

Special Issue

Real-World Perspective on Effectiveness, Safety, and Costs of Orphan Medicines in Neurology

Message from the Guest Editors

Orphan drug can be defined as a medicine for the diagnosis, prevention or treatment of a life-threatening or chronically debilitating condition that is rare (affecting not more than five in 10,000 people) or where the medicine is unlikely to generate sufficient profit to justify research and development costs.

In this Special Issue on “Real-World Perspective on Effectiveness, Safety, and Costs of Orphan Medicines in Neurology”, we aim to publish outstanding contributions in the key fields covered by the journal, which will make a great contribution to the community.

Thus, we believe that the JPM is an excellent platform to support such overview on the current real-world knowledge and novelties regarding effectiveness, safety and costs of orphan medicines in neurology.

We hope that our special issue will draw the attention of the readers, scientists, stakeholders and clinicians in general.

Guest Editors

Prof. Dr. Dinko Vitezić

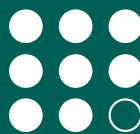
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About the Journal

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

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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).