

## **Opis choroby \***

### Definicja

A rare, multiple congenital anomalies syndrome characterized by craniofacial dysmorphology, congenital heart disease, dermatological abnormalities (most commonly hyperkeratotic skin and sparse, curly hair), neurological manifestations (hypotonia, seizures), failure to thrive and intellectual disability.

### Dane

#### **Klasyfikacja**

Zespół wad wrodzonych CFC syndrome  
Zespół CFC

#### **Kod ORPHA**

1340

#### **Kod OMIM**

615280

#### **Kod ICD10**

Q87.8

#### **Kod ICD11**

LD27.0Y

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\*Źródło

orphanet