



Genetic and Metabolic Molecular Research of Lysosomal Storage Disease 2.0

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Message from the Guest Editor

Lysosomal storage diseases (LSD) are a group of inherited metabolic disorders in which the defects of various lysosomal enzymes and regulatory proteins result in the accumulation of different macromolecules in these organelles. This Special Issue is devoted to publishing the results of such studies, including basic research on the molecular mechanisms of LSD, translational studies on novel therapies, and clinical investigations performed at the molecular level. Review articles on all these aspects are also welcome. Therefore, this Special Issue shall provide a comprehensive view on molecular aspects of various LSD.





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Message from the Editor-in-Chief

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