Techniques in Coloproctology

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Quick Start Guide

- Article Types- Curious about what types of papers Techniques in Coloproctology publishes? <u>Check here</u>.
- Article Details Does your article type need an abstract? What are Word Limits? Check here
- Include a Title Page. <u>Details here</u> if needed.
- Conflicts of Interest *Techniques* uses the ICMJE form to report on COI's . <u>Find the form here</u>. Submit along with your manuscript.
- Submitting a video file? Authors are encouraged to post video to <u>www.figshare.com</u>; and insert a link to the video at Fighsare within their manuscript, to better allow review of video file.
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Original Research presenting study results.

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The abstract does not need to be structured for the reviews; for the meta-analysis can be structured or non-structured. Meta-analyses will also be considered as long as they follow the instructions for reviews.

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Correspondence

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• Editorial

Assigned by the Editors.

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Assigned by editors.

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Original articles where the video plays a major role. See instructions for videos as supplementary material.

• Video Forum

Videos with a short (up to 800 words) description. See instructions for videos as supplementary material.

We no longer accept case reports, Cochrane digests, technical notes, or technical advances. Some of the material previously published in the category Case Report may fit the categories of Short Communication, or Last Image.

Word Limits for Each Article Type

Туре	Abstract?	Keywords?	Word Limit	Reference Limit	Figure/Table/ Illustration Limit
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Review	Yes	Yes	4000	120	12
Controversies	No	No	1500	12	
Short Communication	Structured	No	1500	12	3
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Authors Reply	No	No	600	5	2
Last Image	No	No	100	12	12
Multimedia Article	Yes	Yes	3000	45	12
Video Forum	No	No	800	5	
Invited Editorial	No	No	2500	30	8
Invited Comment	No	No	800	5	3

Manuscript Preparation & Submission

Manuscript Preparation

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Abstract

Please provide a structured abstract of 150 to 250 words which should be divided into the following sections:

- Background (stating the main purposes and research question)
- Methods
- Results
- Conclusions
- Trial registration number and date of registration for prospectively registered trials (when applicable)
- Trial registration number and date of registration, followed by "retrospectively registered", for retrospectively registered trials (*when applicable*)

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At the end of the Methods section, briefly describe the statistical tests used for the analysis. Also include the statistical software used to perform the analysis, including the version and manufacturer, along with any extension packages (eg, the svy suite of commands in Stata or the survival package in R).

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Online document

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4) agree 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.

* Based on/adapted from:

ICMJE, Defining the Role of Authors and Contributors,

<u>Transparency in authors' contributions and responsibilities to promote integrity in scientific publication, McNutt at all,</u> <u>PNAS February 27, 2018</u>

Data transparency

All authors are requested to make sure that all data and materials as well as software application or custom code support their published claims and comply with field standards. Please note that journals may have individual policies on (sharing) research data in concordance with disciplinary norms and expectations.

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One author is assigned as Corresponding Author and acts on behalf of all co-authors and ensures that questions related to the accuracy or integrity of any part of the work are appropriately addressed.

The Corresponding Author is responsible for the following requirements:

- ensuring that all listed authors have approved the manuscript before submission, including the names and order of authors;
- managing all communication between the Journal and all co-authors, before and after publication;*
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• Free text:

All authors contributed to the study conception and design. Material preparation, data collection and analysis were performed by [full name], [full name] and [full name]. The first draft of the manuscript was written by [full name] and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

Example: CRediT taxonomy:

• Conceptualization: [full name], ...; Methodology: [full name], ...; Formal analysis and investigation: [full name], ...; Writing - original draft preparation: [full name, ...]; Writing - review and editing: [full name], ...; Funding acquisition: [full name], ...; Resources: [full name], ...; Supervision: [full name],....

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A Graduate Student's Guide to Determining Authorship Credit and Authorship Order, APA Science Student Council 2006

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- 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.

Recommendations can be downloaded from http://www.icmje.org/recommendations/

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Authors should include the following statements (if applicable) in a separate section entitled "Compliance with Ethical Standards" when submitting a paper:

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- Research involving Human Participants and/or Animals
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The corresponding author should be prepared to collect documentation of compliance with ethical standards and send if requested during peer review or after publication.

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Research involving human participants, their data or biological material

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When reporting a study that involved human participants, their data or biological material, authors should include a statement that confirms that the study was approved (or granted exemption) by the appropriate institutional and/or

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Although retrospective studies are conducted on already available data or biological material (for which formal consent may not be needed or is difficult to obtain) ethics approval may be required dependent on the law and the national ethical guidelines of a country. Authors should check with their institution to make sure they are complying with the specific requirements of their country.

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If human cells are used, authors must declare in the manuscript: what cell lines were used by describing the source of the cell line, including when and from where it was obtained, whether the cell line has recently been authenticated and by what method. If cells were bought from a life science company the following need to be given in the manuscript: name of company (that provided the cells), cell type, number of cell line, and batch of cells.

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Further information is available from the International Cell Line Authentication Committee (ICLAC).

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Examples:

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The trial registration number (TRN) and date of registration should be included as the last line of the manuscript abstract.

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Summary of requirements

The above should be summarized in a statement and placed in a 'Declarations' section before the reference list under a heading of 'Ethics approval'.

Examples of statements to be used when ethics approval has been obtained:

• All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. The study was approved by the Bioethics Committee of the Medical University of A (No. ...).

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• The questionnaire and methodology for this study was approved by the Human Research Ethics committee of the University of D (Ethics approval number: ...).

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• This research study was conducted retrospectively from data obtained for clinical purposes. We consulted extensively with the IRB of XYZ who determined that our study did not need ethical approval. An IRB official waiver of ethical approval was granted from the IRB of XYZ.

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All individuals have individual rights that are not to be infringed. Individual participants in studies have, for example, the right to decide what happens to the (identifiable) personal data gathered, to what they have said during a study or an interview, as well as to any photograph that was taken. This is especially true concerning images of vulnerable people (e.g. minors, patients, refugees, etc) or the use of images in sensitive contexts. In many instances authors will need to secure written consent before including images.

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Summary of requirements

The above should be summarized in a statement and placed in a 'Declarations' section before the reference list under a heading of 'Consent to participate' and/or 'Consent to publish'. Other declarations include Funding, Competing interests, Ethics approval, Consent, Data and/or Code availability and Authors' contribution statements.

Please see the various examples of wording below and revise/customize the sample statements according to your own needs.

Sample statements for "Consent to participate":

Informed consent was obtained from all individual participants included in the study.

Informed consent was obtained from legal guardians.

Written informed consent was obtained from the parents.

Verbal informed consent was obtained prior to the interview.

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The authors affirm that human research participants provided informed consent for publication of the images in Figure(s) 1a, 1b and 1c.

The participant has consented to the submission of the case report to the journal.

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Data availability statements

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