

## Opis choroby \*

### Definicja

A rare acquired endocrine disease related to excessive production of growth hormone (GH) and characterized by progressive somatic disfigurement (mainly involving the face and extremities) and systemic manifestations.

### Dane

### Klasyfikacja

Choroba

**Kod ORPHA**

963

**Kod OMIM**

300943

**Kod ICD10**

E22.0

**Kod ICD11**

5A60.0

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[\\*Źródło](#)

orphanet