

### Product datasheet for TA503421BM

#### 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 (HRP conjugated) [Clone ID: OTI6G6]

#### **Product data:**

**Product Type:** Primary Antibodies

Clone Name: OTI6G6
Applications: FC, IF, WB

Recommended Dilution: WB 1:2000, IF 1:100, FLOW 1:100

Reactivity: Human, Mouse, Rat

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.

**Concentration:** 0.5 mg/ml

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

(protein A/G)

Conjugation: HRP

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 1738 Human

P09622





# Lipoamide Dehydrogenase (DLD) Mouse Monoclonal Antibody (HRP conjugated) [Clone ID: OTI6G6] – TA503421BM

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]

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

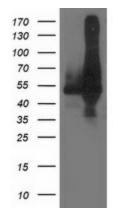
**Protein Families:** Druggable Genome

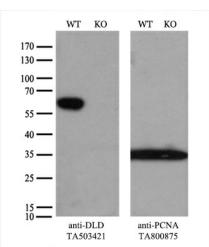
Protein Pathways: Citrate cycle (TCA cycle), Glycine, serine and threonine metabolism, Glycolysis /

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

degradation

## **Product images:**

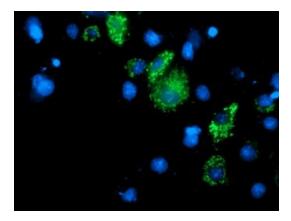




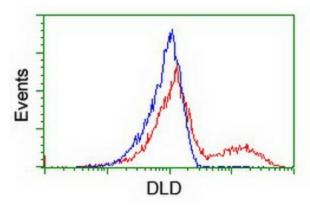
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY DLD ([RC200639], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-DLD. Positive lysates [LY400041] (100ug) and [LC400041]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

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 [TA503421] (1:500). Then the blotted membrane was stripped and reprobed with anti-PCNA antibody as a loading control.





Anti-DLD mouse monoclonal antibody ([TA503421]) 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 ([TA503421]), and then analyzed by flow cytometry.