

## Opis choroby \*

### Definicja

A rare X-linked syndromic intellectual disability characterized by intellectual impairment of variable severity, progressive lower limb spasticity, and diffuse palmoplantar hyperkeratosis. Additional manifestations include pes cavus, extensor plantar responses, hand tremor, and mild dysmorphic facial features.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych Fitzsimmons-McLachlan-Gilbert syndrome  
Zespół Fitzsimmonsa, McLachlan i Gilberta

#### Synonimy

#### Kod ORPHA

2824

#### Kod OMIM

309560

#### Kod ICD10

G82.1

#### Kod ICD11

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#### \*Źródło

orphanet