Opis choroby *

Definicja

A rare genetic, neurodevelopmental syndrome characterized by hypothalamic-pituitary dysfunction with severe hypotonia and feeding deficits during the neonatal period followed by an excessive weight gain period with hyperphagia with a risk of severe obesity during childhood and adulthood, learning difficulties, deficits of social skills and behavioral problems or severe psychiatric problems.

Dane

Klasyfikacja
Choroba

Synonimy
Prader-Labhart-Willi syndrome
Zespół Pradera, Labharta i Williego
Zespół Williego i Pradera

Kod ORPHA
739

Kod OMIM
615547, 176270

Kod ICD10
Q87.1

Kod ICD11
LD90.3

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*Źródło

orphanet