Editorial Process

The manuscripts will be reviewed for possible publication with the understanding that they are being submitted to one journal at a time and have not been published, simultaneously submitted or already accepted for publication elsewhere. The manuscripts are rejected by the editorial office before a formal peer-review.

The Editorial office review all submitted manuscripts initially. Manuscripts with insufficient originality, serious scientific and technical flaws or lack of a significant message are rejected. All manuscripts received are duly acknowledged. Manuscripts are sent to two or more expert reviewers without revealing the identity of the contributors to the reviewers. Each manuscript is also assigned to a member of the editorial team, who based on the comments from the reviewers takes a final decision on the manuscript. The contributors will be informed about the reviewers’ comments and acceptance/rejection of the manuscript. The average submission to first decision time is about 3-4 weeks and about 65-70% of unsolicited manuscripts do not get published.

Articles accepted would be copy edited for grammar, punctuation, print style, and format. Page proofs will be sent to the corresponding author, which has to be returned within three days. Correction received after that period may not be included.

Clinical Trial Registry

All clinical trials from India must be registered with “clinical trials registry – India”. The trials conducted outside India may be registered with any other clinical trial registry. We recommend and making it mandatory to have registration number for all clinical trials submitted for publication from January 2020.

Authorship Criteria

Authorship credit should be based only on substantial contributions

1. Conception and design or acquisition of data or analysis and interpretation of data;
2. Drafting the article or revising it critically for important intellectual content;
3. Final approval of the version to be published.

Conditions 1, 2 and 3 must be met. Participation solely in the acquisition of funding or the collection of data does not justify authorship. General supervision of the research group is not sufficient for authorship. Each contributor should have participated sufficiently in the work to take public responsibility for appropriate
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Contributors should provide a description of what each of them contributed towards the manuscript. Description should be divided in following categories, as applicable: concepts, design, definition of intellectual content, literature search, clinical studies, experimental studies, data acquisition, data analysis, statistical analysis, manuscript preparation, manuscript editing and manuscript review. Author’s contributions will be printed on the first page of the article. One or more author should take responsibility of the integrity of the work as a whole from inception to published article and should be designated as ‘guarantor’.

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This file should provide

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4. If the manuscript was presented as part of a meeting, the organization, place, and exact date on which it was read. A full statement to the editor about all submissions and previous reports that might be regarded as redundant publication of the same or very similar work. Any such work should be referred to specifically, and referenced in the new paper. Copies of such material should be included with the submitted paper, to help the editor decide how to handle the matter.

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These include original research work done in dentistry. The text of original articles amounting to up to 3000 words (excluding Abstract, references, and Tables) should be divided into sections with the headings Abstract (structured) Key-words, Introduction, Material and Methods, Results and Discussion, References, Tables and Figure legends.

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with the ethical standards of the responsible committee on human experimentation (institutional or regional) and with the Helsinki Declaration of 1975, as revised in 2000 (available at http://www.wma.net/e/policy/17-c_e.html). For prospective studies involving human participants, authors are expected to mention the approval of (regional/ national/ institutional or independent Ethics Committee or Review Board, obtaining informed consent from adult research participants, and obtaining assent for children aged over 7 years participating in the trial. The age beyond which assent would be required could vary as per regional and/ or national guidelines. Ensure the confidentiality of subjects by desisting from mentioning participants’ names, initials, or hospital numbers, especially in illustrative material. When reporting experiments on animals, should indicate, whether the institutions or a national research council’s guide for or any national law on the care and use of laboratory animals was followed. Evidence for approval by a local Ethics Committee (for both human as well as animal studies) must be supplied by the authors on demand. Animal experimental procedures should be as humane as possible and the details of anesthetics and analgesics used should be clearly stated. The ethical standards of experiments must be in accordance with the guidelines provided by the CPCSEA and World Medical Association Declaration of Helsinki on Ethical Principles for Medical Research Involving Humans for studies involving experimental animals and human beings, respectively). The journal will not consider any paper which is ethically unacceptable. A statement on ethics committee permission and ethical practices must be included in all research articles under the ‘Materials and Methods’ section.

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Include a summary of key findings (primary outcome measures, secondary outcome measures, results as they relate to a prior hypothesis); Strengths and limitations of the study. Interpretation and implications in the context of the totality of evidence (is there a systematic review to refer to, if not, could one be reasonably done here and now?), what this study adds to the available evidence, any new possible mechanisms, etc); Controversies raised by this study; and Future research directions (for this particular research collaboration, underlying mechanisms, clinical research, etc).

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- Conflicts of interest disclosed

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