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

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*Źródło

orphanet