

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product datasheet for TA501883S

Glutathione S Transferase theta 2 (GSTT2) Mouse Monoclonal Antibody [Clone ID: OTI6C4]

Product data:

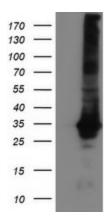
Product Type:	Primary Antibodies
Clone Name:	OTI6C4
Applications:	IF, IHC, WB
Recommended Dilution:	WB 1:200~500, IHC 1:150, IF 1:100
Reactivity:	Human
Host:	Mouse
lsotype:	lgG2b
Clonality:	Monoclonal
Immunogen:	Full length human recombinant protein of human GSTT2(NP_000845) produced in HEK293T cell.
Formulation:	PBS (pH 7.3) containing 1% BSA, 50% glycerol and 0.02% sodium azide.
Concentration:	0.62 mg/ml
Purification:	Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography (protein A/G)
Conjugation:	Unconjugated
Storage:	Store at -20°C as received.
Stability:	Stable for 12 months from date of receipt.
Predicted Protein Size:	27.3 kDa
Gene Name:	glutathione S-transferase theta 2 (gene/pseudogene)
Database Link:	<u>NP_000845</u> Entrez Gene 2953 Human
	POCG29



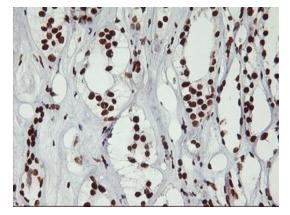
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	Glutathione S Transferase theta 2 (GSTT2) Mouse Monoclonal Antibody [Clone ID: OTI6C4] – TA501883S
Background:	Glutathione S-transferase (GSTs) theta 2 (GSTT2) is a member of a superfamily of proteins that catalyze the conjugation of reduced glutathione to a variety of electrophilic and hydrophobic compounds. Human GSTs can be divided into five main classes: Alpha, Mu, Pi, Theta, and Zeta. The theta class members GSTT1 and GSTT2 share 55% amino acid sequence identity and both are thought to have an important role in human carcinogenesis. The theta genes have a similar structure, being composed of five exons with identical exon/intron boundaries. [provided by RefSeq]
Synonyms:	GSTT2B
Protein Pathwa	ys: Drug metabolism - cytochrome P450, Glutathione metabolism, Metabolism of xenobiotics by cytochrome P450

Product images:



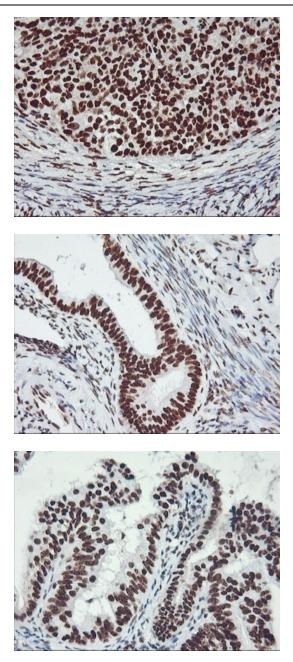
HEK293T cells were transfected with the pCMV6-ENTRY control (Left lane) or pCMV6-ENTRY GSTT2 ([RC200040], Right lane) cDNA for 48 hrs and lysed. Equivalent amounts of cell lysates (5 ug per lane) were separated by SDS-PAGE and immunoblotted with anti-GSTT2. Positive lysates [LY424485] (100ug) and [LC424485] (20ug) can be purchased separately from OriGene.



Immunohistochemical staining of paraffinembedded Human Kidney tissue within the normal limits using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

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Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human ovary tissue using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

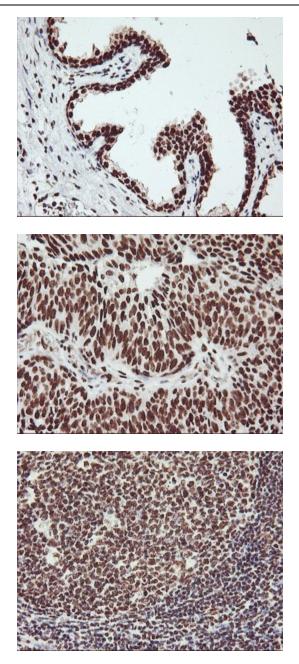
Immunohistochemical staining of paraffinembedded Human endometrium tissue within the normal limits using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

Immunohistochemical staining of paraffinembedded Adenocarcinoma of Human endometrium tissue using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

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Glutathione S Transferase theta 2 (GSTT2) Mouse Monoclonal Antibody [Clone ID: OTI6C4] – TA501883S



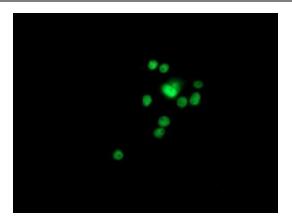
Immunohistochemical staining of paraffinembedded Human prostate tissue within the normal limits using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

Immunohistochemical staining of paraffinembedded Carcinoma of Human bladder tissue using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

Immunohistochemical staining of paraffinembedded Human tonsil within the normal limits using anti-GSTT2 mouse monoclonal antibody. (Heat-induced epitope retrieval by 10mM citric buffer, pH6.0, 100°C for 10min, [TA501883])

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Anti-GSTT2 mouse monoclonal antibody ([TA501883]) immunofluorescent staining of COS7 cells transiently transfected by pCMV6-ENTRY GSTT2 ([RC200040]).

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