## Opis choroby *

Definicja

A rare hereditary ataxia characterized by neurogenic muscular atrophy associated with signs of cerebellar ataxia, hypesthesia, degeneration of the retina, and diabetes mellitus. Onset of the disease is in adolescence and the course is slowly progressive. There have been no further descriptions in the literature since 1983.

Dane

**Klasyfikacja**
Choroba

Synonimy
Furukawa-Takagi-Nakao syndrome
Zespół Furukawa, Takagi i Nakao

**Kod ORPHA**
2579

**Kod OMIM**
158500

**Kod ICD10**
G11.1

**Kod ICD11**
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*Źródło

orphanet