

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	Synonimy
Zespół wad wrodzonych	Infantile cortical hyperostosis Dziecięca hiperostoza korowa

Kod ORPHA	Kod OMIM	Kod ICD10
1310	114000	M89.8

Kod ICD11
LD24.1Y

*Źródło

orphanet