

## SH-PTP2 rabbit pAb

## Cat No.:ES3441

For research use only

## Overview

Product Name	SH-PTP2 rabbit pAb	
Host species	Rabbit	
Applications	WB;IHC;IF;ELISA	
Species Cross-Reactivity	Human;Mouse;Rat;Monkey	
Recommended dilutions	Western Blot: 1/500 - 1/2000.	
	Immunohistochemistry: 1/100 - 1/300. ELISA:	
	1/40000. Not yet tested in other applications.	
Immunogen	The antiserum was produced against synthesized	
	peptide derived from human SH-PTP2. AA	
	range:321-370	
Specificity	SH-PTP2 Polyclonal Antibody detects endogenous	
	levels of SH-PTP2 protein.	
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and	
	0.02% sodium azide.	
Storage	Store at -20 $^\circ\!\!\mathbb{C}$ . Avoid repeated freeze-thaw cycles.	
Protein Name	Tyrosine-protein phosphatase non-receptor type 11	
Gene Name	PTPN11	
Cellular localization	Cytoplasm . Nucleus .	
Purification	The antibody was affinity-purified from rabbit	
	antiserum by affinity-chromatography using	
	epitope-specific immunogen.	
Clonality	Polyclonal	
Concentration	1 mg/ml	
Observed band	68kD	
Human Gene ID	5781	
Human Swiss-Prot Number	Q06124	
Alternative Names	PTPN11; PTP2C; SHPTP2; Tyrosine-protein	
	phosphatase non-receptor type 11; Protein-tyrosine	
	phosphatase 1D; PTP-1D; Protein-tyrosine	
	phosphatase 2C; PTP-2C; SH-PTP2; SHP-2; Shp2;	
	SH-PTP3	
Background	The protein encoded by this gene is a member of	
	the protein tyrosine phosphatase (PTP) family. PTPs	



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are known to be signaling molecules that regulate a variety of cellular processes including cell growth, differentiation, mitotic cycle, and oncogenic transformation. This PTP contains two tandem Src homology-2 domains, which function as phospho-tyrosine binding domains and mediate the interaction of this PTP with its substrates. This PTP is widely expressed in most tissues and plays a regulatory role in various cell signaling events that are important for a diversity of cell functions, such as mitogenic activation, metabolic control, transcription regulation, and cell migration. Mutations in this gene are a cause of Noonan syndrome as well as acute myeloid leukemia. [provided by RefSeq, Aug 2016],



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