Opis choroby *

Definicja

A rare, genetic, non-syndromic, obesity disease characterized by severe, early-onset obesity, associated with major hyperphagia and endocrine abnormalities, resulting from leptin receptor deficiency.

Dane

Klasyfikacja
Podtyp etiologiczny

Kod ORPHA
179494

Kod OMIM
614963

Kod ICD10
E66.8

Kod ICD11
5B81.Y

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*Źródło

orphanet