Glutaric acidemia type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Glutaric acidemia type 3. ORPHA:35706

Glutaryl-CoA oxidase deficiency is a peroxisomal disorder leading to glutaric aciduria. The prevalence is unknown. There is no distinctive phenotype associated with this disorder and one of the reported cases was asymptomatic. Transmission appears to be autosomal recessive.