Dziedziczny zespół mnogich zwapnień tętniczych i stawowych
Kod Orpha: 289601 Kod OMIM: 211800

Opis choroby *

Definicja
Hereditary arterial and articular multiple calcification syndrome is a very rare genetic vascular disease of autosomal recessive inheritance, described in less than 20 patients to date, characterized by adult-onset (as early as the second decade of life) isolated calcification of the arteries of the lower extremities (including the iliac, femoral, and tibial arteries) as well as the capsule joints of the fingers, wrists, ankles and feet, and that usually manifests with mild paresthesias of the lower extremities, intense joint pain and swelling, and early onset arthritis of affected joints.

Dane

Klasyfikacja
Choroba

Synonimy
CALJA
Calcification of joints and arteries
CALJA
Calcification of joints and arteries

Kod ORPHA
289601

Kod OMIM
211800

Kod ICD10
I77.8

Kod ICD11
-

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.