INFORMATION FOR AUTHORS

Acta Pharmaceutica (AP) is a fully open access journal, which publishes original research papers, short communications, preliminary communications and review articles, in the pharmaceutical and related sciences. The Journal also carries book reviews and obituaries. The last issue within each volume (No. 4) gives table of contents throughout the volume, as well as information to authors. Reviewers are acknowledged in the next issue.

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AIMS AND SCOPE

AP is an international, multidisciplinary scientific journal devoted to pharmaceutical and allied sciences which contains articles predominantly on core biomedical and health subjects. The aim of AP is to increase the impact of pharmaceutical research in academia, industry and laboratories. With strong emphasis on quality and originality, AP publishes reports from the discovery of a drug up to clinical practice. Topics covered span from analytics of drugs, through biochemistry, pharmaceutics/biopharmacy and pharmacokinetics, pharmacodynamics, cell biology, clinical pharmacy, drug design/delivery/disposition and stability, genetics, medicine (including diagnostics and therapy), pharmaceutical chemistry/medicinal chemistry, drug metabolism, molecular modelling/docking studies, pharmacology (clinical and animal), peptide and protein chemistry/protein design, nanosystems/pharmaceutical technology/formulation studies, cosmetology, pharmacognosy, radiopharmaceuticals, toxicology to pharmacoepidemiology/pharmacoeconomics, and more.

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Original research papers should contain unpublished results of original research, which should be presented in sufficient detail to ensure the reproducibility of the described experiments. Should not contain more than 60 literature citations and preferably not more than 10 appendices (including schemes, tables and figures).

Short communications provide reports on short, but completed research or descriptions of original laboratory techniques. They should not contain more than 30 literature citations.

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Meta-analysis reviews are also welcome.

Only papers providing previously unpublished scientific information will be entered in the first three of the above categories. Authors should specify to which of the four categories the submitted material should be allocated, but the Editorial Board reserves the right to make the final decision. Authors of review articles are advised to consult the Editorial Board prior to submitting the article.

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Manuscripts submitted to AP are only accepted on the understanding that they are subject to editorial pre-screening (desk evaluation which may result in immediate rejection) and peer-review of at least two independent referees, that they are submitted on an exclusive basis, and that recommendations to comply with ethical standards when performing clinical and other biological experiments have been adhered to. Submission implies that the material submitted to *Acta Pharmaceutica* is original and has not been published and is not in press or under consideration for publication (except in the form of 1-page abstract or as part of academic thesis), whole or in part, in print or electronic format, in any other journal or any other medium, including preprints, electronic journals and computer databases in the public domain.

The authors are advised to provide their papers in accordance with the practice of openness and authenticity, ethics and data transparency (through repositories or supplements). Availability of data should be mentioned.

Reports of biological/clinical research carried out in human subjects must contain a statement indicating approval by the local ethics committee (giving its address) and compliance with the Helsinki Declaration and its revisions, as well as an affirmation that written informed consent has been obtained from each patient. “Approval” is also needed for research with experimental animals, namely, for any use of biological material of human/animal origin.

Manuscripts should be submitted exclusively to the editorial office of *Acta Pharmaceutica via* e-mail address:

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Each article accepted for publication is language edited. The editorial staff reserves the right to make editorial corrections of the manuscript and adjust it to the requirements of the Journal.

During the submittance phase or just before commencing the reviewing process of the manuscript authors should sign *Authors' Agreement Form* and *Open Access License* and return it to the Editor-in-Chief or co-Editor (by e-mail).

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After initial desk review by the Editor-in-Chief, manuscript will be assigned to an appropriate co-Editor for a more competent preliminary review. After passing a preliminary review the manuscript is subject to peer reviewing process. The submitting author will receive acknowledgement of receipt of the

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Bear in mind that the Editor requires a fair, honest and unbiased assessment of the strengths and weaknesses of the manuscript. Both Editor and reviewer responsibilities include accountability, standards of objectivity and fairness, promptness and confidentiality.

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The Authors should give information on the source of funding, author´s contribution and ORCID, and conflict of interest, along with the Acknowledgements.

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S. A. Baron, C. Devaux, P. Colson, D. Raoult and J.-M. Rolain, Teicoplanin: an alternative drug for the treatment of COVID-19?, *Int. J. Antimicrob. Agents* **55**(4) (2020) Article ID 105944 (2 pages); https://doi.org/10.1016/j.ijantimicag.2020.105944

F. Pan-Montojo, O. Anichtchik, Y. Dening, L. Knels, S. Pursche, R. Jung, S. Jackson, G. Gille, M. G. Spillantini, H. Reichmann and R. H. W. Funk, Progression of Parkinson's disease pathology is reproduced by intragastric administration of rotenone in mice, *PLoS One* **5**(1) (2010) e8762 (10 pages); https://doi.org/10.1371/journal.pone.0008762

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E. Mutschler and H. Derendorf, *Drug Actions, Basic Principles and Therapeutic Aspects*, Medpharm Scientific Publishers, Stuttgart 1995, pp. 16–26., or J. Feely, T. Pringle and D. MacLean, *Calcium Channel Blockers,* in *New Drugs* (Ed. J. Feely), 3rd ed., BMJ Publishing Group, London 1994, pp. 82–93.

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International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use, *ICH Harmonised Tripartite Guideline, Validation of Analytical Procedures: Text and Methodology Q2(R1)*, Current *Step 4* version, November 2005; https://www.ich.org/fileadmin/Public_Web_Site/ICH_Products/Guidelines/Quality/Q2_R1/Step4/Q2_R1__Guideline.pdf; last access date June 22, 2012.

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