Opis choroby *

Definicja

Isochromosomy Yq is a rare gonosomy anomaly with a variable phenotype including a female phenotype with sexual development delay, streak gonads, short stature and Turner syndrome features and male phenotype with infertility due to azoospermia.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
98798

Kod OMIM
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Kod ICD10
Q98.6

Kod ICD11
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*Źródło

orphanet