

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

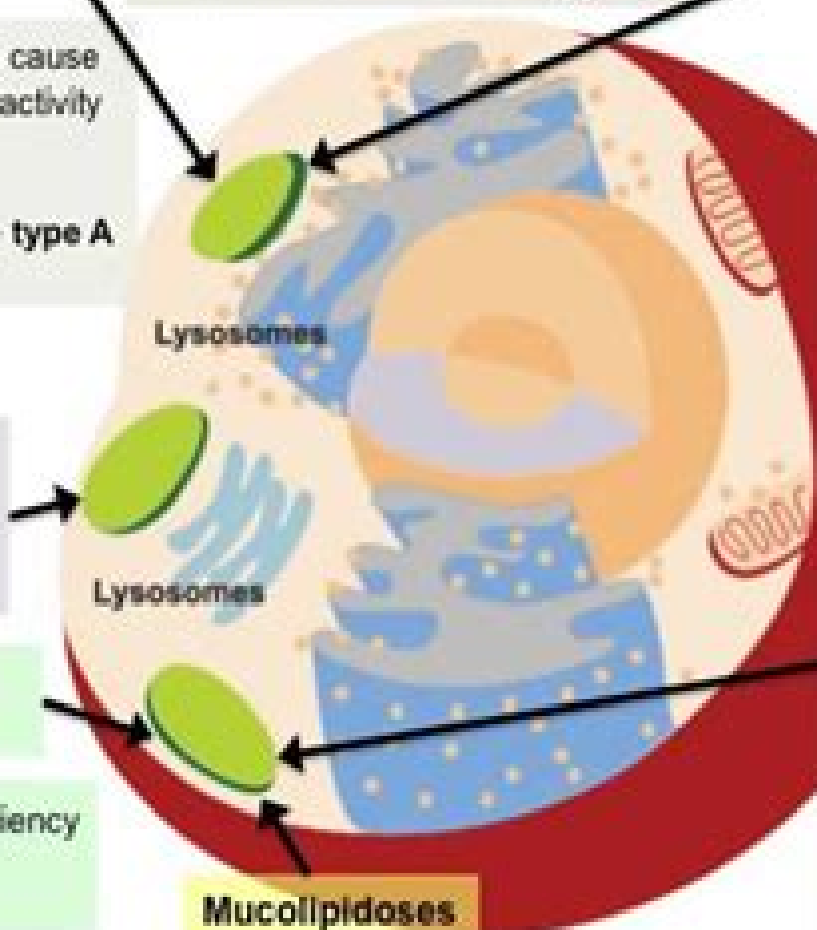


Pompe disease

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



Lysosomes And Lysosomal Storage Diseases

R Sandford



Lysosomes And Lysosomal Storage Diseases:

Unveiling the Energy of Verbal Artistry: An Mental Sojourn through **Lysosomes And Lysosomal Storage Diseases**

In a world inundated with displays and the cacophony of quick conversation, the profound power and mental resonance of verbal artistry usually fade into obscurity, eclipsed by the continuous barrage of sound and distractions. However, set within the lyrical pages of **Lysosomes And Lysosomal Storage Diseases**, a interesting function of literary brilliance that impulses with natural thoughts, lies an wonderful journey waiting to be embarked upon. Penned by a virtuoso wordsmith, that magical opus instructions visitors on a mental odyssey, gently revealing the latent possible and profound impact stuck within the complicated web of language. Within the heart-wrenching expanse of this evocative analysis, we will embark upon an introspective exploration of the book is main subjects, dissect its interesting publishing type, and immerse ourselves in the indelible impact it leaves upon the depths of readers souls.

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