PC gene
pyruvate carboxylase

Normal Function

The PC gene provides instructions for making an enzyme called pyruvate carboxylase. This enzyme is active in mitochondria, which are the energy-producing centers within cells.

Pyruvate carboxylase is responsible for a chemical reaction that converts a molecule called pyruvate to another molecule called oxaloacetate. This reaction is essential for several different cellular functions. In the kidneys and liver, it is the first step in a process called gluconeogenesis. Gluconeogenesis generates glucose, a simple sugar that is the body's main energy source. This chemical reaction also occurs in the pancreas, where it helps regulate the secretion of a hormone called insulin. Insulin controls the amount of glucose in the blood that is passed into cells for conversion to energy.

In fat-storing (adipose) tissue, pyruvate carboxylase is involved in the formation of certain fats (lipogenesis). This enzyme also plays an important role in the nervous system, where it replenishes the building blocks needed to make brain chemicals called neurotransmitters. Additionally, pyruvate carboxylase is necessary for the formation of myelin, which is the fatty covering that insulates and protects certain nerve cells.

Health Conditions Related to Genetic Changes

Pyruvate carboxylase deficiency

More than 30 mutations in the PC gene have been identified in people with pyruvate carboxylase deficiency. This condition causes lactic acid and other potentially toxic compounds to accumulate in the blood. High levels of these substances can damage the body's organs and tissues, particularly in the nervous system.

Most PC gene mutations change a single protein building block (amino acid) in pyruvate carboxylase, which reduces the amount of this enzyme in cells or disrupts its ability to effectively convert pyruvate to oxaloacetate. Other genetic changes lead to the production of an abnormally short version of the enzyme that is completely nonfunctional.

If pyruvate carboxylase is missing or altered, it cannot carry out its role in generating glucose. Any disruption in gluconeogenesis impairs the body's ability to make energy in mitochondria. Additionally, a loss of pyruvate carboxylase allows lactic acid and ammonia, among other compounds, to build up and damage organs and tissues. Researchers suggest that the loss of pyruvate carboxylase function in the nervous
system, particularly the role of the enzyme in myelin formation and neurotransmitter production, also contributes to the neurologic features of pyruvate carboxylase deficiency.

**Chromosomal Location**

Cytogenetic Location: 11q13.2, which is the long (q) arm of chromosome 11 at position 13.2

Molecular Location: base pairs 66,848,522 to 66,958,418 on chromosome 11 (Homo sapiens Annotation Release 109, GRCh38.p12) (NCBI)

Credit: Genome Decoration Page/NCBI

**Other Names for This Gene**

- PCB
- PYC_HUMAN
- Pyruvic carboxylase

**Additional Information & Resources**

**Educational Resources**

- Basic Neurochemistry (sixth edition, 1999): Diseases of Mitochondrial Metabolism
  https://www.ncbi.nlm.nih.gov/books/NBK27914/

- Biochemistry (fifth edition, 2002): Glucose Can Be Synthesized from Noncarbohydrate Precursors
  https://www.ncbi.nlm.nih.gov/books/NBK22591/

**Clinical Information from GeneReviews**

- Pyruvate Carboxylase Deficiency
  https://www.ncbi.nlm.nih.gov/books/NBK6852
Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28pyruvate+carboxylase%5BTIAB%5D%29+AND+english%5BLa%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- PYRUVATE CARBOXYLASE
  http://omim.org/entry/608786

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_PC.html
- ClinVar
  https://www.ncbi.nlm.nih.gov/clinvar?term=PC%5Bgene%5D
- HGNC Gene Symbol Report
- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:5091
- NCBI Gene
- UniProt
  https://www.uniprot.org/uniprot/P11498

Sources for This Summary


Reprinted from Genetics Home Reference:

Reviewed: August 2017
Published: May 28, 2019

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