

**GSTM5 Antibody (N-term) Blocking peptide**  
Synthetic peptide  
Catalog # BP12728a

## Specification

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### GSTM5 Antibody (N-term) Blocking peptide - Product Information

Primary Accession [P46439](#)

### GSTM5 Antibody (N-term) Blocking peptide - Additional Information

Gene ID 2949

#### Other Names

Glutathione S-transferase Mu 5, GST class-mu 5, GSTM5-5, GSTM5

#### Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

#### Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

#### Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

### GSTM5 Antibody (N-term) Blocking peptide - Protein Information

Name GSTM5

#### Function

Conjugation of reduced glutathione to a wide number of exogenous and endogenous hydrophobic electrophiles.

#### Cellular Location

Cytoplasm.

### GSTM5 Antibody (N-term) Blocking peptide - Protocols

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

- [Blocking Peptides](#)

### GSTM5 Antibody (N-term) Blocking peptide - Images

### GSTM5 Antibody (N-term) Blocking peptide - Background

Cytosolic and membrane-bound forms of glutathioneS-transferase are encoded by two distinct

supergene families. At present, eight distinct classes of the soluble cytoplasmic mammalian glutathione S-transferases have been identified: alpha, kappa, mu, omega, pi, sigma, theta and zeta. This gene encodes a glutathione S-transferase that belongs to the mu class. The mu class of enzymes functions in the detoxification of electrophilic compounds, including carcinogens, therapeutic drugs, environmental toxins and products of oxidative stress, by conjugation with glutathione. The genes encoding the mu class of enzymes are organized in a gene cluster on chromosome 1p13.3 and are known to be highly polymorphic. These genetic variations can change an individual's susceptibility to carcinogens and toxins as well as affect the toxicity and efficacy of certain drugs. Diversification of these genes has occurred in regions encoding substrate-binding domains, as well as in tissue expression patterns, to accommodate an increasing number of foreign compounds.

### **GSTM5 Antibody (N-term) Blocking peptide - References**

Wang, Y., et al. J. Hum. Genet. 55(8):490-494(2010) Yu, K.D., et al. Breast Cancer Res. Treat. 121(2):485-496(2010) Davila, S., et al. Genes Immun. 11(3):232-238(2010) Moyer, A.M., et al. Cancer Epidemiol. Biomarkers Prev. 19(3):811-821(2010) Saito, A., et al. J. Hum. Genet. 54(6):317-323(2009)