

GST (3D4): sc-57753

BACKGROUND

Plasmid vectors for the expression of coding regions of eukaryotic genes in *E. coli* are in common usage; such expression vectors often encode hybrid fusion proteins containing part prokaryotic and part eukaryotic specified proteins. For instance, the pGEX.3X expression vector developed by Smith and Johnson allows for synthesis of fusion proteins between glutathione-S-transferase (GST) and proteins encoded by inserted cDNA sequences. Antibodies derived from these GST fusion proteins are useful for checking protein expression both in plaques and on Western blots as well as for immunoaffinity purification of proteins expressed in *E. coli*.

REFERENCES

- Maniatis, T., et al. 1982. Molecular Cloning. Cold Spring Harbor, NY: Cold Spring Harbor Laboratory.
- Smith, D.B. and Johnson, K.S. 1988. Single-step purification of polypeptides expressed in *Escherichia coli* as fusions with glutathione S-transferase. *Gene* 67: 31-40.
- Crabb, B.S. and Studdert, M.J. 1995. Expression of small regions of equine herpesvirus 1 glycoprotein C in *Escherichia coli*. *Vet. Microbiol.* 46: 181-191.
- Soler, D., et al. 1995. Matrilysin: expression, purification and characterization. *J. Protein Chem.* 14: 511-520.
- Yu, L., et al. 1995. Cloning, gene sequencing and expression of the small molecular mass ubiquinone-binding protein of mitochondrial biquinol-cytochrome c reductase. *J. Biol. Chem.* 270: 25634-25638.
- Driscoll, J., et al. 1995. Functional comparison of native and recombinant human salivary histatin 1. *J. Dent. Res.* 74: 1837-1844.
- Chen, Y.R., et al. 1996. Functional expression of subunit IV of *Rhodospira sphaeroides* cytochrome b-c1 complex and reconstitution of recombinant protein with three-subunit core complex. *J. Biol. Chem.* 271: 2057-2062.

SOURCE

GST (3D4) is a mouse monoclonal antibody raised against GST of *Schistosoma japonicum* origin.

PRODUCT

Each vial contains 100 µg IgG₁ in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

GST (3D4) is recommended for detection of GST fusion proteins and glutathione-S-transferase (GST) of *Schistosoma japonicum* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)] and immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

Molecular Weight of GST: 26 kDa.

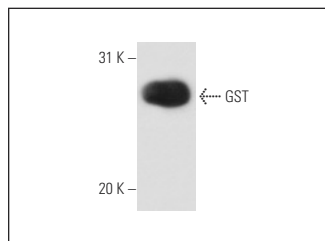
RESEARCH USE

For research use only, not for use in diagnostic procedures.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA



GST (3D4): sc-57753. Western blot analysis of recombinant GST protein.

SELECT PRODUCT CITATIONS

- Dennis, P.B. and Mercer, C.A. 2009. The GST-BHMT assay and related assays for autophagy. *Methods Enzymol.* 452: 97-118.
- Knauer, S.K., et al. 2011. Bioassays to monitor Taspase1 function for the identification of pharmacogenetic inhibitors. *PLoS ONE* 6: e18253.
- Ranneberg-Nilsen, T., et al. 2012. The chromatin remodeling factor SMARCB1 forms a complex with human cytomegalovirus proteins UL114 and UL44. *PLoS ONE* 7: e34119.
- Wünsch, D., et al. 2015. Fly versus man: evolutionary impairment of nucleolar targeting affects the degradome of *Drosophila's* Taspase1. *FASEB J.* 29: 1973-1985.
- Wünsch, D., et al. 2015. Evolutionary divergence of Threonine Aspartase1 leads to species-specific substrate recognition. *Biol. Chem.* 396: 367-376.
- Rui, Y.N., et al. 2015. The GST-BHMT assay reveals a distinct mechanism underlying proteasome inhibition-induced macroautophagy in mammalian cells. *Autophagy* 11: 812-832.
- Cho, E.B., et al. 2018. β -dystroglycan is regulated by a balance between WWP1-mediated degradation and protection from WWP1 by dystrophin and utrophin. *Biochim. Biophys. Acta* 1864: 2199-2213.
- Cramer, M., et al. 2018. MxB is an interferon-induced restriction factor of human herpesviruses. *Nat. Commun.* 9: 1980.
- , A., et al. 2021. Specific inhibition of the survivin-CRM1 interaction by peptide-modified molecular tweezers. *Nat. Commun.* 12: 1505.
- Kim, D.E., et al. 2023. PLK1-mediated phosphorylation of β -catenin enhances its stability and transcriptional activity for extracellular matrix remodeling in metastatic NSCLC. *Theranostics* 13: 1198-1216.

CONJUGATES

See **GST (B-14): sc-138** for GST antibody conjugates, including AC, HRP, FITC, PE, and Alexa Fluor® 488, 546, 594, 647, 680 and 790.