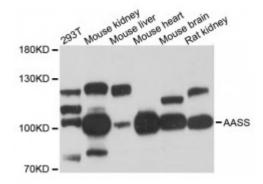


Anti-AASS Antibody





Description This gene encodes a bifunctional enzyme that catalyzes the first two steps

in the mammalian lysine degradation pathway. The N-terminal and the C-terminal portions of this enzyme contain lysine-ketoglutarate reductase and saccharopine dehydrogenase activity, respectively, resulting in the conversion of lysine to alpha-aminoadipic semialdehyde. Mutations in this

gene are associated with familial hyperlysinemia.

Model STJ114729

Host Rabbit

Reactivity Human, Mouse, Rat

Applications WB

Immunogen Recombinant fusion protein containing a sequence corresponding to amino

acids 180-400 of human AASS (NP_005754.2).

Gene ID 10157

Gene Symbol AASS

Dilution range WB 1:500 - 1:2000

Tissue Specificity Expressed in all 16 tissues examined with highest expression in the liver

Purification Affinity purification

Note For Research Use Only (RUO).

Protein Name Alpha-aminoadipic semialdehyde synthase mitochondrial LKR/SDH

Molecular Weight 102.132 kDa

Clonality Polyclonal

Unconjugated Conjugation

IgG **Isotype**

PBS with 0.02% sodium azide, 50% glycerol, pH7.3. Formulation

Store at -20C. Avoid freeze / thaw cycles. **Storage Instruction**

Database Links HGNC:17366OMIM:238700Reactome:R-HSA-71064

Alpha-aminoadipic semialdehyde synthase mitochondrial LKR/SDH **Alternative Names**

Function Bifunctional enzyme that catalyzes the first two steps in lysine degradation,

The N-terminal and the C-terminal contain lysine-ketoglutarate reductase and

saccharopine dehydrogenase activity, respectively

Cellular Localization Mitochondrion

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