

Opis choroby *

Definicja

An heterogeneous group of cases that are clinically diagnosed as Werner syndrome (WS) but do not carry *WRN* gene mutations. Similar to classical WS caused by *WRN* mutations, patients generally exhibit an aged appearance and common age-related disorders at earlier ages compared to the general population.

Dane

Klasyfikacja

Choroba

Synonimy

Atypical progeroid syndrome

Atypowy zespół progeroidalny

Kod ORPHA

79474

Kod OMIM

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Kod ICD10

E34.8

Kod ICD11

LD2B

*Źródło

orphanet