

## Opis choroby \*

### Definicja

A rare, genetic, non-syndromic, obesity disease characterized by severe, early-onset obesity, associated with major hyperphagia and endocrine abnormalities, resulting from leptin receptor deficiency.

### Dane

### Klasyfikacja

Podtyp etiologiczny

#### Kod ORPHA

179494

#### Kod OMIM

614963

#### Kod ICD10

E66.8

#### Kod ICD11

5B81.Y

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### \*Źródło

orphanet