

DAO Antibody (Center)

Affinity Purified Rabbit Polyclonal Antibody (Pab) Catalog # AP13941c

Specification

DAO Antibody (Center) - Product Information

Application WB,E **Primary Accession** P14920 NP 001908.3 Other Accession Reactivity Human Host **Rabbit** Clonality **Polyclonal** Isotype Rabbit IgG Calculated MW 39474 Antigen Region 259-287

DAO Antibody (Center) - Additional Information

Gene ID 1610

Other Names

D-amino-acid oxidase, DAAO, DAMOX, DAO, DAO, DAMOX

Target/Specificity

This DAO antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 259-287 amino acids from the Central region of human DAO.

Dilution

WB~~1:1000

E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

DAO Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures.

DAO Antibody (Center) - Protein Information

Name DAO

Synonyms DAMOX



Function Catalyzes the oxidative deamination of D-amino acids with broad substrate specificity

 $\begin{array}{l} (\text{PubMed:} \underline{16616139}, \text{PubMed:} \underline{17088322}, \text{PubMed:} \underline{17303072}, \text{PubMed:} \underline{18544534}, \\ \text{PubMed:} \underline{20368421}, \text{PubMed:} \underline{20567862}, \text{PubMed:} \underline{20603179}, \text{PubMed:} \underline{22203986}, \\ \text{PubMed:} \underline{23219954}, \text{PubMed:} \underline{23391306}, \text{PubMed:} \underline{25030849}, \text{PubMed:} \underline{25701391}, \\ \text{PubMed:} \underline{29274788}, \text{PubMed:} \underline{29326945}, \text{PubMed:} \underline{30938755}, \text{PubMed:} \underline{31799256}, \\ \text{PubMed:} \underline{32730563}, \text{PubMed:} \underline{33484270}, \text{PubMed:} \underline{34041270}, \text{PubMed:} \underline{37558109}, \\ \end{array}$

PubMed:38035964). Required to catabolize D-amino acids synthesized endogenously, of gastrointestinal bacterial origin or obtained from the diet, and to use these as nutrients (By similarity). Regulates the level of D-amino acid neurotransmitters in the brain, such as D-serine, a co-agonist of N- methyl D-aspartate (NMDA) receptors, and may modulate synaptic transmission (PubMed:17303072). Catalyzes the first step of the racemization of D-DOPA to L-DOPA, for possible use in an alternative dopamine biosynthesis pathway (PubMed:17303072). Also catalyzes the first step of the chiral inversion of N(gamma)-nitro-D-arginine methyl ester (D-NNA) to its L-enantiomer L-NNA that acts as a nitric oxide synthase inhibitor (By similarity). The hydrogen peroxide produced in the reaction provides protection against microbial infection; it contributes to the oxidative killing activity of phagocytic leukocytes and protects against bacterial colonization of the small intestine (By similarity). Enzyme secreted into the lumen of the intestine may not be catalytically active and could instead be proteolytically cleaved into peptides with antimicrobial activity (By similarity). The hydrogen peroxide produced in the reaction may also play a role in promoting cellular senescence in response to DNA damage (PubMed:30659069). Could act as a detoxifying agent which removes D-amino acids accumulated during aging (PubMed:17303072).

Cellular Location

Peroxisome matrix. Cytoplasm, cytosol. Presynaptic active zone {ECO:0000250|UniProtKB:O35078}. Secreted Note=Transiently present in the cytosol before being delivered to the peroxisomes (PubMed:21679769, PubMed:31799256). In the cerebellum, a fraction of protein localizes to the presynaptic active zone, where its activity is regulated by protein BSN (By similarity). Secreted into the lumen of the small intestine (PubMed:27670111) {ECO:0000250|UniProtKB:O35078, ECO:0000269|PubMed:21679769, ECO:0000269|PubMed:27670111, ECO:0000269|PubMed:31799256}

Tissue Location

Expressed in the cerebellum, in astrocytes of the cortex, in motor neurons and fibers of the lumbar spinal cord (at protein level) (PubMed:17880399, PubMed:18544534, PubMed:18560437, PubMed:24138986, PubMed:34041270). Expressed in goblet cells of the small intestine (at protein level) (PubMed:27670111). Increased in the cerebellum of schizophrenic patients (at protein level) (PubMed:17880399, PubMed:18560437). Decreased in motor neurons of the spinal cord of patients with amyotrophic lateral sclerosis (at protein level) (PubMed:24138986). Expressed in the cerebellum, spinal cord, kidney, and thalamus (PubMed:17880399). Abundant in glia of the cerebellum and predominantly neuronal in the dorsolateral prefrontal cortex, hippocampus and substantia nigra (PubMed:17880399)

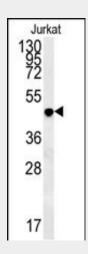
DAO Antibody (Center) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- <u>Immunofluorescen</u>ce
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

DAO Antibody (Center) - Images





DAO Antibody (Center) (Cat. #AP13941c) western blot analysis in Jurkat cell line lysates (35ug/lane). This demonstrates the DAO antibody detected the DAO protein (arrow).

DAO Antibody (Center) - Background

This gene encodes the peroxisomal enzyme D-amino acid oxidase. The enzyme is a flavoprotein which uses flavin adenine dinucleotide (FAD) as its prosthetic group. Its substrates include a wide variety of D-amino acids, but it is inactive on the naturally occurring L-amino acids. Its biological function is not known; it may act as a detoxifying agent which removes D-amino acids that accumulate during aging. In mice, it degrades D-serine, a co-agonist of the NMDA receptor. This gene may play a role in the pathophysiology of schizophrenia.

DAO Antibody (Center) - References

Kim, B., et al. Psychiatry Res 179(2):121-125(2010)
Caldinelli, L., et al. Protein Sci. 19(8):1500-1512(2010)
Ruano, G., et al. Pharmacogenomics 11(7):959-971(2010)
Ohnuma, T., et al. Schizophr. Res. 118 (1-3), 300-302 (2010):
Mitchell, J., et al. Proc. Natl. Acad. Sci. U.S.A. 107(16):7556-7561(2010)