Congenital leptin deficiency

Congenital leptin deficiency is a condition that causes severe obesity beginning in the first few months of life. Affected individuals are of normal weight at birth, but they are constantly hungry and quickly gain weight. Without treatment, the extreme hunger continues and leads to chronic excessive eating (hyperphagia) and obesity. Beginning in early childhood, affected individuals develop abnormal eating behaviors such as fighting with other children over food, hoarding food, and eating in secret.

People with congenital leptin deficiency also have hypogonadotropic hypogonadism, which is a condition caused by reduced production of hormones that direct sexual development. Without treatment, affected individuals experience delayed puberty or do not go through puberty, and may be unable to conceive children (infertile).

Frequency

Congenital leptin deficiency is a rare disorder. Only a few dozen cases have been reported in the medical literature.

Causes

Congenital leptin deficiency is caused by mutations in the \textit{LEP} gene. This gene provides instructions for making a hormone called leptin, which is involved in the regulation of body weight. Normally, the body's fat cells release leptin in proportion to their size. As fat accumulates in cells, more leptin is produced. This rise in leptin indicates that fat stores are increasing.

Leptin attaches (binds) to and activates a protein called the leptin receptor, fitting into the receptor like a key into a lock. The leptin receptor protein is found on the surface of cells in many organs and tissues of the body including a part of the brain called the hypothalamus. The hypothalamus controls hunger and thirst as well as other functions such as sleep, moods, and body temperature. It also regulates the release of many hormones that have functions throughout the body. In the hypothalamus, the binding of leptin to its receptor triggers a series of chemical signals that affect hunger and help produce a feeling of fullness (satiety).

\textit{LEP} gene mutations that cause congenital leptin deficiency lead to an absence of leptin. As a result, the signaling that triggers feelings of satiety does not occur, leading to the excessive hunger and weight gain associated with this disorder. Because hypogonadotrophic hypogonadism occurs in congenital leptin deficiency, researchers suggest that leptin signaling is also involved in regulating the hormones that control sexual development. However, the specifics of this involvement and how it may be altered in congenital leptin deficiency are unknown.
Congenital leptin deficiency is a rare cause of obesity. Researchers are studying the factors involved in more common forms of obesity.

Inheritance Pattern
This condition is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

Other Names for This Condition
• LEPD
• leptin deficiency
• obesity due to congenital leptin deficiency
• obesity, morbid, due to leptin deficiency
• obesity, morbid, nonsyndromic 1
• obesity, severe, due to leptin deficiency

Diagnosis & Management

Genetic Testing Information
• What is genetic testing?
  /primer/testing/genetictesting
• Genetic Testing Registry: Leptin deficiency or dysfunction

Research Studies from ClinicalTrials.gov
• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22leptin+deficiency%22

Other Diagnosis and Management Resources
• Eunice Kennedy Shriver National Institute of Child Health and Human Development: How Are Obesity and Overweight Diagnosed?
  https://www.nichd.nih.gov/health/topics/obesity/conditioninfo/diagnosed
• Genetics of Obesity Study
  https://www.goos.org.uk/home
• National Heart, Lung, and Blood Institute: Overweight and Obesity
  https://www.nhlbi.nih.gov/health-topics/overweight-and-obesity
Additional Information & Resources

**Health Information from MedlinePlus**

- Encyclopedia: Appetite--Increased
  https://medlineplus.gov/ency/article/003134.htm
- Encyclopedia: Hypogonadotropic Hypogonadism
  https://medlineplus.gov/ency/article/000390.htm
- Encyclopedia: Hypothalamus
  https://medlineplus.gov/ency/article/002380.htm
- Health Topic: Obesity
  https://medlineplus.gov/obesity.html

**Additional NIH Resources**

- Eunice Kennedy Shriver National Institute of Child Health and Human Development: How Are Obesity and Overweight Diagnosed?
  https://www.nichd.nih.gov/health/topics/obesity/conditioninfo/diagnosed
- National Heart, Lung, and Blood Institute: Overweight and Obesity
  https://www.nhlbi.nih.gov/health-topics/overweight-and-obesity
- National Institute of Diabetes and Digestive and Kidney Diseases: Active at Any Size!
  https://www.niddk.nih.gov/health-information/weight-management/staying-active-at-any-size

**Educational Resources**

- Centers for Disease Control and Prevention: Obesity and Genetics
  https://www.cdc.gov/genomics/resources/diseases/obesity/
- MalaCards: congenital leptin deficiency
  https://www.malacards.org/card/congenital_leptin_deficiency
- Orphanet: Obesity due to congenital leptin deficiency
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=66628
- TeensHealth: Dealing With Feelings When You’re Overweight
- TeensHealth: Delayed Puberty
- You and Your Hormones: Leptin
  http://www.yourhormones.info/hormones/leptin.aspx

**Patient Support and Advocacy Resources**

- Obesity Action Coalition
  https://www.obesityaction.org/
Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28congenital+leptin+deficiency%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D

Catalog of Genes and Diseases from OMIM

- LEPTIN DEFICIENCY OR DYSFUNCTION
  http://omim.org/entry/614962

Sources for This Summary


  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10523015

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21410864

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22463805

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