abcam

Product datasheet

Recombinant Human GGT1/GGT protein ab190382

Description

Product name Recombinant Human GGT1/GGT protein

Purity > 90 % SDS-PAGE.

The final product was refolded using temperature shift inclusion body refolding technology and

chromatographically purified.

Expression system Escherichia coli

Accession P19440

Protein length Protein fragment

Animal free No

Nature Recombinant

Species Human

Sequence MASMTGGQQMGRGHHHHHHGNLYFQGGESASKEPDNH

VYTRAAVAADAKQ

CSKIGRDALRDGGSAVDAAIAALLCVGLMNAHSMGIGGGL

FLTIYNSTTR

KAEVINAREVAPRLAFATMFNSSEQSQKGGLSVAVPGEIR

GYELAHQRHG

RLPWARLFQPSIQLARQGFPVGKGLAAALENKRTVIEQQP

VLCEVFCRDR

KVLREGERLTLPQLADTYETLAIEGAQAFYNGSLTAQIVKDI

QAAGGIVT

AEDLNNYRAELIEHPLNISLGDVVLYMPSAPLSGPVLALILNI

LKGYNFS

RESVESPEQKGLTYHRIVEAFRFAYAKRTLLGDPKFVDVT

EVVRNMTSEF

FAAQLRAQISDDTTHPISYYKPEFYTPDDGGTAHLSVVAED

GSAVSATST

INLYFGSKVRSPVSGILFNNEMDDFSSPSITNEFGVPPSPA

NFIQPGKQP

LSSMCPTIMVGQDGQVRMVVGAAGGTQITTATALAIIYNLW

FGYDVKRAV

EEPRLHNQLLPNVTTVERNIDQAVTAALETRHHHTQIASTFI

AVVQANR TAGGWAAASDSRKGGEPAGY

Predicted molecular weight 62 kDa

Amino acids 28 to 569

1

Tags His-T7 tag N-Terminus

Additional sequence information NP_038347. Constructed with codon optimization and expressed with a small T7-His-TEV

cleavage site Tag (29aa) fusion at N terminal.

Specifications

Our Abpromise guarantee covers the use of ab190382 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Functional Studies

SDS-PAGE

Form Liquid

Additional notes This product was previously labelled as GGT1

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 2% Tris HCl, 1% Potassium chloride, 0.01% DTT, 0.01% EDTA, 10% Glycerol,

10% Sodium chloride

General Info

Function Initiates extracellular glutathione (GSH) breakdown, provides cells with a local cysteine supply and

contributes to maintain intracelular GSH level. It is part of the cell antioxidant defense mechanism.

Catalyzes the transfer of the glutamyl moiety of glutathione to amino acids and dipeptide acceptors. Alternatively, glutathione can be hydrolyzed to give Cys-Gly and gamma glutamate.

Isoform 3 seems to be inactive.

Tissue specificity Detected in fetal and adult kidney and liver, adult pancreas, stomach, intestine, placenta and lung.

Isoform 3 is lung-specific. There are several other tissue-specific forms that arise from alternative

promoter usage but that produce the same protein.

Pathway Sulfur metabolism; glutathione metabolism.

Involvement in diseaseDefects in GGT1 are a cause of glutathionuria (GLUTH) [MIM:231950]; also known as gamma-

glutamyltranspeptidase deficiency. It is an autosomal recessive disease.

Sequence similarities Belongs to the gamma-glutamyltransferase family.

Post-translational N-glycosylated on both chains. Contains hexoses, hexosamines and sialic acid residues.

modifications Glycosylation profiles tested in kidney and liver tissues reveal the presence of tissue-specific and

site-specific glycan composition, despite the overlap in composition among the N-glycans. A total of 36 glycan compositions, with 40 unique structures are observed. Up to 15 different glycans are observed at a single site, with site-specific variation in glycan composition. The difference in

glycosylation profiles in the 2 tissues do not affect the enzyme activity.

Cellular localization Membrane.

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