Opis choroby *

Definicja

A rare genetic endocrine disease characterized by intrauterine growth restriction, failure of an adolescent growth spurt with proportional adult short stature, insulin resistance, and early adulthood-onset diabetes. Minimal subluxation of the fifth metacarpal-phalangeal joint has been reported, while metaphyseal dysplasia is absent. Testicular volume is low, but fertility is normal. There is no evidence of primary adrenal insufficiency.

Dane

Klasyfikacja
Choroba

Kod ORPHA
436144

Kod OMIM
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Kod ICD10
Q87.1

Kod ICD11
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*Źródło

orphanet