

Argininosuccinate Lyase antibody

Catalog No: #22612



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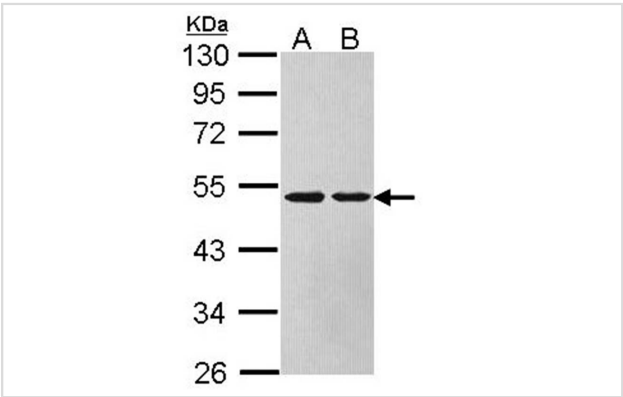
Description

|                       |  |
|-----------------------|--|
| Product Name          | Argininosuccinate Lyase antibody   |
| Host Species          | Rabbit   |
| Clonality             | Polyclonal   |
| Purification          | Purified by antigen-affinity chromatography.   |
| Applications          | WB IHC IF  |
| Species Reactivity    | Hu   |
| Immunogen Type        | Recombinant protein  |
| Immunogen Description | Recombinant protein fragment contain a sequence corresponding to a region within amino acids 13 and 231 of Argininosuccinate Lyase |
| Target Name           | Argininosuccinate Lyase  |
| Accession No.         | Swiss-Prot:P04424Gene ID:435   |
| Uniprot               | P04424   |
| GeneID                | 435;   |
| Concentration         | 1mg/ml   |
| Formulation           | Supplied in 0.1M Tris-buffered saline with 20% Glycerol (pH7.0). 0.01% Thimerosal was added as a preservative.                     |
| Storage               | Store at -20°C for long term preservation (recommended). Store at 4°C for short term use.  |

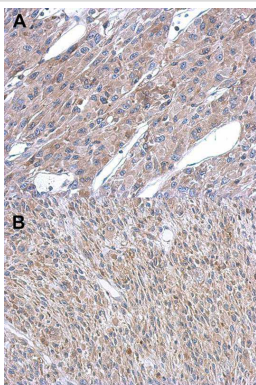
Application Details

Predicted MW: 52kd  
Western blotting: 1:500-1:3000  
Immunohistochemistry: 1:100-1:500  
Immunofluorescence: 1:100-1:200

Images

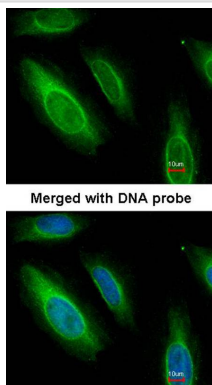


Sample (30 ug of whole cell lysate)  
A: HeLa  
B: Hep G2  
10% SDS PAGE  
Primary antibody diluted at 1: 1000



A:Immunohistochemical analysis of paraffin-embedded U87 xenograft, using ASL antibody at 1: 500 dilution.

B:Immunohistochemical analysis of paraffin-embedded C2C12 xenograft, using ASL antibody at 1: 500 dilution.



Immunofluorescence analysis of paraformaldehyde-fixed HeLa, using Argininosuccinate Lyase antibody at 1: 200 dilution.

## Background

This gene encodes a member of the lyase 1 family. The encoded protein forms a cytosolic homotetramer and primarily catalyzes the reversible hydrolytic cleavage of argininosuccinate into arginine and fumarate, an essential step in the liver in detoxifying ammonia via the urea cycle. Mutations in this gene result in the autosomal recessive disorder argininosuccinic aciduria, or argininosuccinic acid lyase deficiency. A nontranscribed pseudogene is also located on the long arm of chromosome 22. Alternatively spliced transcript variants encoding different isoforms have been described. [provided by RefSeq]

Note: This product is for in vitro research use only