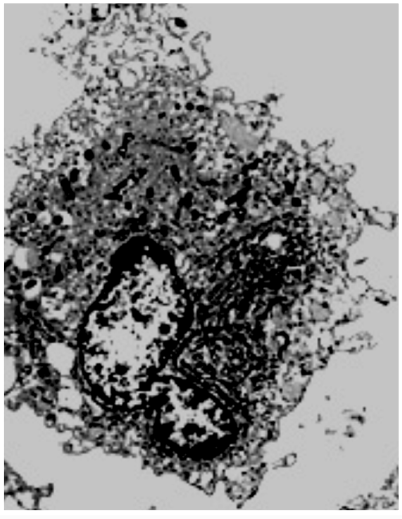
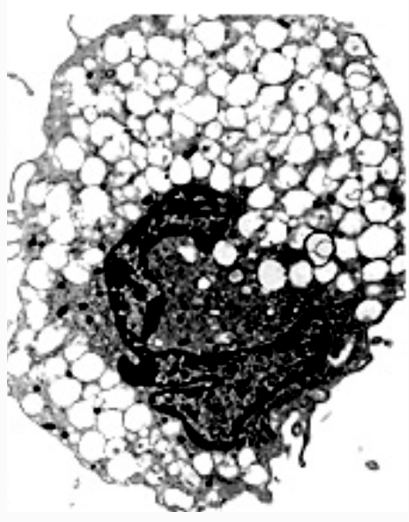
Histological look at the pathophysiology of GAG accumulation in MPS VI [1]

Normal and abnormal lysosomal enzyme function



Cells without GAG accumulation in the lysosomes



Cells with GAG accumulation in the lysosomes

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Abbreviations: GAG, glycosaminoglycan; MPS VI, mucopolysaccharidosis VI.

Reference: 1. Kakkis ED. Enzyme replacement therapy for the mucopolysaccharide storage disorders. *Expert Opin Investig Drugs.* 2002;11(5):675-685.