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