The **EIF2B2** gene provides instructions for making one of five parts of a protein called eIF2B, specifically the beta subunit of this protein. The eIF2B protein helps regulate overall protein production (synthesis) in the cell by interacting with another protein, eIF2. The eIF2 protein is called an initiation factor because it is involved in starting (initiating) protein synthesis.

Under some conditions, eIF2B increases protein synthesis by helping to recycle molecules called GTP, which carry energy to the initiation factor. Under other conditions, it slows protein synthesis by binding tightly to the initiation factor, which converts the eIF2B protein into an inactive form and prevents recycling of GTP.

Proper regulation of protein synthesis is vital for ensuring that the correct levels of protein are available for the cell to cope with changing conditions. For example, cells must synthesize protein much faster if they are multiplying than if they are in a resting state.

### Health Conditions Related to Genetic Changes

**Leukoencephalopathy with vanishing white matter**

Mutations in the **EIF2B2** gene have been identified in a few people with leukencephalopathy with vanishing white matter, including some affected females with a variant of the disorder in which the neurological features are accompanied by ovarian failure (ovarioleukodystrophy). These mutations cause partial loss of eIF2B function. Impairment of eIF2B function makes it more difficult for the body's cells to regulate protein synthesis and deal with changing conditions and stress. Researchers believe that cells in the white matter (nerve fibers covered by a fatty substance called myelin that insulates and protects nerves) may be particularly affected by an abnormal response to stress, resulting in the signs and symptoms of leukencephalopathy with vanishing white matter.
Chromosomal Location

Cytogenetic Location: 14q24.3, which is the long (q) arm of chromosome 14 at position 24.3

Molecular Location: base pairs 75,002,909 to 75,009,591 on chromosome 14 (Homo sapiens Annotation Release 109, GRCh38.p12) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- EI2BB_HUMAN
- EIF-2Bbeta
- EIF2B
- eukaryotic translation initiation factor 2B, subunit 2 (beta, 39kD)
- eukaryotic translation initiation factor 2B, subunit 2 beta, 39kDa

Additional Information & Resources

Educational Resources

- Eurekah Bioscience: Mechanism of Translation Initiation in Eukaryotes
  https://www.ncbi.nlm.nih.gov/books/NBK6597/

  https://www.ncbi.nlm.nih.gov/books/NBK26890/#A1387

Clinical Information from GeneReviews

- Childhood Ataxia with Central Nervous System Hypomyelination/Vanishing White Matter
  https://www.ncbi.nlm.nih.gov/books/NBK1258
Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28EIF2B2%5BTIAB%5D%29+OR+%28EIF2B%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- EUKARYOTIC TRANSLATION INITIATION FACTOR 2B, SUBUNIT 2
  http://omim.org/entry/606454

Research Resources

- ClinVar

- HGNC Gene Symbol Report

- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:8892

- NCBI Gene

- UniProt
  https://www.uniprot.org/uniprot/P49770

Sources for This Summary


- OMIM: EUKARYOTIC TRANSLATION INITIATION FACTOR 2B, SUBUNIT 2
  http://omim.org/entry/606454

- Eurekah Bioscience: Mechanism of Translation Initiation in Eukaryotes
  https://www.ncbi.nlm.nih.gov/books/NBK6597/


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