



GALE gene

UDP-galactose-4-epimerase

Normal Function

The *GALE* gene provides instructions for making an enzyme called UDP-galactose-4-epimerase. This enzyme enables the body to process a simple sugar called galactose, which is present in small amounts in many foods. Galactose is primarily part of a larger sugar called lactose, which is found in all dairy products and many baby formulas.

UDP-galactose-4-epimerase converts a modified form of galactose (UDP-galactose) to another modified sugar (UDP-glucose). Glucose is a simple sugar that is the main energy source for most cells. This enzyme also promotes the reverse chemical reaction, the conversion of UDP-glucose to UDP-galactose. UDP-galactose is used to build galactose-containing proteins and fats, which play critical roles in chemical signaling, building cellular structures, transporting molecules, and producing energy.

Health Conditions Related to Genetic Changes

Galactosemia

More than 20 mutations in the *GALE* gene have been identified in people with a form of galactosemia known as type III or galactose epimerase deficiency. The signs and symptoms of this condition begin shortly after birth and can vary from mild to severe. Most of the genetic changes alter a single protein building block (amino acid) in UDP-galactose-4-epimerase, which makes the enzyme unstable or impairs its usual function.

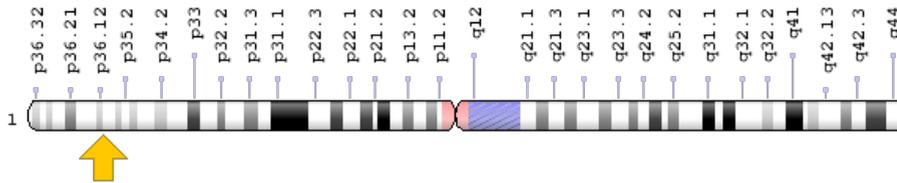
Some *GALE* gene mutations severely reduce or eliminate the activity of UDP-galactose-4-epimerase in all of the body's tissues. These genetic changes lead to a severe form of galactosemia type III described as the generalized form. A loss of enzyme activity prevents cells from processing galactose obtained from the diet. As a result, compounds associated with galactose processing can build up to toxic levels in the body. The accumulation of these substances damages tissues and organs, leading to serious complications such as clouding of the lens of the eye (cataract), intellectual disability, and damage to the liver, kidneys, and brain.

Other mutations in the *GALE* gene reduce the activity of UDP-galactose-4-epimerase in red blood cells only. These genetic changes underlie a much milder form of galactosemia type III described as the peripheral form. Affected individuals may not have any of the complications typically associated with galactosemia and often do not require treatment. Researchers are unclear why the effects of some *GALE* mutations are restricted to blood cells, while other mutations affect all of the body's tissues and cause severe health problems.

Chromosomal Location

Cytogenetic Location: 1p36.11, which is the short (p) arm of chromosome 1 at position 36.11

Molecular Location: base pairs 23,795,599 to 23,800,754 on chromosome 1 (Homo sapiens Updated Annotation Release 109.20190607, GRCh38.p13) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- galactowaldenase
- GALE_HUMAN
- SDR1E1
- UDP Galactose Epimerase
- UDP-Glucose 4-Epimerase
- UDP - Uridyl diphosphate galactose-4-epimerase
- Uridine diphosphate galactose-4-epimerase
- Uridine Diphosphate Glucose Epimerase

Additional Information & Resources

Educational Resources

- Essentials of Glycobiology (second edition, 2009): Figure: UDP-galactose Synthesis and Galactosemia
<https://www.ncbi.nlm.nih.gov/books/NBK1939/figure/ch42.f3/>

Clinical Information from GeneReviews

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

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28UDP-galactose-4-epimerase%5BTIAB%5D%29+OR+%28galactose+epimerase%5BTIAB%5D%29%29+OR+%28%28GALE%5BTIAB%5D%29+AND+%28galactosemia%5BTIAB%5D%29%29+AND+english%5BIa%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- UDP-GALACTOSE-4-EPIMERASE
<http://omim.org/entry/606953>

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
http://atlasgeneticsoncology.org/Genes/GC_GALE.html
- ClinVar
<https://www.ncbi.nlm.nih.gov/clinvar?term=GALE%5Bgene%5D>
- HGNC Gene Symbol Report
https://www.genenames.org/data/gene-symbol-report/#!/hgnc_id/HGNC:4116
- Monarch Initiative
<https://monarchinitiative.org/gene/NCBIGene:2582>
- NCBI Gene
<https://www.ncbi.nlm.nih.gov/gene/2582>
- UniProt
<https://www.uniprot.org/uniprot/Q14376>

Sources for This Summary

- Fridovich-Keil J, Bean L, He M, Schroer R. Epimerase Deficiency Galactosemia. 2011 Jan 25 [updated 2016 Jun 16]. In: Pagon RA, Adam MP, Ardinger HH, Wallace SE, Amemiya A, Bean LJH, Bird TD, Ledbetter N, Mefford HC, Smith RJH, Stephens K, editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2017. Available from <http://www.ncbi.nlm.nih.gov/books/NBK51671/>
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/21290786>
- Pey AL, Padín-Gonzalez E, Mesa-Torres N, Timson DJ. The metastability of human UDP-galactose 4'-epimerase (GALE) is increased by variants associated with type III galactosemia but decreased by substrate and cofactor binding. Arch Biochem Biophys. 2014 Nov 15;562:103-14. doi: 10.1016/j.abb.2014.07.030. Epub 2014 Aug 19.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/25150110>
- Schulz JM, Watson AL, Sanders R, Ross KL, Thoden JB, Holden HM, Fridovich-Keil JL. Determinants of function and substrate specificity in human UDP-galactose 4'-epimerase. J Biol Chem. 2004 Jul 30;279(31):32796-803. Epub 2004 Jun 2.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/15175331>

- Timson DJ, Lindert S. Comparison of dynamics of wildtype and V94M human UDP-galactose 4-epimerase-A computational perspective on severe epimerase-deficiency galactosemia. *Gene*. 2013 Sep 10;526(2):318-24. doi: 10.1016/j.gene.2013.05.027. Epub 2013 May 31.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/23732289>
Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3763920/>
 - Timson DJ. Functional analysis of disease-causing mutations in human UDP-galactose 4-epimerase. *FEBS J*. 2005 Dec;272(23):6170-7.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/16302980>
 - Timson DJ. The structural and molecular biology of type III galactosemia. *IUBMB Life*. 2006 Feb;58(2):83-9. Review.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/16611573>
-

Reprinted from Genetics Home Reference:
<https://ghr.nlm.nih.gov/gene/GALE>

Reviewed: August 2015
Published: September 10, 2019

Lister Hill National Center for Biomedical Communications
U.S. National Library of Medicine
National Institutes of Health
Department of Health & Human Services