

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

*Źródło

orphanet