

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	Kod OMIM	Kod ICD10
369873	-	E66.8
Kod ICD11		
-		

*Źródło

orphanet