

## **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  
BBS  
BBS

#### Synonimy

**Kod ORPHA**  
110

**Kod OMIM**  
617406

**Kod ICD10**  
Q87.8

**Kod ICD11**  
5A61.0

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#### \*Źródło

orphanet