Special Issue

Advances in Lysosomal Disorders: From Molecular Mechanisms to Diagnosis

Message from the Guest Editors

Anderson–Fabry disease is a rare lysosomal storage disorder associated with mutations in the GLA gene. In recent years, thanks to multidisciplinary research efforts involving geneticists, pediatricians, nephrologists, cardiologists, neurologists, internists, and pathologists, among others, much knowledge has been gained about the physiopathogenetic mechanism and the initiators and effectors of tissue damage. New and old biomarkers are reported, which may implement the ability of clinicians to obtain a correct diagnosis and to follow the patients over time. The study of tissues allows us to obtain important information on the pathogenesis and progression of the damage.This issue aims to collect original works and review articles that open new diagnostic and/or pathophysiological perspectives on the disease.

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Biomolecules is a multidisciplinary open-access journal that reports on all aspects of research related to biogenic substances, from small molecules to complex polymers. We invite manuscripts of high scientific quality that pertain to the diverse aspects relevant to organic molecules, irrespective of the biological question or methodology. We aim for a competent, fair peer review and rapid publication. Please look at some of the exciting work that has been published in *Biomolecules* so far. We would be delighted to welcome you as one of our authors.

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