

**HMGCL Antibody (N-term)**  
**Affinity Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP18139a**

**Specification**

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**HMGCL Antibody (N-term) - Product Information**

|                   |                             |
|-------------------|-----------------------------|
| Application       | WB,E                        |
| Primary Accession | <a href="#">P35914</a>      |
| Other Accession   | <a href="#">NP_000182.2</a> |
| Reactivity        | Human                       |
| Host              | Rabbit                      |
| Clonality         | Polyclonal                  |
| Isotype           | Rabbit IgG                  |
| Calculated MW     | 34360                       |
| Antigen Region    | 71-99                       |

**HMGCL Antibody (N-term) - Additional Information**

**Gene ID** 3155

**Other Names**

Hydroxymethylglutaryl-CoA lyase, mitochondrial, HL, HMG-CoA lyase,  
3-hydroxy-3-methylglutarate-CoA lyase, HMGCL

**Target/Specificity**

This HMGCL antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 71-99 amino acids from the N-terminal region of human HMGCL.

**Dilution**

WB~~1:1000

**Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

**Storage**

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

HMGCL Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

**HMGCL Antibody (N-term) - Protein Information**

**Name** HMGCL

**Function** Mitochondrial 3-hydroxymethyl-3-methylglutaryl-CoA lyase that catalyzes a

cation-dependent cleavage of (S)-3-hydroxy-3-methylglutaryl-CoA into acetyl-CoA and acetoacetate, a key step in ketogenesis. Terminal step in leucine catabolism. Ketone bodies (beta-hydroxybutyrate, acetoacetate and acetone) are essential as an alternative source of energy to glucose, as lipid precursors and as regulators of metabolism.

#### Cellular Location

Mitochondrion matrix {ECO:0000250|UniProtKB:P38060}. Peroxisome {ECO:0000250|UniProtKB:P38060}. Note=Unprocessed form is peroxisomal {ECO:0000250|UniProtKB:P38060}

#### Tissue Location

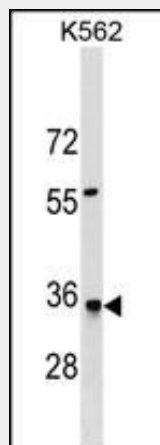
Highest expression in liver. Expressed in pancreas, kidney, intestine, testis, fibroblasts and lymphoblasts. Very low expression in brain and skeletal muscle. The relative expression of isoform 2 (at mRNA level) is highest in heart (30%), skeletal muscle (22%), and brain (14%).

### HMGCL Antibody (N-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

### HMGCL Antibody (N-term) - Images



HMGCL Antibody (N-term) (Cat. #AP18139a) western blot analysis in K562 cell line lysates (35ug/lane). This demonstrates the HMGCL antibody detected the HMGCL protein (arrow).

### HMGCL Antibody (N-term) - Background

The protein encoded by this gene belongs to the HMG-CoA lyase family. It is a mitochondrial enzyme that catalyzes the final step of leucine degradation and plays a key role in ketone body formation. Mutations in this gene are associated with HMG-CoA lyase deficiency. Alternatively spliced transcript variants encoding

different isoforms have been found for this gene. [provided by RefSeq].

#### **HMGCL Antibody (N-term) - References**

Fu, Z., et al. J. Biol. Chem. 285(34):26341-26349(2010)  
Pierron, S., et al. Arch Pediatr 17(1):10-13(2010)  
Menao, S., et al. Hum. Mutat. 30 (3), E520-E529 (2009) :  
Lin, W.D., et al. Clin. Chim. Acta 401 (1-2), 33-36 (2009) :  
Carrasco, P., et al. Mol. Genet. Metab. 91(2):120-127(2007)