



## Anti-GCSH (full length) polyclonal antibody (DPAB-DC1311)

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

### PRODUCT INFORMATION

<b>Antigen Description</b>	Degradation of glycine is brought about by the glycine cleavage system, which is composed of four mitochondrial protein components: P protein (a pyridoxal phosphate-dependent glycine decarboxylase), H protein (a lipoic acid-containing protein), T protein (a tetrahydrofolate-requiring enzyme), and L protein (a lipoamide dehydrogenase). The protein encoded by this gene is the H protein, which transfers the methylamine group of glycine from the P protein to the T protein. Defects in this gene are a cause of nonketotic hyperglycinemia (NKH). Two transcript variants, one protein-coding and the other probably not protein-coding, have been found for this gene. Also, several transcribed and non-transcribed pseudogenes of this gene exist throughout the genome.[provided by RefSeq, Jan 2010]
<b>Immunogen</b>	GCSH (AAH00790.1, 1 a.a. ~ 173 a.a) full-length recombinant protein with GST tag. The sequence is  MALRVVRSVRALLCTLRAVPLAAPCPPRWQLGVGAVRTLRTGPALLSVRKFTEKHEWV TTENGIGTVGISNFAQEALGDVYCSLPEVGTKLNKQDEFGALESVKAASELYSPLSGEV TEINEALAENPGLVNKSCYEDGWLIKMTLSNPSELDELMSEEAYEKYIKSIEE
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB (Recombinant protein), ELISA,
<b>Size</b>	50 µl
<b>Buffer</b>	50 % glycerol
<b>Preservative</b>	None

**Storage**

Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

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## GENE INFORMATION

<b>Gene Name</b>	<a href="#">GCSH glycine cleavage system protein H (aminomethyl carrier) [ Homo sapiens (human) ]</a>
<b>Official Symbol</b>	GCSH
<b>Synonyms</b>	GCSH; glycine cleavage system protein H (aminomethyl carrier); GCE; NKH; glycine cleavage system H protein, mitochondrial; lipoic acid-containing protein; mitochondrial glycine cleavage system H-protein;
<b>Entrez Gene ID</b>	<a href="#">2653</a>
<b>Protein Refseq</b>	<a href="#">NP_004474</a>
<b>UniProt ID</b>	<a href="#">P23434</a>
<b>Chromosome Location</b>	16q23.2
<b>Pathway</b>	Glycine, serine and threonine metabolism; Glyoxylate and dicarboxylate metabolism.
<b>Function</b>	aminomethyltransferase activity; enzyme binding;

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