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anti-GPX5 antibody (Internal Region)

2 Images



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Overview	
Quantity:	100 μL
Target:	GPX5
Binding Specificity:	Internal Region
Reactivity:	Human, Rat, Mouse
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GPX5 antibody is un-conjugated
Application:	ELISA, Western Blotting (WB), Immunohistochemistry (IHC)
Product Details	
Immunogen:	A synthesized peptide derived from human GPX5, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	GPX5 Antibody detects endogenous levels of total GPX5.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	GPX5

Target Details

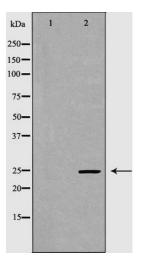
Alternative Name:	GPX5 (GPX5 Products)
Background:	Description: Protects cells and enzymes from oxidative damage, by catalyzing the reduction of
	hydrogen peroxide, lipid peroxides and organic hydroperoxide, by glutathione. May constitute a
	glutathione peroxidase-like protective system against peroxide damage in sperm membrane
	lipids.
	Gene: GPX5
Molecular Weight:	25kDa
Gene ID:	2880
UniProt:	075715
Pathways:	Thyroid Hormone Synthesis

Application Details

Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

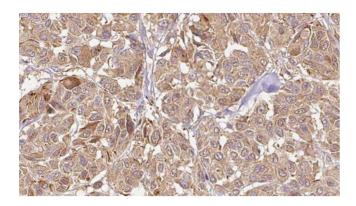
Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



Western Blotting

Image 1. Western blot analysis of testis using GPX5 antibody. The lane on the left is treated with the antigenspecific peptide.



Immunohistochemistry

Image 2. ABIN6276489 at 1/100 staining Human Melanoma tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22¡ãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary