Special Issue

Advances in Diagnosis and Management of Multisystem Wilson's Disease

Message from the Guest Editors

Wilson's disease (WD) is an autosomal recessive, multisystem disorder of copper metabolism characterized by pathological copper accumulation in various tissues (primarily the liver and brain), leading to organ damage and clinical symptoms—predominantly hepatic and neuropsychiatric. As such, the management and treatment of WD require the involvement of multiple medical specialists (e.g., hepatologists, neurologists, psychiatrists, ophthalmologists, speech therapists, dietitians, etc.).

Advances in WD diagnosis are being made, particularly through improvements in genetics, neuroradiology, and serum copper metabolism biomarkers. Treatment options are also evolving, with new developments in gene therapy, anti-copper drugs, and symptomatic treatment.

The aim of this Special Issue is to provide a platform for the latest research on the diagnosis and treatment of WD, emphasizing the importance of multidisciplinary involvement. We invite submissions of various types of papers, including research articles, up-to-date reviews, case reports, and commentaries. We also encourage the formulation of hypotheses based on current knowledge that may inspire future research.

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Deadline for manuscript submissions

30 April 2026

Diagnostics

an Open Access Journal by MDPI

Impact Factor 3.3
CiteScore 5.9
Indexed in PubMed

mdpi.com/si/256348

Diagnostics
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You are cordially invited to submit research articles, short communications, comprehensive reviews, case reports or interesting images for consideration and publication in *Diagnostics* (ISSN 2075-4418). *Diagnostics* is published in open access format – research articles, reviews and other contents are released on the Internet immediately after acceptance. The scientific community and the general public have unlimited and free access to the content as soon as it is published. We would be pleased to welcome you as one of our authors.

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