

Recombinant

DGRmAb®

DMGDH (DGR32797) Rabbit mAb

db13362

Package : 10μL 20μL 50μL 100μL

Product Name : DMGDH (DGR32797) Rabbit mAb**Cat.No.:** db13362**Synonyms :** DMGDHD; ME2GLYDH**Application :** WB**Reactivity :** Human,Mouse,Rat**Host species :** Rabbit**Background**

This gene encodes an enzyme involved in the catabolism of choline, catalyzing the oxidative demethylation of dimethylglycine to form sarcosine. The enzyme is found as a monomer in the mitochondrial matrix, and uses flavin adenine dinucleotide and folate as cofactors. Mutation in this gene causes dimethylglycine dehydrogenase deficiency, characterized by a fishlike body odor, chronic muscle fatigue, and elevated levels of the muscle form of creatine kinase in serum. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jul 2013]

Immunogen

A synthetic peptide of human DMGDH

Gene ID

29958

Swiss Prot

Q9UI17

Synonyms

DMGDHD; ME2GLYDH

Reactivity

Human,Mouse,Rat

Application

WB

Recommended dilution

WB: 1:1000

Calculated MW

97 kDa

Observed MW

97 kDa

Host species

Rabbit

Clonality

Monoclonal

Clonality No.

DGR32797

Isotype

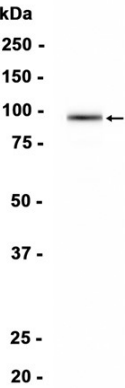
IgG

Purity

Affinity Purification

Conjugation	Un-conjugated
Storage Stability	Store at -20°C. Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium azide and 0.05% BSA. Stable for 12 months from date of receipt.

Human fetal liver



Western blot analysis of extracts from Human fetal liver tissue using db13362 at 1:1000.