**DESCRIPTION**

Revista Española de Anestesiología y Reanimación (REDAR) is the scientific publication of the Sociedad Española de Anestesiología, Reanimación y Terapéutica del Dolor (SEDAR) (Spanish Society of Anaesthesiology, Resuscitation and Pain Relief).

REDAR, a monthly journal (10 issues), will consider for publication original scientific studies in relation to clinical anesthesia, resuscitation, critical/intensive care, the treatment of acute and chronic pain and emergency care.

All scientific contributions will be subject to an external anonymous peer review process (double blind). REDAR publishes mostly original scientific studies with clinical and experimental content, reviews, and consensus documents. Clinical cases, opinion articles and any other information of interest to specialists may also be published. The journal accepts work in both Spanish and English. The Journal is indexed in MEDLINE/PubMed, EMBASE, and SCOPUS.

For further information, consult [http://www.elsevier.es/redar](http://www.elsevier.es/redar).

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INTRODUCTION

*Revista Española de Anestesiología y Reanimación* (REDAR) is the scientific publication of the *Sociedad Española de Anestesiología, Reanimación y Terapéutica del Dolor* (SEDAR) (Spanish Society of Anaesthesiology, Resuscitation and Pain Relief).

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**Types of article**

**Originals.** Clinical research, or experimental work on animals, or basic sciences associated with any aspect of the in the field of the specialty (see "General Aspects"), with the following sections: Abstract, Introduction, Material and methods, Results, and Discussion. The length of the text will be limited to 12 DIN-A4 pages including a structured Abstract of 250 words, the Key Words, and up to a maximum of 30 literature references. Besides the text, up to 6 Figures or Tables will be accepted. This section requires graphical abstract and highlights. Please, see these sections further on in these guide for authors. A maximum of 6 will be allowed, except where justified. Retrospective, descriptive works that do not include the statistical treatment of the results will not be accepted. Clinical trials must be registered in a public data base before they start place and patients are recruited, following approval by the institutional or regional Clinical Research Ethics Committee (CEIC). The registration number and data base in which they are registered must be supplied. It will be obligatory for all clinical trials which start to recruit patients after 1 January 2017 to be registered in a public data base. Trials which recruited patients previously may still be sent to the REDAR for evaluation. When preparing controlled clinical trials, the CONSORT standards must be followed, which are available at [http://www.consort-statement.org](http://www.consort-statement.org). For observational studies, the points listed in the checklist available at [http://www.strobe-statement.org](http://www.strobe-statement.org) must be followed. Studies on the validity of diagnostic tests must follow the STARD standards available at [http://www.stard-statement.org](http://www.stard-statement.org).

**Short Originals.** Research works that due to their characteristics can be published in a shorter form. They will have the same structure as Originals, and will have a maximum length of 5 DIN-A4 pages including the structured abstract of 250 words, the Key Words, and up to a maximum of 15 literature references. As well as the text, up to 4 Figures or Tables will be accepted. A maximum of 6 authors will be permitted (see the "Originals" section above).

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Also a questionnaire of five questions with five possible responses, (with only one being true and with its corresponding explanation) must be provided. It is recommended to contact the Editor in advance if this type of work is going to be submitted.

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All authors should have made substantial contributions to all of the following: (1) the conception and design of the study, 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 submitted.

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Randomized controlled trials should be presented according to the CONSORT guidelines. At manuscript submission, authors must provide the CONSORT checklist accompanied by a flow diagram that illustrates the progress of patients through the trial, including recruitment, enrollment, randomization, withdrawal and completion, and a detailed description of the randomization procedure. The CONSORT checklist and template flow diagram are available online.

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If there is more than one appendix, they should be identified as A, B, etc. Formulae and equations in appendices should be given separate numbering: Eq. (A.1), Eq. (A.2), etc.; in a subsequent appendix, Eq. (B.1) and so on. Similarly for tables and figures: Table A.1; Fig. A.1, etc.
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