Opis choroby *

Definicja

A rare genetic multisystem disorder characterized by the variable association of retinal dystrophy, obesity, polydactyly, genitourinary and kidney anomalies, learning disability and hypogonadism, with a wide spectrum of other minor manifestations.

Dane

Klasyfikacja
Choroba

Synonimy
BBS
BBS

Kod ORPHA
110

Kod OMIM
617406

Kod ICD10
Q87.8

Kod ICD11
5A61.0

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*Źródło

orphanet