

Opis choroby *

Definicja

Glycogenesis due to glucose-6-phosphatase (G6P) deficiency or glycogen storage disease, (GSD), type 1, is a group of inherited metabolic diseases, including types a and b (see these terms), and characterized by poor tolerance to fasting, growth retardation and hepatomegaly resulting from accumulation of glycogen and fat in the liver.

Dane

Klasyfikacja

Choroba

Synonimy

G6P deficiency

Choroba spichrzania glikogenu spowodowana

niedoborem G6P

Choroba spichrzania glikogenu typu 1

Choroba Von Gierke

Glikogenoza typu 1

Glikogenoza wątrobowo-nerkowa

GSD spowodowane niedoborem G6P

GSD typu 1

Niedobór G6P

GSD due to G6P deficiency

GSD type 1

GSD type I

Glycogen storage disease due to G6P deficiency

Glycogen storage disease type 1

Glycogen storage disease type I

Glycogenesis type 1

Glycogenesis type I

Hepatorenal glycogenosis

Von Gierke disease

Kod ORPHA

364

Kod OMIM

232240

Kod ICD10

E74.0

Kod ICD11

5C51.3

[*Źródło](#)

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