HBB gene
hemoglobin subunit beta

Normal Function
The HBB gene provides instructions for making a protein called beta-globin. Beta-globin is a component (subunit) of a larger protein called hemoglobin, which is located inside red blood cells. In adults, hemoglobin normally consists of four protein subunits: two subunits of beta-globin and two subunits of another protein called alpha-globin, which is produced from another gene called HBA. Each of these protein subunits is attached (bound) to an iron-containing molecule called heme; each heme contains an iron molecule in its center that can bind to one oxygen molecule. Hemoglobin within red blood cells binds to oxygen molecules in the lungs. These cells then travel through the bloodstream and deliver oxygen to tissues throughout the body.

Health Conditions Related to Genetic Changes

Beta thalassemia
Nearly 400 mutations in the HBB gene have been found to cause beta thalassemia. Most of the mutations involve a change in a single DNA building block (nucleotide) within or near the HBB gene. Other mutations insert or delete a small number of nucleotides in the HBB gene.

HBB gene mutations that decrease beta-globin production result in a type of the condition called beta-plus (B+) thalassemia. Mutations that prevent cells from producing any beta-globin result in beta-zero (B0) thalassemia.

Problems with the subunits that make up hemoglobin, including low levels of beta-globin, reduce or eliminate the production of this molecule. A lack of hemoglobin disrupts the normal development of red blood cells. A shortage of mature red blood cells can reduce the amount of oxygen that is delivered to tissues to below what is needed to satisfy the body's energy needs. A lack of oxygen in the body's tissues can lead to poor growth, organ damage, and other health problems associated with beta thalassemia.

Methemoglobinemia, beta-globin type
More than 10 mutations in the HBB gene have been found to cause methemoglobinemia, beta-globin type, which is a condition that alters the hemoglobin within red blood cells. These mutations often affect the region of the protein that binds to heme. For hemoglobin to bind to oxygen, the iron within the heme molecule needs to be in a form called ferrous iron (Fe^{2+}). The iron within the heme can
change to another form of iron called ferric iron (Fe\(^{3+}\)), which cannot bind oxygen. Hemoglobin that contains ferric iron is known as methemoglobin and is unable to efficiently deliver oxygen to the body's tissues.

In methemoglobinemia, beta-globin type, mutations in the \textit{HBB} gene alter the beta-globin protein and promote the heme iron to change from ferrous to ferric. This altered hemoglobin gives the blood a brown color and causes a bluish appearance of the skin, lips, and nails (cyanosis). The signs and symptoms of methemoglobinemia, beta-globin type are generally limited to cyanosis, which does not cause any health problems. However, in rare cases, severe methemoglobinemia, beta-globin type can cause headaches, weakness, and fatigue.

\textbf{Sickle cell disease}

Sickle cell anemia, a common form of sickle cell disease, is caused by a particular mutation in the \textit{HBB} gene. This mutation results in the production of an abnormal version of beta-globin called hemoglobin S or HbS. In this condition, hemoglobin S replaces both beta-globin subunits in hemoglobin. The mutation changes a single protein building block (amino acid) in beta-globin. Specifically, the amino acid glutamic acid is replaced with the amino acid valine at position 6 in beta-globin, written as Glu6Val or E6V. Replacing glutamic acid with valine causes the abnormal hemoglobin S subunits to stick together and form long, rigid molecules that bend red blood cells into a sickle (crescent) shape. The sickle-shaped cells die prematurely, which can lead to a shortage of red blood cells (anemia). The sickle-shaped cells are rigid and can block small blood vessels, causing severe pain and organ damage.

Mutations in the \textit{HBB} gene can also cause other abnormalities in beta-globin, leading to other types of sickle cell disease. These abnormal forms of beta-globin are often designated by letters of the alphabet or sometimes by a name. In these other types of sickle cell disease, just one beta-globin subunit is replaced with hemoglobin S. The other beta-globin subunit is replaced with a different abnormal variant, such as hemoglobin C or hemoglobin E.

In hemoglobin SC (HbSC) disease, the beta-globin subunits are replaced by hemoglobin S and hemoglobin C. Hemoglobin C results when the amino acid lysine replaces the amino acid glutamic acid at position 6 in beta-globin (written Glu6Lys or E6K). The severity of hemoglobin SC disease is variable, but it can be as severe as sickle cell anemia. Hemoglobin E (HbE) is caused when the amino acid glutamic acid is replaced with the amino acid lysine at position 26 in beta-globin (written Glu26Lys or E26K). In some cases, the hemoglobin E mutation is present with hemoglobin S. In these cases, a person may have more severe signs and symptoms associated with sickle cell anemia, such as episodes of pain, anemia, and abnormal spleen function.

Other conditions, known as hemoglobin sickle-beta thalassemias (HbSBetaThal), are caused when mutations that produce hemoglobin S and beta thalassemia occur together. Mutations that combine sickle cell disease with beta-zero (\(\beta^0\)) thalassemia
lead to severe disease, while sickle cell disease combined with beta-plus (B+) thalassemia is milder.

Other disorders

Hundreds of variations have been identified in the HBB gene. These changes result in the production of different versions of beta-globin. Some of these variations cause no noticeable signs or symptoms and are found when blood work is done for other reasons, while other variations may affect a person's health. Two of the most common variants are hemoglobin C and hemoglobin E.

Hemoglobin C (HbC), caused by the Glu6Lys mutation in beta-globin, is more common in people of West African descent than in other populations. People who have two hemoglobin C subunits in their hemoglobin, instead of normal beta-globin, have a mild condition called hemoglobin C disease. This condition often causes chronic anemia, in which the red blood cells are broken down prematurely.

Hemoglobin E (HbE), caused by the Glu26Lys mutation in beta-globin, is a variant of hemoglobin most commonly found in the Southeast Asian population. When a person has two hemoglobin E subunits in their hemoglobin in place of beta-globin, a mild anemia called hemoglobin E disease can occur. In some cases, the mutations that produce hemoglobin E and beta thalassemia are found together. People with this hemoglobin combination can have signs and symptoms ranging from mild anemia to severe thalassemia major.

Chromosomal Location

Cytogenetic Location: 11p15.4, which is the short (p) arm of chromosome 11 at position 15.4

Molecular Location: base pairs 5,225,464 to 5,227,071 on chromosome 11 (Homo sapiens Updated Annotation Release 109.20200228, GRCh38.p13) (NCBI)

Other Names for This Gene

- beta globin
- beta-globin
- HBB_HUMAN
- hemoglobin beta gene
- hemoglobin--beta locus
- hemoglobin, beta

**Additional Information & Resources**

**Educational Resources**

**Clinical Information from GeneReviews**
- Beta-Thalassemia https://www.ncbi.nlm.nih.gov/books/NBK1426
- Sickle Cell Disease https://www.ncbi.nlm.nih.gov/books/NBK1377

**Scientific Articles on PubMed**
- PubMed https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28HBB+gene%5BTI%5D+%29+OR+%28beta+hemoglobin%5BTI%5D%29+OR+%28beta+globin%5BTI%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

**Catalog of Genes and Diseases from OMIM**
- HEMOGLOBIN--BETA LOCUS http://omim.org/entry/141900

**Research Resources**
- Monarch Initiative https://monarchinitiative.org/gene/NCBIGene:3043
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/10791557

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16472167

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15561674

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15481886

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12126952

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16769681

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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18796252

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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15474138

• Thein SL. Genetic insights into the clinical diversity of beta thalassaemia. Br J Haematol. 2004 Feb; 
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14717773

• Thom CS, Dickson CF, Gell DA, Weiss MJ. Hemoglobin variants: biochemical properties and 
  cshperspect.a011858. Review. 
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23388674 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3579210/

• do Nascimento TS, Pereira RO, de Mello HL, Costa J. Methemoglobinemia: from diagnosis to 
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19082413

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