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
Choroba

Synonimy
DMG dehydrogenase deficiency
Niedobór dehydrogenazy DMG
Niedobór DMGDH
DMGDH deficiency

Kod ORPHA
243343

Kod OMIM
605850

Kod ICD10
E72.5

Kod ICD11
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*Źródło

orphanet