The *JDR Clinical & Translational Research* (JDR CTR) adheres to the CSE (8th Edition) editorial style. All submitted manuscripts should be formatted in this style.

The *JDR Clinical & Translational Research* (JDR CTR) is a peer-reviewed scientific journal dedicated to the dissemination of new knowledge and information on all science relevant to dentistry and to the oral cavity and associated structures in health and disease. The *JDR Clinical & Translational Research*’s primary readership consists of researchers, clinician scientists, clinicians, educators, policy-makers and the public. The *JDR Clinical & Translational Research* is published quarterly and offers OnlineFirst, by which forthcoming articles are published online before they are scheduled to appear in print.

*JDR Clinical & Translational* operates a conventional single-blind reviewing policy in which the reviewer’s name is always concealed from the submitting Author. All manuscripts are reviewed initially by the Editor and only those papers that meet the scientific aim and scope and editorial standards of the Journal will be sent of outside review. Each original manuscript is reviewed by at least two reviewers. Manuscript reviewers are given 3 weeks to complete their reviews and an editorial decision is generally reached within 4-6 weeks of submission.

Authors of all articles should be aware of the following guidelines when submitting to *JDR CTR*. Manuscripts missing any required components of submission will be returned to the authors for correction prior to peer-review.

**GENERAL INFORMATION FOR SUBMITTING AUTHORS**

**Prior Publication:** Manuscripts submitted to the *JDR Clinical & Translational Research* are accepted for consideration giving the understanding that it contains original material that has not been submitted for publication or has been previously published elsewhere. Any form of publication other than an abstract only constitutes prior publication.

**Research Ethics and Patient consent:** Manuscript submission guidelines for the *JDR Clinical & Translational Research* follow the “Uniform Requirements for Manuscripts Submitted to Biomedical Journals” set forth by the International Committee of Medical Journal Editors (ICMJE). For additional information please visit the ICMJE web site at [http://www.icmje.org/](http://www.icmje.org/). Submitted manuscripts should conform to the ICMJE Recommendations for the Conduct, Reporting, Editing, and Publication of Scholarly Work in Medical Journals. All papers reporting animal and/or human studies must state in the methods section that the relevant Ethics Committee or Institutional Review Board provided (or waived) approval. Ensure you have provided the full name and institution of the review committee, in addition to the approval number.

For research articles, authors are required to state in the methods section whether participants provided informed consent and whether the consent was written or verbal.
Authors are required to follow the ICJME Recommendations for the Protection of Research Participants. Identifying information, including patients’ names, initials, or hospital numbers, should not be published in written descriptions, photographs, and pedigrees unless the information is essential for scientific purposes and the patient (or parent or guardian) gives written informed consent for publication. Informed consent for this purpose requires that a patient who is identifiable be shown the manuscript to be published.

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**Declaration of Conflicting Interests:** It is the policy of *JDR CTR* to require a declaration of conflicting interests from all authors enabling a statement to be carried within the paginated pages of all published articles. Ensure any perceived or actual conflicts of interest are included at the end of your manuscript, after the acknowledgements and prior to the references. If no conflict exists, include the statement “The Author(s) declare(s) that there is no conflict of interest.” The *JDR CTR* abides by the International Committee of Medical Journal Editors guidelines for the Ethical Considerations in the Conduct and Report of Research ([http://www.icmje.org](http://www.icmje.org)), which can be referenced for additional guidance.

**Definition of Authorship & Contributorship:** Papers should be submitted for consideration once consent is given by all contributing authors. Those submitting papers should carefully check that all those whose work contributed to the paper are acknowledged as contributing authors. The list of authors should include all those who can legitimately claim authorships. This is all who:

- a. Made a substantial contribution to the concept and design, acquisition of data or analysis and interpretation of data,
- b. Drafted the article or revised it critically for important intellectual content,
- c. Approved the version to be published.

Authors should meet the conditions of all points about. Each author should have participated sufficiently in the work to take public responsibility for the manuscript. These individuals should fully meet the criteria for authorship.

As stated in the Uniform Requirements for Manuscripts Submitted to Biomedical Journals, put forth by the ICMJE, the Journal considers the following as an accurate definition of contributorship:

**Contributors Listing in 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 assistance, or a department
chairperson who provided only general support. Editors should ask corresponding authors to declare whether they had assistance with study design, data collection, data analysis, or manuscript preparation.

If such assistance was available, the authors should disclose the identity of the individuals who provided this assistance and the entity that supported it in the published article. Financial and material support should also be acknowledged.

Groups of persons who have contributed materially to the paper but whose contributions do not justify authorship may be listed under such headings as “clinical investigators” or “participating investigators,” and their function or contribution should be described—for example, “served as scientific advisors,” “critically reviewed the study proposal,” “collected data,” or “provided and cared for study patients.” Because readers may infer their endorsement of the data and conclusions, these persons must give written permission to be acknowledged.

**Contributor Forms:** All rights to manuscripts will be transferred to the *JDR Clinical & Translational Research* upon submission. Submission of a manuscript will constitute each author’s agreement that the Journal holds all propriety rights in the manuscript submitted, including all copyrights. Upon acceptance, the corresponding author will be asked to sign a formal transfer of copyright. Only the corresponding author is required to complete a contributor form unless any co-authors are work-for-hire or government employees. If co-authors fall into either of these categories, the corresponding author should contact the editorial office at jdrctr@iadr.org for additional instruction.

Please note that the *JDR Clinical & Translational Research* secures completed contributor forms electronically via the SAGETrack online submission and review system. Without the completion of the contributor form for all co-authors listed, accepted manuscripts cannot continue into production, delaying publication.

**Reporting Guidelines:** The relevant EQUATOR Network reporting guidelines should be followed depending on the type of study. Other resources can be found at the National Library of Medicine’s Research Reporting Guidelines and Initiatives.

The following completed reporting guidelines are required to be uploaded with submissions based on the type of research the manuscript contains:

- **Health Technology Assessment (HTA) Reports** must follow the INAHTA Checklist. The checklist needs to be completed by authors submitting HTA reports.

- **Randomized clinical trials** must follow CONSORT guidelines (see www.consort-statement.org). The checklist needs to be completed by authors submitting clinical trials.

- **Authors of randomized clinical trials** are required to register their clinical trials in a public trials registry. Authors of manuscripts describing such studies are asked to submit the name of the registry and the study registration number prior to publication. Authors are asked to include their clinical trial registration number at the end of their abstracts. In accordance with the aforementioned “Uniform Requirements for Manuscripts Submitted to Biomedical Journals,” clinical trials will only be considered for publication if they are registered. The trial registry name, registry URL and registration number must be included at the end of the abstract.
• Non-randomized study designs require completion and submission of the appropriate STROBE checklist (http://www.strobe-statement.org/index.php?id=available-checklists).

• For epidemiologic reports on population estimates of health conditions, the GATHER checklist must be completed and submitted.

• Multi-methods Research studies must follow the MMAT 2011 checklist. The checklist needs to be completed by authors submitting mixed methods reports.

• Qualitative studies must follow the SRQR guidelines. The checklist needs to be completed by authors submitting qualitative studies.

• Systematic reviews and meta-analyses must follow the PRISMA checklist. The checklist needs to be completed by authors submitting reports on systematic reviews and meta-analyses.

Public Gene Data: Prior to submission, the JDR Clinical & Translational Research asks that novel gene sequences be deposited in a public database and the accession number provided to the Journal. Authors may want to use the following Journal approved databases:


EMBL: www.ebi.ac.uk/embl/Submission/index.html

DDBJ: www.ddbj.nig.ac.jp/index-e.html

Manuscript submissions including microarray data should include the information recommended by the MIAME guidelines in their submission, and/or identify the submission details for the experiments details to one of the publicly available databases such as Array Express or GEO. Information on MIAME, Array Express and GEO can be found by clicking on the corresponding links below:

MIAME: www.mged.org/Workgroups/MIAME/miame.html

ArrayExpress: www.ebi.ac.uk/arrayexpress


REQUIREMENTS FOR MAIN TEXT OF MANUSCRIPTS

The research must be original, not under publication consideration elsewhere and free of conflict of interest. Authors will be asked to verify this information at the time of submission.

Cover letters: are not required, unless there is something specific authors would like the Editor to know.

Title Pages: All submissions must include a title page. Title pages must contain all author contact information and institutional affiliations in your main text document. Title pages should include: abstract
Title: Titles can consist of a maximum of 100 characters (including spaces). Titles do not normally include numbers, acronyms, abbreviations or punctuation. The title should include sufficient detail for indexing purposes but be general enough for readers outside the field to appreciate what the paper is about.

Knowledge Transfer Statement: All manuscripts must include a statement for dissemination to patients, clinicians and policy makers on how the results of the study can contribute to improved health. Statements should appear above the abstract of the paper. This statement should not exceed 75 words.

Example of Knowledge Transfer Statement: “The results of this study can be used by clinicians when deciding which approach they wish to use when planning placement of mini-dental implants in an individual’s edentulous mandible. With consideration of cost and patient preference, this information could lead to more appropriate therapeutic decisions.”

Suggested Reviewers: Please submit the names and email addresses of four preferred reviewers when prompted by the SAGETrack system. Preferred reviewers cannot be colleagues in the contributors’ institution or present collaborators.

ONLINE SUBMISSION

Submissions to the JDR CTR are only accepted for consideration via the SAGETrack online manuscript submission site. Authors who do not have an active account within the system are required to create a new account by clicking, “Create Account,” on the log-in page. The system will prompt the authors through a step by step process to create their account. Once created authors can submit their manuscripts by entering their “Author Center” and clicking the button by “Click Here to Submit a New Manuscript.”

If any difficulty is encountered at any time during the account creation or submission process, authors are encouraged to contact JDR Clinical & Translational Research Clinical & Translational Research editorial office staff at jdrctr@iadr.org.
MANUSCRIPT REQUIREMENTS BY TYPE

The JDR Clinical & Translational Research Clinical & Translational Research accepts the following types of manuscripts for consideration:

Original Reports: The various categories of Original Report manuscripts each have maximum word limits that include the introductions, materials and methods, results and discussion. Abstracts, acknowledgements, figure legends and references are not included in the word limits. Each Original Report must contain a 300 word abstract. Figures or tables are limited to a total of 5. A maximum of 40 references are allowed.

Original Report Categories:

a. Epidemiologic Research, including Randomized Clinical Trials – 4,000 maximum word limit
   i. Randomized clinical trials must follow CONSORT guidelines. Completed checklists must be uploaded at the time of submission.
   ii. Non-randomized study designs require completion and submission of the appropriate STROBE checklists.
   iii. For epidemiologic reports on population estimates of health conditions, the GATHER checklist must be completed and submitted.

b. Health Services Research – 4,000 maximum word limit

c. Economic Research (including cost-effectiveness; willingness to pay, cost-benefit, etc.) – 4,000 maximum word limit

d. Health Technology Assessment Reports – 8,000 maximum word limit
   i. Health Technology Assessment Reports must follow the INAHTA or CHEERS checklist. Completed checklists must be uploaded at the time of submission.

e. Multi-methods Research – 8,000 maximum word limit*
   i. Multi-methods Research must follow the MMAT 2011 checklist. Completed checklists must be uploaded at the time of submission.

f. Qualitative Research – 8,000 maximum word limit
   i. Qualitative studies must follow the SRQR or the CIPH reporting guidelines. Completed checklists must be uploaded at the time of submission.

g. Participatory Research – 5,000 maximum word limit

* If you would like to submit a Multi-methods Research report that has more than 8,000 words, you must correspond with and receive permission from the Editor prior to submission. Inclusion of written permission from Editor will be required at the time of submission

Review Manuscripts: These systematic reviews & meta-analyses have a 5,000 maximum word limit including introductions, materials and methods, results and discussion. Abstracts, acknowledgements, figure legends and references are not included in the word limits. Each Review must contain a 300 word abstract. Figures or tables are limited to a total of 5. A maximum of 50 references are allowed.

a. Systematic Reviews and meta-analyses must follow the PRISMA checklist. Completed checklists must be uploaded at the time of submission.
**Invited Commentary**: A point of view on any relevant topic by journalists, politicians, policy makers, educators, opinion leaders, organizations, researchers. Etc. These commentaries have a 1,000 maximum word limit. A maximum of 5 references are allowed. No figures or tables are permitted.

**Perspectives**: Reflections written by any stakeholder (e.g. on being a dentist, patient, researcher, etc. The perspectives are monographs taken from one’s personal experiences and have a 1,000 maximum word limit.

**Special Communications**: These are reports that stem from organizations, committees, consensus meetings, etc.; they describe the group’s outlook on an important issue regarding care provision, education, public health, health policy or research. Authors are requested to contact the Editor prior to submission. These submissions have a 2,000 maximum word limit. Figures or table are limited to a total of 2. A maximum of 5 references are allowed.

**Letters to the Editor**: Letters must include evidence to support a position about the scientific or editorial content of the *JDR CTR*. Manuscripts submitted as a letter to editor have a limit of 250 words and 3 references. No figures or tables are permitted. Letters on published articles must be submitted within 3 months of the article’s print publication date.

**Guest Editorials**: A clear and substantiated position on issues of interest to the readership community can be considered for this manuscript type. Guest Editorials are limited to 1,000 words. A maximum of 5 references are allowed. No figures or tables are permitted.
FIGURE AND TABLE REQUIREMENTS

These guidelines are intended to aid authors in providing figures that will reproduce well in both print and online media. Submitting digital image files that conform to these guidelines will prevent delays in the review and publication processes, and maximize the published quality of your figures.

Files containing figures and tables should be clearly labeled to indicate their placement in the text or appendix. Tables should be viewable in a portrait view. Tables that are created in a landscape view are more suitable for an appendix.

**Figure Types:** JDR CTR figures can fall into one of three categories: **Continuous-tone images**, **Line-art images**, and **Combination images**. Each image type has specific requirements in terms of the resolution needed for publication and the file type’s best suited for the figure. See the following panels for examples and requirements.

**Continuous-tone Image**  
Minimum resolution: 300dpi.  
Preferred File Formats: TIFF, Bitmap.

**Line-art Image**  
Minimum resolution: 800dpi.  
Preferred File Formats: EPS, PowerPoint, Illustrator.

**Combination Image**  
Minimum resolution: 800dpi.  

**Figure Resolution:** In order for a figure to be used in publication, its Digital Image File must have the required resolution when it is created. The resolution cannot be raised after the original image is made. Attempting to do so (for example, with Adobe Photoshop’s “Image Size” command) results in the addition of artificial pixels that distort the image and lower its sharpness. The figures on the right show an example of this reduced sharpness.

Line-art supplied at high resolution (1000dpi).  
Using “Image Size” to go from 300 DPI to 1000 DPI.
Figure Fonts: Limit fonts used in any figure to Times, Times New Roman, Arial, Frutiger, and Sabon. Other fonts cannot be guaranteed to reproduce properly.

Color Figure Charges: If the online version is in color and the printed version in black and white, please submit separate files for each version. Figures should be identical except in color or grayscale. The cost of color figures in the print version will be borne by the authors. Rates for color reproduction are $300 per initial page of color and $150 for each additional page of color. However, there are no charges for figures and diagrams printed in black and white. Color figures may be included in the online version of JDR CTR with no extra charges.

REFERENCES

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SUPPLEMENTAL FILES

Additional supporting data may be referenced as a supplemental appendix for publication online only. All supplemental appendix files must be submitted with the manuscript for review. Supplementary files will be subjected to peer-review alongside the article.

Supplementary files will be uploaded as supplied. They will not be checked for accuracy, copyedited, typeset or proofread. The responsibility for scientific accuracy and file functionality remains with the authors. A disclaimer will be displayed to this effect with any supplementary material published. Supplemental files may include additional figures or tables that exceed the Journal's limit. Material intended for the supplemental appendix must have “supplemental” or “appendix” in the file name upon upload. When formatting your supplemental files, please follow these instructions:

- Authors should provide a single Word file with all Appendix content. Figures and tables should be included in the main Appendix file so they can appear immediately alongside their captions. High resolution figures may also be supplied separately if authors wish, but they also must be copied into the Word file so everything can be kept together.

- Remove all highlighting/other colors. Use one font throughout.

- Be sure to run spell check and proofread the text.

- The Appendix should include the title of the article and all authors. Page numbers are recommended.

- Figures and Tables should be labeled Appendix Figure/Table 1, Appendix Figure/Table 2, etc. Avoid labeling as S1, S2, and so forth.

- All table footnotes and figure legends should be included.
• Preferably, authors shouldn’t label separate parts as “Appendix 1”, “Appendix 2”, etc.; just use section heads as in a regular article.

**LANGUAGE EDITING**

Manuscripts submitted for publication consideration should be written in English. Prior to submission, if a manuscript would benefit from professional editing, authors may consider using a language-editing service. Suggestions for this type of service can be found at www.iadr.org/EditingServices. The *JDR Clinical & Translational Research* does not take responsibility for, or endorse these services, and their use has no bearing on acceptance of a manuscript for publication.

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Authors are required to specify during the submission process if their paper received funding from NIH, NIDCR, or the Wellcome Trust and provide the grant number.

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