

Special Issue

Advanced Research in Lysosomal Storage Disorders

Message from the Guest Editors

Lysosomal diseases (LDs) are a group of about 70 hereditary metabolic disorders caused by the inactivation of one of the genes encoding components of lysosomal machinery. LDs are rare diseases (combined incidence of about 1:5500) with mainly recessive inheritance. This Special Issue of *Biomedicines* will showcase novel developments in the field, presenting the state of the art on the screening, diagnosis, pathophysiology, and treatment of lysosomal diseases from all points of view (clinical, biochemical, genomic, and biomarkers) and will highlight questions that need to be addressed in the field.

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Deadline for manuscript submissions

closed (30 April 2024)



Biomedicines

an Open Access Journal
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Impact Factor 3.9
CiteScore 6.8
Indexed in PubMed



mdpi.com/si/109383

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Message from the Editor-in-Chief

Biomedicines (ISSN 2227-9059) is an open access journal devoted to all aspects of research on human health and disease, the discovery and characterization of new therapeutic targets, therapeutic strategies, and research of naturally driven biomedicines, pharmaceuticals, and biopharmaceutical products. Topics include pathogenesis mechanisms of diseases, translational medical research, biomaterial in biomedical research, natural bioactive molecules, biologics, vaccines, gene therapies, cell-based therapies, targeted specific antibodies, recombinant therapeutic proteins, nanobiotechnology driven products, targeted therapy, bioimaging, biosensors, biomarkers, and biosimilars. The journal is open for publication of studies conducted at the basic science and preclinical research levels. We invite you to consider submitting your work to *Biomedicines*, be it original research, review articles, or developing Special Issues of current key topics.

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