Opis choroby *

Definicja

A rare, genetic, syndromic intellectual disability characterized by severe intellectual disability, distinctive craniofacial features and variable multiple congenital anomalies including ocular, brain, urogenital and skeletal abnormalities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
2707

Kod OMIM
244450

Kod ICD10
Q87.0

Kod ICD11
-

---

*Źródło

orphanet