Opis choroby *

Definicja

Crandall syndrome is characterized by progressive sensorineural deafness, alopecia and hypogonadism with LH and GH deficiencies. It has been described in three brothers. It resembles Björnstad's syndrome (see this term) that combines irregular pili torti and deafness. It is probably inherited as and autosomal recessive disorder.

Dane

Klasyfikacja
Choroba

Synonimy
Alopecia-deafness-hypogonadism syndrome
Zespół łysienia, głuchoty i hipogonadyzmu
Alopecia-hearing loss-hypogonadism syndrome
Alopecia-sensorineural deafness-hypogonadism syndrome
Alopecia-sensorineural hearing loss-hypogonadism syndrome

Kod ORPHA
202

Kod OMIM
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Kod ICD10
E23.0

Kod ICD11
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*Źródło

orphanet