried out experiments and collected and analyzed data with S.F. and M.M.M. S.B. carried out experiments, adapted the rapid TALEN assembly protocol, and analyzed data with Y.S. M.S.W. and M.M.M. designed the vector for the repair experiment. M.S.W. constructed the repair vector. S.F. and Y.S. carried out the repair experiment. S.F., Y.S., M.S.W., and M.M.M. discussed and edited the paper. M.M.M. supervised this study, designed and performed experiments, analyzed data, and wrote the paper.

(Cited from: Fanucchi S, Shibayama Y, Burd S, *et al*. Chromosomal contact permits transcription between coregulated genes. Cell 2013;155:606-20.)

DISCLOSURE

At the time of submission, each author must disclose and describe any involvement, financial or otherwise, that might potentially pose conflict of interest. Disclosure must be included in the text of the manuscript.

REFERENCES

The Vancouver system of referencing should be used (examples are given below). In the text, references should be identified using numbers in round brackets in which they appear consecutively [e.g., "cancer-related mortality (19)"; "denocarcinoma (29,30)"; "malignancies (14-18)"]. 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**. 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.

• Journal article
1. 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.

1. 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
2. Ernstoff M. Urologic Cancer. Blackwell Science, Boston, 1997.

• Chapter in a Book
3. 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.

TABLES

Tables should be self-contained and complement, but not duplicate, information contained in the text. Number tables consecutively in the text in Arabic numerals. Type tables on a separate page with the legend above. Legends should be concise but comprehensive – the table, legend and footnotes must be understandable without reference to the text. Vertical lines should not be used to separate columns. 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 from the copyright holder (usually the Publisher), stating authorization to reproduce the material, must be attached to the covering letter.

FIGURES

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