ABCG2 gene
ATP binding cassette subfamily G member 2 (Junior blood group)

Normal Function

The ABCG2 gene belongs to a group of genes called the ATP-binding cassette family; genes in this family provide instructions for making proteins that transport molecules across cell membranes. In the intestines, the ABCG2 protein helps release (secrete) a substance called urate into the urine. Urate is a byproduct of certain normal biochemical reactions in the body. In the bloodstream it acts as an antioxidant, protecting cells from the damaging effects of unstable molecules called free radicals. Urate levels are regulated by the kidneys and, to a lesser extent, by the intestines.

The ABCG2 protein also transports certain drugs out of cells. For example, this protein clears some chemotherapy drugs from organs and tissues. Transport of these drugs allows them to have their intended effects and be eliminated from the body.

Health Conditions Related to Genetic Changes

Gout

Genetic changes in the ABCG2 gene have been found to be associated with a condition called gout, which is a form of arthritis that causes painful joint inflammation.

ABCG2 gene changes associated with gout decrease the protein's ability to release urate. One variant replaces the protein building block (amino acid) glutamine with the amino acid lysine at position 141 in the protein (written as Gln141Lys or Q141K). This change reduces the protein's ability to secrete urate by half. Another variant creates a premature stop signal in the instructions for making the ABCG2 protein (written as Gln126Ter or Q126X), which results in no functional ABCG2 protein. Variants in the ABCG2 gene reduce the removal of urate in the blood, which causes the blood level of urate to rise. The excess urate can accumulate in the body's joints in the form of crystals, triggering an inflammatory response from the immune system and leading to gout.

While changes in the ABCG2 gene can alter urate levels in the body, they are not enough to cause gout by themselves. A combination of dietary, genetic, and other environmental factors play a part in determining the risk of developing this complex disorder.
Chromosomal Location

Cytogenetic Location: 4q22.1, which is the long (q) arm of chromosome 4 at position 22.1

Molecular Location: base pairs 88,090,264 to 88,231,626 on chromosome 4 (Homo sapiens Updated Annotation Release 109.20190607, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- ABC15
- ABCP
- ATP-binding cassette transporter G2
- ATP-binding cassette, sub-family G (WHITE), member 2 (Junior blood group)
- BCRP
- BCRP1
- BMDP
- breast cancer resistance protein
- CD338
- CDw338
- EST157481
- mitoxantrone resistance-associated protein
- MRX
- multi drug resistance efflux transport ATP-binding cassette sub-family G (WHITE) member 2
- MXR
- MXR-1
- MXR1
- placenta-specific ATP-binding cassette transporter
• placenta specific MDR protein
• UAQTL1

Additional Information & Resources

Educational Resources
• Biochemistry (fifth edition, 2002): Purines Are Degraded to Urate in Human Beings
  https://www.ncbi.nlm.nih.gov/books/NBK22372/#A3526
• Informed Health Online: Gout: Overview
  https://www.ncbi.nlm.nih.gov/books/NBK284934/

Scientific Articles on PubMed
• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28ABCG2%5BTI%5D%29+OR+%28ATP+binding+cassette+subfamily+G+member+2%5BTI%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29+AND+english%5Blia%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
• ATP-BINDING CASSETTE, SUBFAMILY G, MEMBER 2
  http://omim.org/entry/603756

Research Resources
• Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_ABCG2.html
• ClinVar
  https://www.ncbi.nlm.nih.gov/clinvar?term=ABCG2%5Bgene%5D
• HGNC Gene Symbol Report
• Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:9429
• NCBI Gene
• UniProt
  https://www.uniprot.org/uniprot/Q9UNQ0
Sources for This Summary

- OMIM: ATP-BINDING CASSETTE, SUBFAMILY G, MEMBER 2
  http://omim.org/entry/603756


