

Choroba Caffey'a

Kod Orpha: 1310 Kod OMIM: 114000

Opis choroby *

Definicja

Caffey disease is an osteosclerotic dysplasia characterized by acute inflammation with massive subperiosteal new bone formation usually involving the diaphyses of the long bones, as well as the ribs, mandible, scapulae, and clavicles. The disease is associated with fever, irritability pain and soft tissue swelling, with onset around the age of 2 months and resolving spontaneously by the age of 2 years. However, prenatal disease onset has also been described.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

Infantile cortical hyperostosis
Dziecięca hiperostoza korowa

Kod ORPHA

1310

Kod OMIM

114000

Kod ICD10

M89.8

Kod ICD11

LD24.1Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.