**FMO3 gene**
flavin containing monooxygenase 3

**Normal Function**

The *FMO3* gene provides instructions for making an enzyme that is part of a larger enzyme family called flavin-containing monooxygenases (FMOs). These enzymes break down compounds that contain nitrogen, sulfur, or phosphorus. The FMO3 enzyme, which is made chiefly in the liver, is responsible for breaking down nitrogen-containing compounds derived from the diet. One of these compounds is trimethylamine, which is the molecule that gives fish their fishy smell. Trimethylamine is produced as bacteria in the intestine help digest certain proteins obtained from eggs, liver, legumes (such as soybeans and peas), certain kinds of fish, and other foods. The FMO3 enzyme normally converts fishy-smelling trimethylamine into another compound, trimethylamine-N-oxide, which has no odor. Trimethylamine-N-oxide is then excreted from the body in urine.

Researchers believe that the FMO3 enzyme also plays a role in processing some types of drugs. For example, this enzyme is likely needed to break down the anticancer drug tamoxifen, the pain medication codeine, the antifungal drug ketoconazole, and certain medications used to treat depression (antidepressants). The FMO3 enzyme may also be involved in processing nicotine, an addictive chemical found in tobacco. Normal variations (polymorphisms) in the *FMO3* gene may affect the enzyme's ability to break down these substances. Researchers are working to determine whether *FMO3* polymorphisms can help explain why people respond differently to certain drugs.

**Health Conditions Related to Genetic Changes**

**Trimethylaminuria**

More than 25 mutations in the *FMO3* gene have been identified in people with trimethylaminuria. Most of these mutations lead to the production of a small, nonfunctional version of the FMO3 enzyme. Other mutations change single building blocks (amino acids) used to build the enzyme, which alters its shape and disrupts its function. Without enough functional FMO3 enzyme, the body is unable to convert trimethylamine into trimethylamine-N-oxide effectively. As a result, trimethylamine builds up in the body and is released in an affected person's sweat, urine, and breath. The excretion of this compound is responsible for the strong body odor characteristic of trimethylaminuria. Studies suggest that diet and stress also play a role in determining the intensity of the fish-like odor.
Chromosomal Location

Cytogenetic Location: 1q24.3, which is the long (q) arm of chromosome 1 at position 24.3

Molecular Location: base pairs 171,090,873 to 171,117,819 on chromosome 1 (Homo sapiens Updated Annotation Release 109.20190905, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

• Dimethylaniline monooxygenase [N-oxide-forming] 3
• Dimethylaniline oxidase 3
• FMO3_HUMAN
• FMOII

Additional Information & Resources

Clinical Information from GeneReviews

• Primary Trimethylaminuria
  https://www.ncbi.nlm.nih.gov/books/NBK1103

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28FMO3%5BTIAB%5D%29+OR+%28flavin+containing+monooxygenase+3%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

• FLAVIN-CONTAINING MONOOXYGENASE 3
  http://omim.org/entry/136132
Research Resources

- **Atlas of Genetics and Cytogenetics in Oncology and Haematology**
  http://atlasgeneticsoncology.org/Genes/GC_FMO3.html
- **ClinVar**
  https://www.ncbi.nlm.nih.gov/clinvar?term=FMO3%5Bgene%5D
- **HGNC Gene Symbol Report**
- **Monarch Initiative**
  https://monarchinitiative.org/gene/NCBIGene:2328
- **NCBI Gene**
- **UniProt**
  https://www.uniprot.org/uniprot/P31513

Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15975041
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