

#### OriGene Technologies, Inc.

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# Product datasheet for TA344335

### Lipoamide Dehydrogenase (DLD) Rabbit Polyclonal Antibody

### **Product data:**

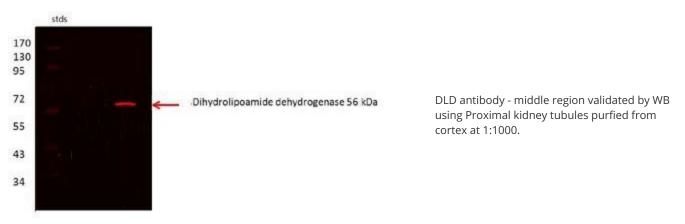
Product Type:	Primary Antibodies
Applications:	IHC, WB
Recommended Dilution:	WB
Reactivity:	Human, Rat
Host:	Rabbit
lsotype:	IgG
Clonality:	Polyclonal
Immunogen:	The immunogen for anti-DLD antibody: synthetic peptide directed towards the middle region of human DLD. Synthetic peptide located within the following region: AGEMVNEAALALEYGASCEDIARVCHAHPTLSEAFREANLAASFGKSINF
Formulation:	Liquid. Purified antibody supplied in 1x PBS buffer with 0.09% (w/v) sodium azide and 2% sucrose. Note that this product is shipped as lyophilized powder to China customers.
Purification:	Affinity Purified
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	56 kDa
Gene Name:	dihydrolipoamide dehydrogenase
Database Link:	<u>NP_000099</u> <u>Entrez Gene 298942 RatEntrez Gene 1738 Human</u> <u>P09622</u>

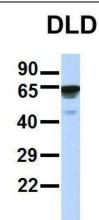


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	Lipoamide Dehydrogenase (DLD) Rabbit Polyclonal Antibody – TA344335
Background:	DLD is the L protein of the mitochondrial glycine cleavage system. The L protein, also named dihydrolipoamide dehydrogenase, is also a component of the pyruvate dehydrogenase 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. 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, 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. Publication Note: This RefSeq record includes a subset of the publications that are available for this gene. Please see the Entrez Gene record to access additional publications.
Synonyms:	DLDD; DLDH; E3; GCSL; LAD; PHE3
Note:	lmmunogen Sequence Homology: Dog: 100%; Horse: 100%; Human: 100%; Bovine: 100%; Rabbit: 100%; Pig: 93%; Rat: 93%; Zebrafish: 93%; Guinea pig: 93%; Mouse: 86%; Yeast: 86%
Protein Families	: Druggable Genome
Protein Pathway	<b>ys:</b> Citrate cycle (TCA cycle), Glycine, serine and threonine metabolism, Glycolysis / Gluconeogenesis, Metabolic pathways, Pyruvate metabolism, Valine, leucine and isoleucine degradation

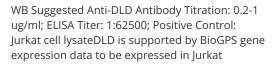
## **Product images:**

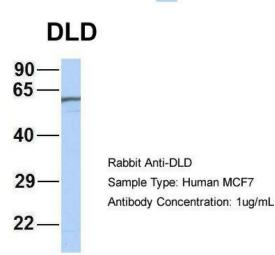


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Rabbit Anti-DLD Sample Type: Human Adult Placenta Antibody Concentration: 1ug/mL

2Hum. Adult Placenta; Host: Rabbit; Target Name: SERPINA3; Sample Tissue: Human Adult Placenta; Antibody Dilution: 1.0 ug/ml





90 kDa\_ 65 kDa

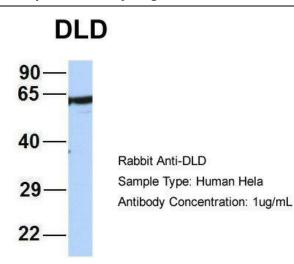
40 kDa

29 kDa\_

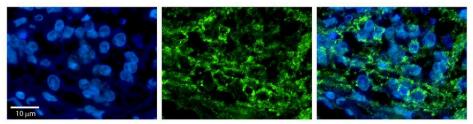
22 kDa

4Human MCF7; Host: Rabbit; Target Name: NOP56; Sample Tissue: MCF7; Antibody Dilution: 1.0 ug/ml; DLD is supported by BioGPS gene expression data to be expressed in MCF7

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5Human Hela; Host: Rabbit; Target Name: EGFL8; Sample Tissue: Hela; Antibody Dilution: 1.0 ug/ml; DLD is supported by BioGPS gene expression data to be expressed in HeLa



Rabbit Anti-DLD Antibody; Formalin Fixed Paraffin Embedded Tissue: Human Pineal Tissue; Observed Staining: Cytoplasmic in cell bodies and processes of pinealocytes; Primary Antibody Concentration: 1:100; Secondary Antibody: Donkey anti-Rabbit-Cy3; Seconda

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