

Opis choroby *

Definicja

A rare genetic disease characterized by lethal non-spherocytic, non-immune hemolytic anemia, in association with abnormalities of the external genitalia (such as micropenis and hypospadias). Reported dysmorphic features include flat occiput, dimpled earlobes, deep plantar creases, and increased space between the first and second toes. There have been no further descriptions in the literature since 1995.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	Water-West syndrome
	Zespół Watera i Westa

Kod ORPHA	Kod OMIM	Kod ICD10
1046	600461	D58.8

Kod ICD11

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*Źródło

orphanet