

Special Issue

Clinical Application of Personalized Genomic Medicine

Message from the Guest Editor

The aim of this Special Issue is to present current best practices, emerging areas of development, and informatics approaches in applying personalized genomic medicine into the clinical environment. We are inviting the submission of original articles and reviews for this Special Issue. Articles may be focused on (though not limited to) the clinical application of: bioinformatics, phenotype-based approaches to diagnostic sequencing, genomic screening, polygenic risk scores, risk alleles and low penetrance variants, genomic-based blood typing, integrative reporting, and return of genomic results, including applications and methods communicating with the HER.

Guest Editor

Dr. Matthew Lebo

Department of Pathology, Harvard Medical School, Boston, MA 02115, USA

Deadline for manuscript submissions

closed (15 April 2021)



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*Journal of Personalized
Medicine*
Editorial Office
MDPI, Grosspeteranlage 5
4052 Basel, Switzerland
Tel: +41 61 683 77 34
jpm@mdpi.com

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About the Journal

Message from the Editor-in-Chief

Journal of Personalized Medicine is one of the few journals that covers the diverse areas involved in the field, including research at basic, translational, and clinical levels. It focuses on “omics”-level studies that seek to define the basis of interindividual variation in susceptibility for a disease, its prognosis or definition of clinical subsets, and response to therapy (pharmacogenomics). We are also interested in systems biology as it relates to interindividual variation, and research on new methodologies, informatics, and biostatistics, in the aforementioned areas.

Editor-in-Chief

Prof. Dr. Kenneth P.H. Pritzker

Department of Laboratory Medicine and Pathobiology, Department of Surgery, University of Toronto, 6 Queens Pk Crescent W.F, Toronto, ON M5S 3H2, Canada

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High Visibility:

indexed within Scopus, PubMed, PMC, Embase, and other databases.

Journal Rank:

CiteScore - Q1 (Medicine (miscellaneous))

Rapid Publication:

manuscripts are peer-reviewed and a first decision is provided to authors approximately 25 days after submission; acceptance to publication is undertaken in 5.8 days (median values for papers published in this journal in the second half of 2025).