Special Issue

Molecular Mechanisms of Kidney Diseases

Message from the Guest Editors

Both genetic mutations and epigenetic modifications are now understood to play critical roles in inherited and acquired kidney disorders. Variants in genes encoding essential structural or regulatory proteins can increase susceptibility to specific nephropathies, while epigenetic factors can influence how these diseases develop and progress. We invite researchers to contribute original research articles and comprehensive reviews that explore the molecular basis of kidney diseases. Relevant topics may include, but are not limited to, the following:

- Molecular signaling pathways in kidney injury and repair;
- Podocyte biology and glomerular barrier dysfunction;
- Mitochondrial and metabolic dysregulation in renal disease;
- Inflammatory and fibrotic mechanisms:
- Genetic and epigenetic contributors to kidney pathology;
- Novel molecular targets for therapy;
- Omics approaches and systems biology in nephrology.

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

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Rapid Publication:

manuscripts are peer-reviewed and a first decision is provided to authors approximately 21.5 days after submission; acceptance to publication is undertaken in 3.5 days (median values for papers published in this journal in the first half of 2025).

