

## Opis choroby \*

### Definicja

A rare, genetic form of obesity characterized by severe early-onset obesity, hyperphagia, and variable presence of cognitive impairment and behavioral disorder, including autistic spectrum behavior, impaired concentration and memory deficit. Some patients present with Prader-Willi-like features such as hypotonia, developmental delay, intellectual disability, short stature, hypopituitarism and dysmorphic facial features.

### Dane

### Klasyfikacja

Podtyp etiologiczny

**Kod ORPHA**

369873

**Kod OMIM**

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**Kod ICD10**

E66.8

**Kod ICD11**

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### \*Źródło

orphanet