## Opis choroby *

Definicja

A rare genetic disease characterized by early-onset severe obesity due to mutations in single genes acting on the development and function of the hypothalamus or the leptin-melanocortin pathway, leading to disruption of energy homeostasis and endocrine dysfunction. Patients present with a body mass index over three standard deviations above normal at less than five years of age, accompanied by a variety of signs and symptoms according to the mutated gene, including hyperphagia, insulin resistance, reduced basal metabolic rate, or hypogonadism, among others.

Dane

**Klasyfikacja**
Choroba

Synonimy
Monogenic obesity due to a leptin-melanocortin pathway anomaly
Otyłość monogenowa z powodu anomalii szlaku leptyna-monokortyna

**Kod ORPHA**
98267

**Kod OMIM**
-

**Kod ICD10**
E66.8

**Kod ICD11**
-

---

*Źródło

orphanet