

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 an 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