## Opis choroby *

Definicja

A rare genetic neurodegenerative disease characterized by sudden onset of progressive motor deterioration and regression of developmental milestones. Manifestations include dystonia and muscle spasms, dysphagia, dysarthria, and eventually loss of speech and ambulation. Brain MRI shows predominantly striatal abnormalities. The disease is potentially associated with a fatal outcome.

Dane

**Klasyfikacja**
Choroba

Synonimy
Lenk-Ploski syndrome
Zespół Lenkego i Ploskiego

**Kod ORPHA**
497906

**Kod OMIM**
617054

**Kod ICD10**
G31.8

**Kod ICD11**
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*Źródło

orphanet