G6PD gene
glucose-6-phosphate dehydrogenase

Normal Function

The *G6PD* gene provides instructions for making an enzyme called glucose-6-phosphate dehydrogenase. This enzyme, which is active in virtually all types of cells, is involved in the normal processing of carbohydrates. It plays a critical role in red blood cells, which carry oxygen from the lungs to tissues throughout the body. This enzyme helps protect red blood cells from damage and premature destruction.

Glucose-6-phosphate dehydrogenase is responsible for the first step in the pentose phosphate pathway, a series of chemical reactions that convert glucose (a type of sugar found in most carbohydrates) to another sugar, ribose-5-phosphate. Ribose-5-phosphate is an important component of nucleotides, which are the building blocks of DNA and its chemical cousin RNA. This chemical reaction produces a molecule called NADPH, which plays a role in protecting cells from potentially harmful molecules called reactive oxygen species. These molecules are byproducts of normal cellular functions. Reactions involving NADPH produce compounds that prevent reactive oxygen species from building up to toxic levels within cells. The production of NADPH by glucose-6-phosphate dehydrogenase is essential in red blood cells, which are particularly susceptible to damage by reactive oxygen species because they lack other NADPH-producing enzymes.

Health Conditions Related to Genetic Changes

Glucose-6-phosphate dehydrogenase deficiency

More than 200 mutations that cause glucose-6-phosphate dehydrogenase deficiency have been identified in the *G6PD* gene. Almost all of these mutations lead to changes in single building blocks (amino acids) in the glucose-6-phosphate dehydrogenase enzyme. These changes disrupt the normal structure and function of the enzyme or reduce the amount of the enzyme produced in cells.

Without enough functional glucose-6-phosphate dehydrogenase, red blood cells are unable to protect themselves from the damaging effects of reactive oxygen species. The damaged cells are likely to rupture and break down prematurely (undergo hemolysis). Factors such as infections, certain drugs, and ingesting fava beans can increase the levels of reactive oxygen species, causing red blood cells to undergo hemolysis faster than the body can replace them. This loss of red blood cells causes the signs and symptoms of hemolytic anemia, which is a characteristic feature of glucose-6-phosphate dehydrogenase deficiency.
Chromosomal Location

Cytogenetic Location: Xq28, which is the long (q) arm of the X chromosome at position 28

Molecular Location: base pairs 154,531,390 to 154,547,569 on the X chromosome (Homo sapiens Updated Annotation Release 109.20200228, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

• G6PD1
• G6PD_HUMAN

Additional Information & Resources

Educational Resources

• Biochemistry (fifth edition, 2002): Glucose 6-Phosphate Dehydrogenase Plays a Key Role in Protection Against Reactive Oxygen Species
  https://www.ncbi.nlm.nih.gov/books/NBK22389/

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28G6PD%5BTIAB%5D+%29+OR+%28glucose-6-phosphate+dehydrogenase%5BTIAB%5D%29%29+AND+%28%28glucose-6-phosphate+dehydrogenase%5BMAJR%5D%29+OR+%28glucose-6-phosphate:nadp++1-oxidoreductase%5BMAJR%5D%29%29+AND+%28%28Genes%5BMH%5D+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• GLUCOSE-6-PHOSPHATE DEHYDROGENASE
  http://omim.org/entry/305900
Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_G6PD.html
- ClinVar
- HGNC Gene Symbol Report
- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:2539
- NCBI Gene
- UniProt
  https://www.uniprot.org/uniprot/P11413

Sources for This Summary

- Biochemistry (fifth edition, 2002): Glucose 6-Phosphate Dehydrogenase Plays a Key Role in Protection Against Reactive Oxygen Species
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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18177777
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16311511
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/27040960
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