Opis choroby *

Definicja

A rare neurologic disease that is characterized by the early-onset of cerebellar signs, eye movement abnormalities and pyramidal signs.

Dane

Klasyfikacja
Choroba

Synonimy
SCA11
SCA11

Kod ORPHA
98767

Kod OMIM
604432

Kod ICD10
G11.8

Kod ICD11
8A03.16

---

*Źródło

orphanet