



# Rabbit Anti-Human DLD Polyclonal Antibody (CABT-L2119)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Product Overview</b>	Polyclonal Antibody to Dihydrolipoyl Dehydrogenase (Knockout Validated)
<b>Specificity</b>	The antibody is a rabbit polyclonal antibody raised against DLD. It has been selected for its ability to recognize DLD in immunohistochemical staining and western blotting.
<b>Target</b>	DLD
<b>Immunogen</b>	Recombinant fragment corresponding to human DLD (Gly280~His487)
<b>Isotype</b>	IgG
<b>Source/Host</b>	Rabbit
<b>Species Reactivity</b>	Human, Mouse
<b>Purification</b>	Antigen-specific affinity chromatography followed by Protein A affinity chromatography
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB
<b>Format</b>	Liquid
<b>Concentration</b>	Lot specific
<b>Size</b>	200 µg
<b>Buffer</b>	Supplied as solution form in 0.01M PBS with 50% glycerol, pH7.4.
<b>Preservative</b>	0.05% Proclin-300

<b>Storage</b>	Avoid repeated freeze/thaw cycles. Store at 4°C for frequent use. Aliquot and store at -20°C for 12 months.
<b>Ship</b>	4°C with ice bags

## BACKGROUND

<b>Introduction</b>	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. [provided by RefSeq, Jan 2014]
<b>Keywords</b>	LAD;GCSL;Dihydrolipoamide Dehydrogenase;E3 Component Of Pyruvate Dehydrogenase Complex;2-Oxo-Glutarate Complex;Branched Chain Keto Acid Dehydrogenase Complex

## GENE INFORMATION

<b>Gene Name</b>	DLD dihydrolipoamide dehydrogenase [ Homo sapiens (human) ]
<b>Official Symbol</b>	DLD
<b>Synonyms</b>	DLD; dihydrolipoamide dehydrogenase; E3; LAD; DLDD; DLDH; GCSL; PHE3; dihydrolipoyl dehydrogenase, mitochondrial; diaphorase; lipoamide reductase; lipoyl dehydrogenase; lipoamide dehydrogenase; glycine cleavage system L protein; glycine cleavage system protein L; E3 component of pyruvate dehydrogenase complex, 2-oxo-glutarate complex, branched chain keto acid dehydrogenase complex;
<b>Entrez Gene ID</b>	<a href="#">1738</a>
<b>Protein Refseq</b>	NP_000099
<b>UniProt ID</b>	<a href="#">A0A024R713</a>
<b>Chromosome Location</b>	7q31-q32
<b>Pathway</b>	2-oxobutanoate degradation; 2-oxobutanoate degradation I; 2-oxoglutarate decarboxylation to succinyl-CoA; 2-oxoisovalerate decarboxylation to isobutanoyl-CoA; Branched-chain amino acid catabolism; Carbon metabolism; Citrate cycle (TCA cycle); Citrate cycle (TCA cycle, Krebs cycle);

**Function**

NADP binding; dihydrolipoyl dehydrogenase activity; flavin adenine dinucleotide binding;  
mercury (II) reductase activity; mercury ion binding;

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