

Lipid Storage

including sphingomyelin and cholesterol accumulation



SMPD1 gene mutation cause acid sphingomyelinase activity deficiency



Niemann-Pick disease type A and B

Glycolipid Storage

Glucocerebrosidase deficiency



Gaucher's disease

Glycosphingolipid Storage

α-galactosidase A deficiency



Fabry diseases

Mutations of NPC1 and NPC2 genes cause impairment in cellular processing and transport of LDL-cholesterol



Niemann-Pick disease type C

Glycogen Storage

Acid alpha-glucosidase or acid maltase deficiency

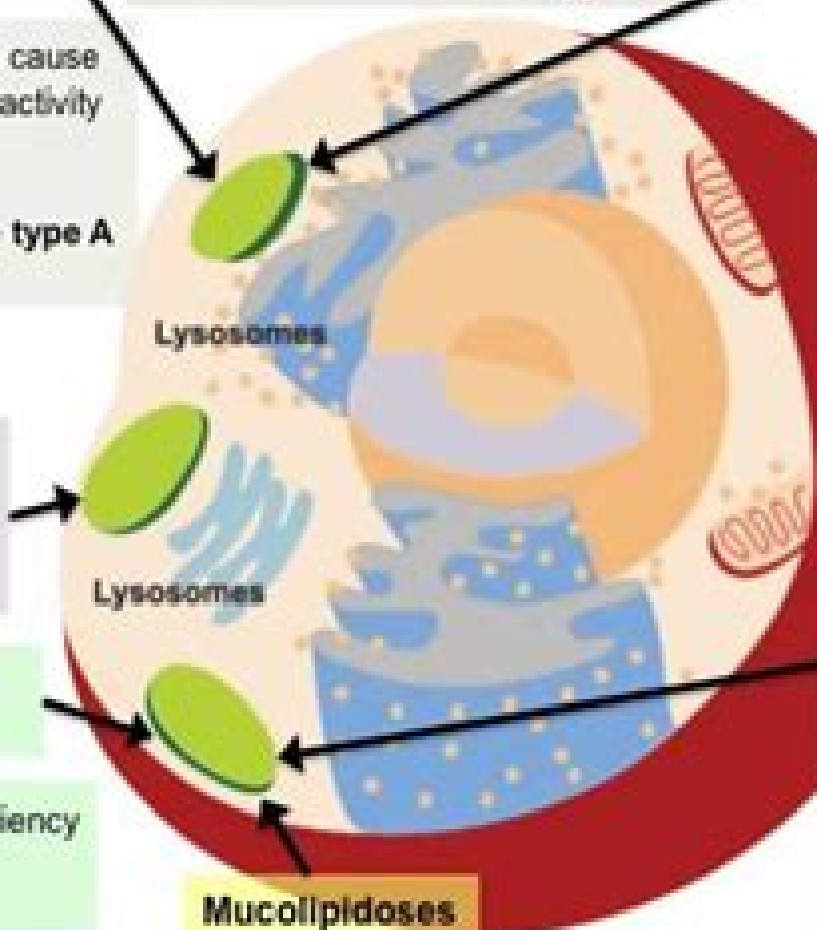


Pompe disease

Glycosaminoglycans (mucopolysaccharides) Storage

- α-L-iduronidase deficiency → mucopolysaccharidosis I
- Iduronate sulfatase deficiency → mucopolysaccharidosis II
- Heparan sulfamidase deficiency → mucopolysaccharidosis IIIA
- N-acetylglucosaminidase deficiency → mucopolysaccharidosis IIIB
- Heparan-α-glucosaminide N-acetyltransferase deficiency → mucopolysaccharidosis IIIC
- N-acetylglucosamine 6-sulfatase deficiency → mucopolysaccharidosis IIID
- Galactose-6-sulfate sulfatase deficiency → mucopolysaccharidosis IVA
- β-galactosidase deficiency → mucopolysaccharidosis IVB
- N-acetylgalactosamine-4-sulfatase deficiency → mucopolysaccharidosis VI
- β-glucuronidase deficiency → mucopolysaccharidosis VII
- Hyaluronidase deficiency → mucopolysaccharidosis IX

- α-N-acetyl neuraminidase (sialidase) → mucopolipidosis type I
- N-acetylglucosamine-1-phosphotransferase deficiency → mucopolipidosis type II and type III
- MCONLN1 gene mutation encoding mucopolipin1 → mucopolipidosis type IV



Lysosomal Storage Disease

Lei Huang



Lysosomal Storage Disease:

Unveiling the Energy of Verbal Art: An Emotional Sojourn through **Lysosomal Storage Disease**

In some sort of inundated with displays and the cacophony of instant interaction, the profound power and emotional resonance of verbal beauty usually disappear into obscurity, eclipsed by the continuous barrage of noise and distractions. Yet, nestled within the lyrical pages of **Lysosomal Storage Disease**, a fascinating work of literary beauty that impulses with raw thoughts, lies an wonderful journey waiting to be embarked upon. Published with a virtuoso wordsmith, this magical opus instructions visitors on a psychological odyssey, gently revealing the latent potential and profound influence stuck within the elaborate web of language. Within the heart-wrenching expanse of the evocative analysis, we shall embark upon an introspective exploration of the book is main subjects, dissect its charming publishing style, and immerse ourselves in the indelible effect it leaves upon the depths of readers souls.

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