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

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*Źródło

orphanet