Belgian Genetic Tests database

DISEASE:
Obesity due to SIM1 deficiency

NAME:	Obesity due to SIM1 deficiency
DESCRIPTION:	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.
ORPHACODE:	369873
XREF(S):	Orphanet
ANALYTE(S):	SIM1
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- Obesitas, early onset (gene panel)
- Obesity (gene panel)

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- Centre de Génétique Humaine - CHU Sart-Tilman
- Centrum Medische Genetica - UZ Antwerpen
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Related Gene Panels

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