Zespół Weismanna i Nettera
Kod Orpha: 3344 Kod OMIM: 112350

Opis choroby *

Definicja
Weismann-Netter syndrome is a rare, genetic, primary, bent bone dysplasia characterized by anterior diaphyseal bowing of the tibia and fibula, broadening of the fibula, posterior cortical thickening of both bones and short stature. Additional skeletal abnormalities include scoliosis with marked lumbar lordosis, horizontal sacrum and square iliac wings and/or, less frequently, vertebral malformations, abnormal shape of the clavicles and ribs, calvarial hyperostosis and delayed eruption of permanent teeth. Delayed ambulation is also frequently associated.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Anterior bowing of legs with dwarfism
WNS
Weismann-Netter-Stuhl syndrome

Kod ORPHA
3344

Kod OMIM
112350

Kod ICD10
Q77.8

Kod ICD11
LD24.C

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.