Glycogen Storage Disease Type VI

National Cancer Institute

Source

National Cancer Institute. Glycogen Storage Disease Type VI. NCI Thesaurus. Code C126875.

An autosomal recessive sub-type of glycogen storage disease caused by mutation(s) in the PYGL gene, encoding glycogen phosphorylase, liver form. The condition is characterized by mild-moderate hypoglycemia, growth retardation and hepatomegaly.