

Opis choroby *

Definicja

A rare genetic disease characterized by mild intellectual disability, osteoporosis, delayed bone age, macrocephaly with wormian bones and frontal bossing, anomalies of fingers, nails, and teeth, thoracic deformities, hyperextensibility of joints, as well as congenital amaurosis and paraplegia. There have been no further descriptions in the literature since 1981.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych Heide	Heide syndrome
	Zespół Heidego

Kod ORPHA	Kod OMIM	Kod ICD10
2787	-	Q87.5

Kod ICD11
LD24.KY

*Źródło

orphanet