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Advance in the Mechanism and Treatment of Lysosomal Storage Disorders

Guest Editor:

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Deadline for manuscript submissions:
closed (15 August 2022)

Message from the Guest Editor

Lysosomal storage disorders (LSDs) are inherited metabolic diseases usually caused by the deficiency of an enzyme, resulting in substrate accumulation. The advent of enzyme replacement therapy (ERT) in the 1990s changed the natural history of Gaucher disease, the first treated LSD. Nowadays, many other ERTs are commercially available, but the morbidity and mortality of LSDs are still high. We invite researchers to contribute original studies, as well as review articles, addressing recent advances in the treatment of LSDs. Your ground-breaking research will contribute to the improvement of patients' lives in the near future

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Message from the Editor-in-Chief

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