



SGSH gene

N-sulfoglucosamine sulfohydrolase

Normal Function

The *SGSH* gene provides instructions for producing an enzyme called sulfamidase. This enzyme is located in lysosomes, compartments within cells that digest and recycle different types of molecules. Sulfamidase is involved in the step-wise breakdown of large molecules called glycosaminoglycans (GAGs). GAGs are composed of sugar molecules that are linked together to form a long string. To break down these large molecules, individual sugars are removed one at a time from one end of the molecule. Sulfamidase removes a chemical group known as a sulfate from a sugar called glucosamine when it is at the end of the GAG chain.

Health Conditions Related to Genetic Changes

Mucopolysaccharidosis type III

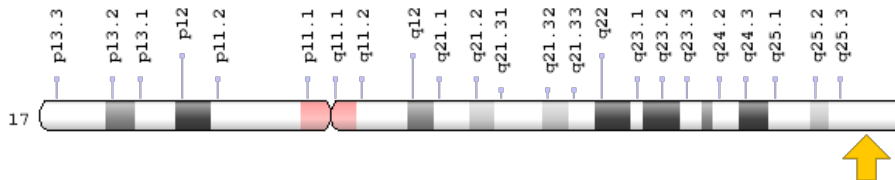
More than 80 mutations in the *SGSH* gene have been found to cause mucopolysaccharidosis type IIIA (MPS IIIA). Most of these mutations change single DNA building blocks (nucleotides) in the gene. All of the mutations that cause MPS IIIA reduce or eliminate the function of sulfamidase.

The lack of sulfamidase activity disrupts the breakdown of a subset of GAGs called heparan sulfate. As a result, partially broken down heparan sulfate accumulates within lysosomes. Researchers believe that the accumulation of GAGs interferes with the functions of other proteins inside the lysosomes and disrupts the normal functions of cells. It is unknown why the buildup of heparan sulfate mostly affects the central nervous system in MPS IIIA.

Chromosomal Location

Cytogenetic Location: 17q25.3, which is the long (q) arm of chromosome 17 at position 25.3

Molecular Location: base pairs 80,200,668 to 80,220,333 on chromosome 17 (Homo sapiens Updated Annotation Release 109.20190905, GRCh38.p13) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- heparan N-sulfatase
- heparan sulfate sulfatase
- HSS
- N-sulphoglucosamine sulphohydrolase
- N-sulphoglucosamine sulphohydrolase precursor
- SFMD
- SPHM_HUMAN
- sulfamidase
- sulfoglucosamine sulfamidase
- sulphamidase

Additional Information & Resources

Educational Resources

- Eureka Bioscience Collection: Defects in Glycosaminoglycan Degradation (Mucopolysaccharidoses)
<https://www.ncbi.nlm.nih.gov/books/NBK6177/#A53462>

Clinical Information from GeneReviews

- Mucopolysaccharidosis Type III
<https://www.ncbi.nlm.nih.gov/books/NBK546574>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28SGSH%5BTIAB%5D%29+OR+%28N-sulfo-glucosamine+sulfohydrolase%5BTIAB%5D%29%29+OR+%28%28heparan+sulfate+sulfatase%5BTIAB%5D%29+OR+%28MPS3A%5BTIAB%5D%29+OR+%28N-sulpho-glucosamine+sulphohydrolase%5BTIAB%5D%29+OR+%28sulphamidase%5BTIAB%5D%29+OR+%28heparan+N-sulfatase%5BTIAB%5D%29+OR+%28sulfamidase%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- N-SULFOGLUCOSAMINE SULFOHYDROLASE
<http://omim.org/entry/605270>

Research Resources

- ClinVar
<https://www.ncbi.nlm.nih.gov/clinvar?term=SGSH%5Bgene%5D>
- HGNC Gene Symbol Report
https://www.genenames.org/data/gene-symbol-report/#!/hgnc_id/HGNC:10818
- Monarch Initiative
<https://monarchinitiative.org/gene/NCBIGene:6448>
- NCBI Gene
<https://www.ncbi.nlm.nih.gov/gene/6448>
- UniProt
<https://www.uniprot.org/uniprot/P51688>

Sources for This Summary

- Meyer A, Kossow K, Gal A, Steglich C, Mühlhausen C, Ullrich K, Braulke T, Muschol N. The mutation p.Ser298Pro in the sulphamidase gene (SGSH) is associated with a slowly progressive clinical phenotype in mucopolysaccharidosis type IIIA (Sanfilippo A syndrome). *Hum Mutat.* 2008 May;29(5):770. doi: 10.1002/humu.20738.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/18407553>
- Muschol N, Storch S, Ballhausen D, Beesley C, Westermann JC, Gal A, Ullrich K, Hopwood JJ, Winchester B, Braulke T. Transport, enzymatic activity, and stability of mutant sulfamidase (SGSH) identified in patients with mucopolysaccharidosis type III A. *Hum Mutat.* 2004 Jun;23(6):559-66.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/15146460>
- OMIM: N-SULFOGLUCOSAMINE SULFOHYDROLASE
<http://omim.org/entry/605270>
- Valstar MJ, Ruijter GJ, van Diggelen OP, Poorthuis BJ, Wijburg FA. Sanfilippo syndrome: a mini-review. *J Inherit Metab Dis.* 2008 Apr;31(2):240-52. doi: 10.1007/s10545-008-0838-5. Epub 2008 Apr 4. Review.
Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/18392742>

Reprinted from Genetics Home Reference:
<https://ghr.nlm.nih.gov/gene/SGSH>

Reviewed: August 2010
Published: November 12, 2019

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