## PTEN(Ab-380/382/383) Antibody

Catalog No: #21056

Package Size: #21056-1 50ul #21056-2 100ul



Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

Description	
Product Name	PTEN(Ab-380/382/383) Antibody
Host Species	Rabbit
Clonality	Polyclonal
Purification	Antibodies were produced by immunizing rabbits with synthetic peptide and KLH conjugates. Antibodies were
	purified by affinity-chromatography using epitope-specific peptide.
Applications	WB IHC
Species Reactivity	Hu Ms Rt
Specificity	The antibody detects endogenous level of total PTEN protein.
Immunogen Type	Peptide-KLH
Immunogen Description	Peptide sequence around aa.378~382/380~384/381~385 (R-Y-S-D-T-T-D-S) derived from Human PTEN.
Target Name	PTEN
Other Names	MMAC1; Mutated in multiple advanced cancers 1; Protein-tyrosine phosphatase PTEN; TEP1;
Accession No.	Swiss-Prot: P60484NCBI Protein: NP_000305.3

P60484

5728;

1.0mg/ml

sodium azide and 50% glycerol.

Application Details			
Predicted MW: 54kd			
Western blotting: 1:500~1:1000			
Immunohistochemistry: 1:50~1:1	00		

## Images

Uniprot GenelD

Concentration

Formulation

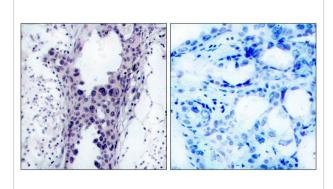
Storage

кд   НТ29   1 1 8 - 85 —	
47 PTEN	
36-	
26- Peptide - +	
47 PTEN 36- 26-	

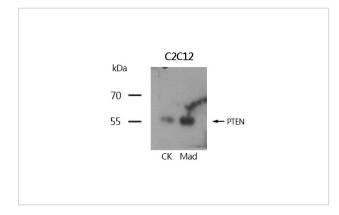
Western blot analysis of extracts from HT29 cells using PTEN(Ab-380/382/383) Antibody #21056 and the same antibody preincubated with blocking peptide.

Supplied at 1.0mg/mL in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl, 0.02%

Store at -20°C for long term preservation (recommended). Store at 4°C for short term use.



Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue using PTEN(Ab-380/382/383) Antibody #21056(left) or the same antibody preincubated with blocking peptide(right).



Western blotting analysis using PTEN(Ab-380/382/383) Antibody #21056.

## Background

Tumor suppressor. Acts as a dual-specificity protein phosphatase, dephosphorylating tyrosine-, serine- and threonine-phosphorylated proteins. Also acts as a lipid phosphatase, removing the phosphate in the D3 position of the inositol ring from phosphatidylinositol 3,4,5-trisphosphate, phosphatidylinositol 3,4-diphosphate, phosphatidylinositol 3-phosphate and inositol 1,3,4,5-tetrakisphosphate with order of substrate preference in vitro PtdIns(3,4,5)P3 > PtdIns(3,4)P2 > PtdIns3P > Ins(1,3,4,5)P4. The lipid phosphatase activity is critical for its tumor suppressor function. Antagonizes the PI3K-AKT/PKB signaling pathway by dephosphorylating phosphoinositides and thereby modulating cell cycle progression and cell survival. The unphosphorylated form cooperates with AIP1 to suppress AKT1 activation. Dephosphorylates tyrosine-phosphorylated focal adhesion kinase and inhibits cell migration and integrin-mediated cell spreading and focal adhesion formation. May be a negative regulator of insulin signaling and glucose metabolism in adipose tissue.

Al-Khouri AM, et al. (2005) J Biol Chem. 280(42):35195-35202.

Torres J, et al. (2001) J Biol Chem. 276(2): 993-998.

Vazquez F, et al. (2000) Mol Cell Biol. 20(14): 5010-5018.

Note: This product is for in vitro research use only