Galactosemia

Galactosemia is a disorder that affects how the body processes a simple sugar called galactose. A small amount of galactose is present in many foods. It is primarily part of a larger sugar called lactose, which is found in all dairy products and many baby formulas. The signs and symptoms of galactosemia result from an inability to use galactose to produce energy.

Researchers have identified several types of galactosemia. These conditions are each caused by mutations in a particular gene and affect different enzymes involved in breaking down galactose.

Classic galactosemia, also known as type I, is the most common and most severe form of the condition. If infants with classic galactosemia are not treated promptly with a low-galactose diet, life-threatening complications appear within a few days after birth. Affected infants typically develop feeding difficulties, a lack of energy (lethargy), a failure to gain weight and grow as expected (failure to thrive), yellowing of the skin and whites of the eyes (jaundice), liver damage, and abnormal bleeding. Other serious complications of this condition can include overwhelming bacterial infections (sepsis) and shock. Affected children are also at increased risk of delayed development, clouding of the lens of the eye (cataract), speech difficulties, and intellectual disability. Females with classic galactosemia may develop reproductive problems caused by an early loss of function of the ovaries (premature ovarian insufficiency).

Galactosemia type II (also called galactokinase deficiency) and type III (also called galactose epimerase deficiency) cause different patterns of signs and symptoms. Galactosemia type II causes fewer medical problems than the classic type. Affected infants develop cataracts but otherwise experience few long-term complications. The signs and symptoms of galactosemia type III vary from mild to severe and can include cataracts, delayed growth and development, intellectual disability, liver disease, and kidney problems.

**Frequency**

Classic galactosemia occurs in 1 in 30,000 to 60,000 newborns. Galactosemia type II and type III are less common; type II probably affects fewer than 1 in 100,000 newborns and type III appears to be very rare.

**Causes**

Mutations in the *GALT*, *GALK1*, and *GALE* genes cause galactosemia. These genes provide instructions for making enzymes that are essential for processing galactose obtained from the diet. These enzymes break down galactose into another simple sugar, glucose, and other molecules that the body can store or use for energy.
Mutations in the *GALT* gene cause classic galactosemia (type I). Most of these genetic changes almost completely eliminate the activity of the enzyme produced from the *GALT* gene, preventing the normal processing of galactose and resulting in the life-threatening signs and symptoms of this disorder. Another *GALT* gene mutation, known as the Duarte variant, reduces but does not eliminate the activity of the enzyme. People with the Duarte variant tend to have much milder features of galactosemia.

Galactosemia type II results from mutations in the *GALK1* gene, while mutations in the *GALE* gene underlie galactosemia type III. Like the enzyme produced from the *GALT* gene, the enzymes made from the *GALK1* and *GALE* genes play important roles in processing galactose. A shortage of any of these critical enzymes allows galactose and related compounds to build up to toxic levels in the body. The accumulation of these substances damages tissues and organs, leading to the characteristic features of galactosemia.

**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**

- classic galactosemia
- epimerase deficiency galactosemia
- galactokinase deficiency disease
- galactose-1-phosphate uridylyl-transferase deficiency disease
- galactose epimerase deficiency
- GALE deficiency
- GALK deficiency
- GALT deficiency
- UDP-galactose-4-epimerase deficiency disease
- UTP hexose-1-phosphate uridylyltransferase deficiency

**Diagnosis & Management**

**Formal Diagnostic Criteria**

- ACT Sheet: Absent/Reduced Galactose-1-Phosphate Uridyltransferase (GALT)
  https://www.ncbi.nlm.nih.gov/books/NBK55827/bin/Galactose_GALT.pdf
- ACT Sheet: Increased Total Galactose with normal GALT
  https://www.ncbi.nlm.nih.gov/books/NBK55827/bin/Galactose.pdf
Genetic Testing Information

- What is genetic testing?
  /primer/testing/genetictesting

- Genetic Testing Registry: Deficiency of galactokinase

- Genetic Testing Registry: Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase

- Genetic Testing Registry: Galactosemia

- Genetic Testing Registry: UDPglucose-4-epimerase deficiency

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22galactosemia%22

Other Diagnosis and Management Resources

- Baby’s First Test: Classic Galactosemia
  https://www.babysfirsttest.org/newborn-screening/conditions/classic-galactosemia

- Baby’s First Test: Galactoepimerase Deficiency
  https://www.babysfirsttest.org/newborn-screening/conditions/galactoepimerase-deficiency

- Baby’s First Test: Galactokinase Deficiency
  https://www.babysfirsttest.org/newborn-screening/conditions/galactokinase-deficiency

- GeneReview: Classic Galactosemia and Clinical Variant Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK1518

- GeneReview: Duarte Variant Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK258640

- GeneReview: Epimerase Deficiency Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK51671

- MedlinePlus Encyclopedia: Galactose-1-phosphate uridylyltransferase
  https://medlineplus.gov/ency/article/003636.htm

- MedlinePlus Encyclopedia: Galactosemia
  https://medlineplus.gov/ency/article/000366.htm
Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Galactose-1-phosphate uridyltransferase
  https://medlineplus.gov/ency/article/003636.htm
- Encyclopedia: Galactosemia
  https://medlineplus.gov/ency/article/000366.htm
- Health Topic: Carbohydrate Metabolism Disorders
  https://medlineplus.gov/carbohydratemetabolismdisorders.html
- Health Topic: Genetic Brain Disorders
  https://medlineplus.gov/geneticbraindisorders.html
- Health Topic: Liver Diseases
  https://medlineplus.gov/liverdiseases.html
- Health Topic: Newborn Screening
  https://medlineplus.gov/newbornscreening.html

Genetic and Rare Diseases Information Center

- Galactosemia
  https://rarediseases.info.nih.gov/diseases/2424/galactosemia

Educational Resources

- Genetic Science Learning Center, University of Utah
  https://learn.genetics.utah.edu/content/disorders/singlegene/
- MalaCards: galactosemia
  https://www.malacards.org/card/galactosemia
- Merck Manual Consumer Version: Overview of Hereditary Metabolic Disorders
- Minnesota Department of Health: Duarte Galactosemia
  http://www.health.state.mn.us/divs/cfh/topic/diseasesconds/duartegalact.cfm
- New England Consortium of Metabolic Programs
  http://newenglandconsortium.org/for-families/galactosemia/galactosemia-guide-for-parents-of-babies/
- Orphanet: Galactosemia
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=352
- Virginia Department of Health
Patient Support and Advocacy Resources

- American Liver Foundation
  https://liverfoundation.org/for-patients/about-the-liver/diseases-of-the-liver/galactosemia/

- Galactosemia Foundation
  http://www.galactosemia.org/

- Metabolic Support UK
  https://www.metabolicsupportuk.org/

- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/galactosemia/

- Resource List from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/galactos.html

Clinical Information from GeneReviews

- Classic Galactosemia and Clinical Variant Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK1518

- Duarte Variant Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK258640

- Epimerase Deficiency Galactosemia
  https://www.ncbi.nlm.nih.gov/books/NBK51671

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Galactosemias%5BMAJR%5D%29+AND+%28galactosemia%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- GALACTOKINASE DEFICIENCY
  http://omim.org/entry/230200

- GALACTOSE EPIMERASE DEFICIENCY
  http://omim.org/entry/230350

- GALACTOSEMIA
  http://omim.org/entry/230400
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301691

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22483615

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16838075

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21290786

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25473725

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20978943
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3063539/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24273939

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16385452
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1380226/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/26143117

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