

## BACKGROUND

Glutathione S-transferases (GSTs) are a large and diverse family of enzymes, and in humans, there are at least 13 GST enzymes belonging to five families, namely alpha (GSTA), mu (GSTM), pi (GSTP), theta (GSTS), and tao (GSTT). GSTs play an important role in detoxification by catalyzing the conjugation of many hydrophobic and electrophilic compounds with reduced glutathione.

Among this family of isozymes, GSTP1 is generally the most prevalent in mammalian cells. The link between GST overexpression, especially with respect to GSTP1, and anticancer drug resistance has been extensively studied. Because of the defined role of GST in drug metabolism, elevated expression of GSTP1 in solid tumors or in drug-resistant cells has been associated frequently with detoxification reactions even in instances where there is no evidence that the selecting drug is a substrate for GSTP1. More recently, however, the link between the redox active components of GSTP1 and stress-activated kinases, such as JNK, has been redefined as a non-catalytic, ligand binding activity that mediates both stress and apoptotic responses. In a parallel series of studies, either GST $\mu$  or thioredoxin has also been identified as a ligand-binding partner for apoptosis-signaling kinase (ASK1), extending the role that "redox proteins" may have in kinase regulation. It was demonstrated that GSTP1 has significant affinity for the C terminus of JNK and confirms the ligand-binding regulatory role for this ubiquitously expressed protein.<sup>1</sup> In addition; GSTP1 activity can be regulated by interacting with other proteins. The Fanconi anemia group C protein (FANCC) interacts with GSTP1 and prevents the formation of inactivating disulfide bonds within GSTP1 and increases GSTP1 activity during apoptosis.<sup>2</sup>

GSTP1 is not only involved in cancers, but also in other diseases. GSTP1 gene polymorphisms affect substrate selectivity and stability, and the oxidative milieu in dopaminergic neurons, which increases the susceptibility to Parkinson's disease. In addition, it is possible that intermediate electrophilic metabolites, arising in the first phase of detoxification, are not metabolized by GST enzymes in asthmatic patients and are not excreted. These intermediate metabolites may damage cells and generate oxidative stress, and so contribute to the pathogenesis of asthma.<sup>3</sup>

### References:

1. Wang, T. et al: J. Biol. Chem. 276:20999-21003, 2001
2. Cummings, R.C. et al: Nature Med. 7:814-20, 2001
3. Tamer, L. et al: Respiration 9:493-8, 2004

## TECHNICAL INFORMATION

### Source:

GSTP1 antibody is a mouse monoclonal antibody raised against purified recombinant human GSTP1 proteins expressed in *E. coli*.

### Specificity and Sensitivity:

This antibody detects endogenous GSTP1 proteins without cross-reactivity with other family members.

**Storage Buffer:** PBS and 30% glycerol

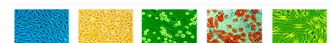
### Storage:

Store at -20°C for at least one year. Store at 4°C for frequent use. Avoid repeated freeze-thaw cycles.

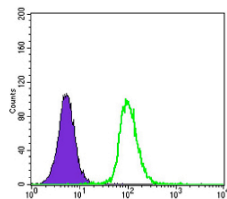
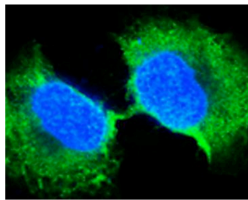
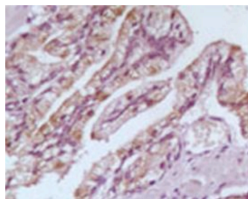
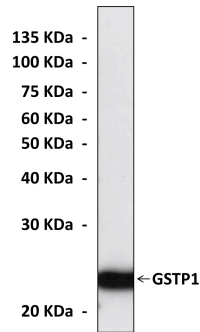
## APPLICATIONS

Application:	*Dilution:
WB	1:1000
IP	n/d
IHC	1:200
ICC	1:200
FACS	1:200

*\*Optimal dilutions must be determined by end user.*



## QUALITY CONTROL DATA



**Top:** Western Blot detection of GSTP1 proteins in PC-3 cell lysate using GSTP1 Antibody. **Middle, upper:** This antibody stains paraffin-embedded human prostate cancer tissue in immunohistochemical analysis. **Middle, lower:** It also stains HepG2 cells in confocal immunofluorescent testing (GSTP1 Antibody: Green; Actin filament: Red; DRAQ5 DNA dye: Blue). **Bottom:** This antibody specifically reacts with GSTP1 protein in K562 cells vs. negative control mouse IgG (Blue) in FACS testing.

