

# ALDH5A1 gene

aldehyde dehydrogenase 5 family member A1

#### **Normal Function**

The *ALDH5A1* gene provides instructions for producing the succinic semialdehyde dehydrogenase enzyme. This enzyme is found in the energy-producing centers of cells (mitochondria). Succinic semialdehyde dehydrogenase is involved in the breakdown of a chemical that transmits signals in the brain (neurotransmitter) called gamma-amino butyric acid (GABA). The primary role of GABA is to prevent the brain from being overloaded with too many signals. Once GABA molecules have been released from nerve cells, they are broken down by succinic semialdehyde dehydrogenase and other enzymes.

# **Health Conditions Related to Genetic Changes**

Succinic semialdehyde dehydrogenase deficiency

Variants (also called mutations) in the *ALDH5A1* gene have been found to cause succinic semialdehyde dehydrogenase deficiency. This is a disorder that can cause a variety of neurological problems. Most of the variants that cause this condition lead to the production of an enzyme with little or no activity. A lack of functional succinic semialdehyde dehydrogenase disrupts the conversion of succinic semialdehyde to succinic acid. Instead, succinic semialdehyde is converted back into GABA or a related molecule, gamma-hydroxybutyrate (GHB). It is unclear how increases in GHB and GABA cause developmental delays, seizures, and other features of succinic semialdehyde dehydrogenase deficiency.

Research has shown that people with *ALDH5A1* gene variants that result in a lack of functional succinic semialdehyde dehydrogenase tend to have more severe signs and symptoms than people who have gene variants that impair protein activity.

#### Other Names for This Gene

- aldehyde dehydrogenase 5 family, member A1
- aldehyde dehydrogenase 5 family, member A1 (succinate-semialdehyde dehydrogenase)
- aldehyde dehydrogenase 5A1

- mitochondrial succinate semialdehyde dehydrogenase
- NAD(+)-dependent succinic semialdehyde dehydrogenase
- SSADH
- SSDH
- SSDH\_HUMAN

#### Additional Information & Resources

# Tests Listed in the Genetic Testing Registry

Tests of ALDH5A1 (https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=7915[geneid])

## Scientific Articles on PubMed

 PubMed (https://pubmed.ncbi.nlm.nih.gov/?term=%28ALDH5A1%5BTIAB%5D%29 +OR+%28%28SSDH%5BTIAB%5D%29+OR+%28SSADH%5BTIAB%5D%29%29+ AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D %29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+2880 +days%22%5Bdp%5D)

# Catalog of Genes and Diseases from OMIM

ALDEHYDE DEHYDROGENASE 5 FAMILY, MEMBER A1; ALDH5A1 (https://omim.org/entry/610045)

## Gene and Variant Databases

- NCBI Gene (https://www.ncbi.nlm.nih.gov/gene/7915)
- ClinVar (https://www.ncbi.nlm.nih.gov/clinvar?term=ALDH5A1[gene])

#### References

- Akaboshi S, Hogema BM, Novelletto A, Malaspina P, Salomons GS, Maropoulos GD, Jakobs C, Grompe M, Gibson KM. Mutational spectrum of the succinate semialdehydedehydrogenase (ALDH5A1) gene and functional analysis of 27 novel disease-causingmutations in patients with SSADH deficiency. Hum Mutat. 2003 Dec; 22(6):442-50.doi: 10.1002/humu.10288. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/14635103)
- Blasi P, Boyl PP, Ledda M, Novelletto A, Gibson KM, Jakobs C, Hogema B, Akaboshi S, Loreni F, Malaspina P. Structure of human succinic semialdehydedehydrogenase gene: identification of promoter region and alternatively processedisoforms. Mol Genet Metab. 2002 Aug;76(4):348-62. doi:10.

- 1016/s1096-7192(02)00105-1. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/1 2208142)
- Gordon N. Succinic semialdehyde dehydrogenase deficiency (SSADH)(4-hydroxybutyric aciduria, gamma-hydroxybutyric aciduria). Eur J PaediatrNeurol. 2004;8(5):261-5. doi: 10.1016/j.ejpn.2004.06.004. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/15341910)
- Pearl PL, Gibson KM, Acosta MT, Vezina LG, Theodore WH, Rogawski MA, NovotnyEJ, Gropman A, Conry JA, Berry GT, Tuchman M. Clinical spectrum of succinicsemialdehyde dehydrogenase deficiency. Neurology. 2003 May 13;60(9): 1413-7. doi:10.1212/01.wnl.0000059549.70717.80. Citation on PubMed (https://pub.med.ncbi.nlm.nih.gov/12743223)
- Pearl PL, Novotny EJ, Acosta MT, Jakobs C, Gibson KM. Succinic semialdehydedehydrogenase deficiency in children and adults. Ann Neurol. 2003;54 Suppl6:S73-80. doi: 10.1002/ana.10629. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/12891657)
- Tokatly Latzer I, Roullet JB, Cesaro S, DiBacco ML, Arning E, Rotenberg A, LeeHHC, Opladen T, Jeltsch K, Garcia-Cazorla A, Julia-Palacios N, Gibson KM, Bertoldi M, Pearl PL. Phenotypic correlates of structural and functional proteinimpairments resultant from ALDH5A1 variants. Hum Genet. 2023Dec;142(12): 1755-1776. doi: 10.1007/s00439-023-02613-6. Epub 2023 Nov 14. Citation on PubMed (https://www.ncbi.nlm.nih.gov/pubmed/37962671)

## **Genomic Location**

The *ALDH5A1* gene is found on chromosome 6 (https://medlineplus.gov/genetics/chromosome/6/).

Last updated March 6, 2024