Zespół Wernera
Kod Orpha: 902 Kod OMIM: 277700

Opis choroby *

Definicja
Werner syndrome (WS) is a rare inherited syndrome characterized by premature aging with onset in the third decade of life and with cardinal clinical features including bilateral cataracts, short stature, graying and thinning of scalp hair, characteristic skin disorders and premature onset of additional age-related disorders.

Dane

Klasyfikacja
Choroba

Synonimy
Adult progeria
Progeria dorosłych
WS
WS

Kod ORPHA
902

Kod OMIM
277700

Kod ICD10
E34.8

Kod ICD11
LD2B

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.