

Opis choroby *

Definicja

A rare autosomal anomaly defined by the presence of three copies of chromosome 8 in some cells of the body, and clinically characterized by facial dysmorphism, typically deep palmar and plantar creases, mild intellectual deficit and joint, urinary, cardiac and skeletal anomalies.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	Mosaic trisomy chromosome 8
	Zespół Warkany
	Trisomy 8 mosaicism
	Warkany syndrome

Kod ORPHA	Kod OMIM	Kod ICD10
96061	-	Q92.1

Kod ICD11
LD40.Y

*Źródło

orphanet