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.

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.

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.

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exceed five. A justification should be included if the number of contributors exceeds these limits. Two/three additional authors from other departments/specialties would be permissible if they have contributed significantly.

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**Contribution Details**

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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**Preparation Of The Manuscript**

**A. Title Page**

The Title page should carry

1. Types of manuscript : Original article, Case Report
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9. Acknowledgement, if any; one or more statements should specify 1) contributions that need acknowledge but do not justify authorship, such as general support by a departmental chair, 2) acknowledgments of technical help; and 3) acknowledgement of financial and material support, which should specify the nature of the support. This should be included in the title page of the manuscript and not in the main article file.
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11. Registration number of clinical trials.

**B. Abstract Page**

The second page should carry the full title of the manuscript and an abstract (of no more than 150 words for brief report and 250 words for original articles and other article types). The abstract should be structured for original articles. State the context (background), aims, settings and design, material and methods, statistical analysis used, results and conclusions. Below the abstract should provide 3 to 8 keyword, arranged alphabetically. The abstract should not be structured for a brief report, review article, brief communication and research methodology. Don’t consider reference in abstract.
C. Introduction

State the purpose and summarize the study or observation.

D. Materials and Methods

The Methods section should only include information that was available at the time the study was planned or protocol written; all information obtained during the conduct of the study belongs to the results section.

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Reporting Guidelines for Specific Study Designs

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<td>QUOROM</td>
<td>Systematic reviews and meta-analyses</td>
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When reporting studies on human indicate whether the procedures followed were in accordance 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 https://www.wma.net/what-we-do/education/medical-ethics-manual/). Do not use patients’ names, initials, or hospital numbers, especially in illustrative material. When reporting experiments on animals, indicate whether the institution’s or a national research council’s guide for, or any national law on the care and use of laboratory animals was followed.

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H. Discussion
Include summary of key findings (primary outcome measures, secondary outcome measures, results as they relate to a prior hypothesis); Strengths and limitations of the study (study question, study design, data collection, analysis and interpretation); 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, effects on patient care and health policy, possible mechanism); Controversies raised by this study; and Future research directions (for this particular research collaboration, underlying mechanisms, clinical research).

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Include clinical and imagine photographs in the article to have better impact on the readers.

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Preparation of Case Report

Follow the standard format for the article (Abstract, Key-words, Introduction, Cases History, Discussion and References).

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- Conflicts of interest disclosed

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