Niepełnosprawność intelektualna sprzężona z chromosomem X - ataksja - apraksja
Kod Orpha: 85338 Kod OMIM:

Opis choroby *

Definicja
A rare, X-linked syndromic intellectual disability disorder characterized by non-progressive ataxia, apraxia, variable intellectual disability and/or visuospatial, visuographic and visuoconstructive dysfunctions in male patients. Seizures, congenital clubfoot and macroorchidism have also been associated. Partial clinical expression was noted in obligate female carriers. There have been no further descriptions in the literature since 1992.
Dane

Klasyfikacja
Choroba

Kod ORPHA
85338

Kod OMIM
-

Kod ICD10
G31.8

Kod ICD11
-

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.