

## **Product datasheet for TA503391**

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Lipoamide Dehydrogenase (DLD) Mouse Monoclonal Antibody [Clone ID: OTI5G7]

**Product data:** 

**Product Type:** Primary Antibodies

Clone Name: OTI5G7
Applications: FC, IF, WB

Recommended Dilution: WB 1:500, IF 1:100, FLOW 1:100

Reactivity: Human, Dog, Rat, Monkey, Mouse

Host: Mouse Isotype: IgG1

Clonality: Monoclonal

Immunogen: Full length human recombinant protein of human DLD(NP\_000099) produced in HEK293T

cell

**Formulation:** PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.

**Concentration:** 0.98 mg/ml

**Purification:** Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

Storage: Store at -20°C as received.

**Stability:** Stable for 12 months from date of receipt.

**Predicted Protein Size:** 50.1 kDa

**Gene Name:** dihydrolipoamide dehydrogenase

Database Link: NP 000099

Entrez Gene 13382 MouseEntrez Gene 298942 RatEntrez Gene 403978 DogEntrez Gene

700494 MonkeyEntrez Gene 1738 Human

P09622





**Background:** This gene encodes the L protein of the mitochondrial glycine cleavage system. The L protein,

also named dihydrolipoamide dehydrogenase, is also a component of the pyruvate dehydrogenase complex, the alpha-ketoglutarate dehydrogenase complex, and the branched-chain alpha-keto acide dehydrogenase complex. Mutations in this gene have been

identified in patients with E3-deficient maple syrup urine disease and lipoamide

dehydrogenase deficiency. [provided by RefSeq, Jul 2008]

Synonyms: DLDD; DLDH; E3; GCSL; LAD; PHE3

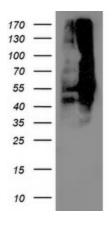
**Protein Families:** Druggable Genome

Protein Pathways: Citrate cycle (TCA cycle), Glycine, serine and threonine metabolism, Glycolysis /

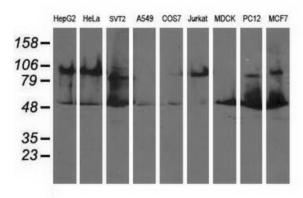
Gluconeogenesis, Metabolic pathways, Pyruvate metabolism, Valine, leucine and isoleucine

degradation

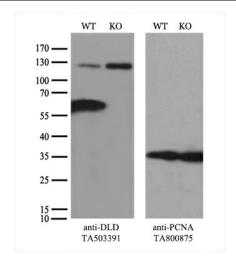
## **Product images:**



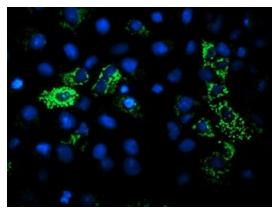
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY DLD ([RC200639], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-DLD. Positive lysates [LY400041] (100ug) and [LC400041] (20ug) can be purchased separately from OriGene.



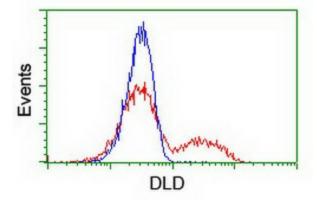
Western blot analysis of extracts (35ug) from 9 different cell lines by using anti-DLD monoclonal antibody (HepG2: human; HeLa: human; SVT2: mouse; A549: human; COS7: monkey; Jurkat: human; MDCK: canine; PC12: rat; MCF7: human).



Equivalent amounts of cell lysates (10 ug per lane) of wild-type HeLa cells (WT, Cat# LC810HELA) and DLD-Knockout HeLa cells (KO, Cat# [LC832718]) were separated by SDS-PAGE and immunoblotted with anti-DLD monoclonal antibody TA503391 (1:500). Then the blotted membrane was stripped and reprobed with anti-PCNA antibody as a loading control.



Anti-DLD mouse monoclonal antibody (TA503391) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY DLD ([RC200639]).



HEK293T cells transfected with either [RC200639] overexpress plasmid (Red) or empty vector control plasmid (Blue) were immunostained by anti-DLD antibody (TA503391), and then analyzed by flow cytometry.