Opis choroby *

Definicja

A rare, genetic form of obesity characterized by severe early-onset obesity, hyperphagia, insulin resistance with hyperinsulinemia, reduced adult final height, delayed speech and language development and a tendency for social isolation and aggressive behavior.

Dane

Klasyfikacja
Podtyp etiologiczny

Kod ORPHA
329249

Kod OMIM
-

Kod ICD10
E66.8

Kod ICD11
-

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*Źródło

orphanet