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