

Glycogen Storage

Acid alpha-glucosidase or acid maltase deficiency

Pompe disease

Glycosaminoglycans (mucopolysaccharides) Storage

- α-L-iduronidase deficiency → mucopolyssacharidosis I
- Iduronate sulfatase deficiency → mucopolyssacharidosis II
- Heparan sulfamidase deficiency → mucopolyssacharidosis IIIA
- N-acetylglucosaminidase deficiency → mucopolyssacharidosis IIIB
 - Heparan-α-glucosaminide Nacetyltransferase deficiency →
- mucopolyssacharidosis IIIC
 N-acetylglucosamine 6-sulfatase
- Galactose-6-sulfate sulfatase deficiency->mucopolyssacharidosis IVA

deficiency→mucopolyssacharidosis IIID

- β-galactosidase deficiency → mucopolyssacharidosis IVB
- N-acetylgalactosamine-4-sulfatase deficiency →mucopolyssacharidosis VI
 - β-glucoronidase deficiency → mucopolyssacharidosis VII
 - Hyaluronidase deficiency -> mucopolyssacharidosis IX

Lysosomal Storage Disease

JS Bruner

Lysosomal Storage Disease:

Reviewing Lysosomal Storage Disease: Unlocking the Spellbinding Force of Linguistics

In a fast-paced world fueled by information and interconnectivity, the spellbinding force of linguistics has acquired newfound prominence. Its capacity to evoke emotions, stimulate contemplation, and stimulate metamorphosis is really astonishing. Within the pages of "Lysosomal Storage Disease," an enthralling opus penned by a very acclaimed wordsmith, readers attempt an immersive expedition to unravel the intricate significance of language and its indelible imprint on our lives. Throughout this assessment, we shall delve to the book is central motifs, appraise its distinctive narrative style, and gauge its overarching influence on the minds of its readers.

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