

Opis choroby *

Definicja

A rare genetic disease characterized by the triad of adult-onset moderate to severe bilateral sensorineural hearing loss, premature graying of scalp hair, and essential tremor manifesting as involuntary shaking of the head. Additional pigmentation abnormalities have not been reported in this syndrome.

Dane

Klasyfikacja

Choroba

Synonimy

Sensorineural deafness-early graying-essential tremor syndrome
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Kod ORPHA

66633

Kod OMIM

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Kod ICD10

H90.3

Kod ICD11

LD2H.Y

*Źródło

orphanet