Opis choroby *

Definicja

A rare genetic disease characterized by slowly progressive cerebellar degeneration resulting in ataxia, oculomotor apraxia, and other cerebellar symptoms. There is an increased frequency of spontaneous chromosomal aberrations, as well as hypersensitivity to ionizing radiation, while telangiectasia is absent.

Dane

Klasyfikacja
Choroba

Synonimy
ATLD
ATLD

Kod ORPHA
251347

Kod OMIM
604391

Kod ICD10
G11.3

Kod ICD11
4A01.31

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*Źródło

orphanet