

Opis choroby *

Definicja

A rare sex chromosome number anomaly disorder characterized, genetically, by the presence of an extra X and Y chromosome in males and, clinically, by tall stature, dysfunctional testes associated with infertility and insufficient testosterone production, cognitive, affective and social functioning impairments, global developmental delay, and an increased risk of congenital malformations.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

10

Kod OMIM

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Kod ICD10

Q98.8

Kod ICD11

LD50.3Y

*Źródło

orphanet