DISEASE:
Obesity due to congenital leptin deficiency

NAME:	Obesity due to congenital leptin deficiency
DESCRIPTION:	Congenital leptin deficiency is a form of monogenic obesity characterised by severe early-onset obesity and marked hyperphagia.
ORPHACODE:	66628
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	LEP
CREATED:	13 May 2019 - 01:02
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- Obesitas, early onset (gene panel)
- Obesity (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centrum Medische Genetica - UZ Antwerpen
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- leptin

Related Gene Panels

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- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Obesitas (genepanel) - UZA
- Obesitas, early onset (8 genes) - VUB