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

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*Źródło

orphanet