## Opis choroby *

Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by the association of intellectual deficit, facial dysmorphism (a highly arched palate, pointed chin, and small mouth, hypotelorism, a long nose and large protruding ears), arachnodactyly, hypogenitalism (undescended testes and hypospadias) and failure to thrive.

Dane

**Klasyfikacja**
Zespół wad wrodzonych

Synonimy
Cranio-facio-digito-genital syndrome
Zespół czaszkowo-twarzowo-palcowo-płciowy

**Kod ORPHA**
2115

**Kod OMIM**
601095

**Kod ICD10**
Q87.8

**Kod ICD11**
-

---

*Źródło

orphanet