

## **Product datasheet for TP317886**

## OriGene Technologies, Inc.

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## GCSH (NM 004483) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human glycine cleavage system protein H (aminomethyl carrier)

(GCSH), 20 µg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC217886 representing NM\_004483

or AA Sequence: Red=Cloning site Green=Tags(s)

MALRVVRSVRALLCTLRAVPLPAAPCPPRPWQLGVGAVRTLRTGPALLSVRKFTEKHEWVTTENGIGTVG ISNFAQEALGDVVYCSLPEVGTKLNKQDEFGALESVKAASELYSPLSGEVTEINEALAENPGLVNKSCYE

**DGWLIKMTLSNPSELDELMSEEAYEKYIKSIEE** 

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 13.8 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

**RefSeq:** NP 004474

**Locus ID:** 2653

UniProt ID: P23434





RefSeq Size: 1161

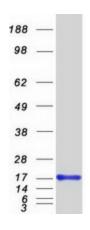
Cytogenetics: 16q23.2 RefSeq ORF: 519

Synonyms: GCE; NKH

**Summary:** 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]

## **Product images:**



Coomassie blue staining of purified GCSH protein (Cat# TP317886). The protein was produced from HEK293T cells transfected with GCSH cDNA clone (Cat# [RC217886]) using MegaTran 2.0 (Cat# [TT210002]).