Opis choroby *

Definicja

A rare endocrine disease characterized by excessively tall stature and rapid growth velocity due to growth hormone excess from a pituitary adenoma/hyperplasia occurring before closure of the epiphyseal growth plates. Additional features may include pubertal delay, visual defects, headache, excessive appetite, hyperhidrosis, menstrual irregularity, prognathism, coarse facial features and large hands/feet.

Dane

Klasyfikacja
Choroba

Synonimy
Hypophyseal gigantism
Dziecięce i młodzieńcze formy akromegalii
Infantile and juvenile forms of acromegaly

Kod ORPHA
99725

Kod OMIM
102200

Kod ICD10
E22.0

Kod ICD11
5A60.0

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*Źródło

orphanet