

Opis choroby *

Definicja

Dimethylglycine dehydrogenase deficiency is an extremely rare autosomal recessive glycine metabolism disorder characterized clinically in the single reported case to date by muscle fatigue and a fish-like odor.

Dane

Klasyfikacja	Synonimy
Choroba	DMG dehydrogenase deficiency Niedobór dehydrogenazy DMG Niedobór DMGDH DMGDH deficiency

Kod ORPHA	Kod OMIM	Kod ICD10
243343	605850	E72.5

Kod ICD11

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*Źródło

orphanet