Rodzinny łagodny Niedobór miedzi
Kod Orpha: 1551 Kod OMIM: 121270

Opis choroby *

Definicja
Familial benign copper deficiency is a rare disorder of mineral absorption and transport characterized by hypocupremia that manifests as failure to thrive, mild anemia, repeated seizures, hypotonia, and seborrheic skin. Spurring of the femur and tibia are also noted on radiographic imaging. Symptoms are reversible or improve with supplements of oral copper. There have been no further descriptions in the literature since 1988.
Dane

Klasyfikacja
Choroba

Synonimy
Familial benign hypocupremia

Kod ORPHA
1551

Kod OMIM
121270

Kod ICD10
E83.0

Kod ICD11
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.