Otyłość spowodowana niedoborem SIM1
Kod Orpha: 369873 Kod OMIM:

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
-

Kod ICD10
E66.8

Kod ICD11
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.