Opis choroby *

Definicja

A complex form of hereditary spastic paraplegia characterized by spastic paraplegia, demyelinating peripheral sensorimotor neuropathy, poikiloderma (manifesting with loss of eyebrows and eyelashes in childhood in addition to delicate, smooth, and wasted skin) and distal amyotrophy (presenting after puberty). There have been no further descriptions in the literature since 1992.

Dane

Klasyfikacja
Choroba

Synonimy
Antinolo-Nieto-Borrego syndrome
Zespół Antinolo, Nieto i Borrego

Kod ORPHA
2821

Kod OMIM
182815

Kod ICD10
G11.4

Kod ICD11
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*Źródło

orphanet