Obesity due to SIM1 deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Obesity due to SIM1 deficiency. ORPHA:369873

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.