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
Sensorineural deafness-early graying-essential tremor syndrome

Kod ORPHA
66633

Kod OMIM
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Kod ICD10
H90.3

Kod ICD11
LD2H.Y

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*Źródło

orphanet