# **DLDH/DLD Rabbit mAb**

Catalog No.: A5220 Recombinant 2 Publications



# **Basic Information**

### **Observed MW**

60kDa

### **Calculated MW**

54kDa

### Category

Primary antibody

### **Applications**

ELISA, WB, IHC-P, IF/ICC

### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC1224

# **Background**

This gene encodes a member of the class-I pyridine nucleotide-disulfide oxidoreductase family. The encoded protein has been identified as a moonlighting protein based on its ability to perform mechanistically distinct functions. In homodimeric form, the encoded protein functions as a dehydrogenase and is found in several multi-enzyme complexes that regulate energy metabolism. However, as a monomer, this protein can function as a protease. Mutations in this gene have been identified in patients with E3-deficient maple syrup urine disease and lipoamide dehydrogenase deficiency. Alternative splicing results in multiple transcript variants.

# **Recommended Dilutions**

WB	1:500 - 1:1000	
IHC-P	1:50 - 1:200	
IF/ICC	1:50 - 1:200	

# **Immunogen Information**

Gene ID	Swiss Prot
1738	P09622

### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 401-500 of human DLDH/DLD (P09622).

## Synonyms

E3; LAD; DLDD; DLDH; GCSL; PHE3; OGDC-E3; DLDH/DLD

# **Contact**

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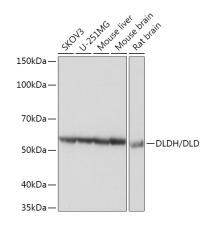
## **Product Information**

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of extracts of various cell lines, using DLDH/DLD Rabbit mAb (A5220) at 1:3000 dilution

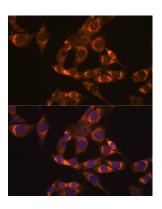
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

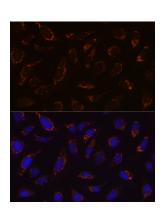
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

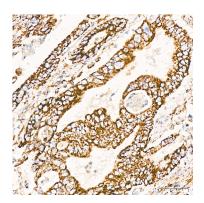
Exposure time: 1s.



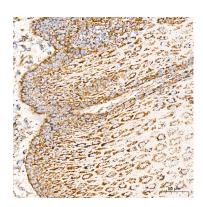
Immunofluorescence analysis of NIH-3T3 cells using DLDH/DLD Rabbit mAb (A5220) at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



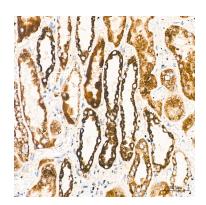
Immunofluorescence analysis of U-2 OS cells using DLDH/DLD Rabbit mAb (A5220) at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



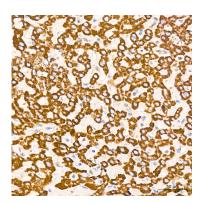
Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded human colon carcinoma tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



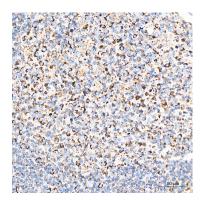
Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded human esophagus tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



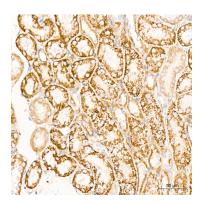
Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded human kidney tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



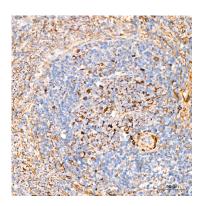
Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded human liver tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded human tonsil tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded mouse kidney tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens).High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of DLDH/DLD in paraffin-embedded rat spleen tissue using DLDH/DLD Rabbit mAb (A5220) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.