

DISEASE:
Obesity due to pro-opiomelanocortin deficiency

NAME:	Obesity due to pro-opiomelanocortin deficiency
DESCRIPTION:	Pro-opiomelanocortin (POMC) deficiency is a form of monogenic obesity resulting in severe early-onset obesity, adrenal insufficiency, red hair and pale skin.
ORPHACODE:	71526
SYNONYMS:	POMC deficiency
XREF(S):	Orphanet OMIM ICD-10 OMIM
ANALYTE(S):	POMC
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/2910>

RELATED CONTENT

Related Genetic Tests

- [Early-onset severe obesity](#)
- [Obesitas, early onset \(gene panel\)](#)
- [Obesity \(gene panel\)](#)

Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)
- [Centrum Medische Genetica - UZ Antwerpen](#)
- [Centrum Medische Genetica - UZ Brussel VUB](#)

Related Analytes

- [proopiomelanocortin](#)

Related Gene Panels

- [Early-onset severe obesity \(44 genes\) - ULG](#)
- [Obesitas \(genepanel\) - UZA](#)
- [Obesitas, early onset \(8 genes\) - VUB](#)