

Special Issue

Advancing Nephrogenetics: Diagnosis, Genetic Testing, and Treatment Innovations

Message from the Guest Editors

This Special Issue's goal is to showcase state-of-the-art approaches to diagnosing hereditary kidney diseases, innovations in genetic testing technologies, and emerging therapeutic strategies grounded in molecular findings. We welcome contributions addressing a broad spectrum of topics within nephrogenetics, including but not limited to the following: Identification of novel genetic variants and mechanisms underlying monogenic or complex renal disorders; Diagnostic utility and clinical integration of next-generation sequencing (NGS), including whole-exome and whole-genome sequencing;

Genotype–phenotype correlations and their implications for disease stratification and management;

Advances in prenatal and preimplantation genetic diagnostics for hereditary kidney diseases;

Functional studies of disease-associated variants and their relevance to kidney pathophysiology;

Precision medicine and targeted therapies informed by genetic findings;

Ethical, psychological, and practical challenges in genetic testing and counseling in nephrology;

Bioinformatics tools and databases relevant to nephrogenetic research.

Guest Editors

Prof. Dr. Katarina Vukojević

Dr. Anita Racetin

Dr. Nela Kelam

Deadline for manuscript submissions

25 February 2026

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Editorial Office

MDPI, Grosspeteranlage 5

4052 Basel, Switzerland

Tel: +41 61 683 77 34

genes@mdpi.com

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About the Journal

Message from the Editor-in-Chief

Genes is central to our understanding of biology, and modern advances such as genomics and genome editing have maintained genetics as a vibrant, diverse and fast-moving field. There is a need for good quality, open access journals in this area, and the *Genes* team aims to provide expert manuscript handling, serious peer review, and rapid publication across the whole discipline of genetics. Starting in 2010, the journal is now well established and recognised. Why not consider *Genes* for your next genetics paper?

Editor-in-Chief

Prof. Dr. Selvarangan Ponnazhagan
Department of Pathology, The University of Alabama at Birmingham,
1825 University Blvd, SHEL 814, Birmingham, AL 35294-2182, USA

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