Zespół Weavera i Williamsa
Kod Orpha: 3448 Kod OMIM:

Opis choroby *

Definicja
Weaver-Williams syndrome is a multiple congenital anomalies syndrome characterized by moderate-to-severe intellectual disability, decreased muscle mass, microcephaly, facial dysmorphism (prominent ears, midfacial hypoplasia, small mouth and cleft palate), clinodactyly of the fingers, delayed osseous maturation and generalized bone hypoplasia. The syndrome has been described in a brother and sister and an autosomal recessive mode of inheritance has been suggested. There have been no further descriptions in the literature since 1977.
Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
3448

Kod OMIM
-

Kod ICD10
Q87.8

Kod ICD11
-

---

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.