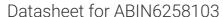
# antibodies .- online.com





### anti-PLK3 antibody (Internal Region)

3 Images

Overview



Go to Product page

Quantity:	100 μL
Target:	PLK3
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This PLK3 antibody is un-conjugated

## Application: Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF)

#### **Product Details**

Immunogen:	A synthesized peptide derived from human PLK3, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	PLK3 Antibody detects endogenous levels of total PLK3.
Predicted Reactivity:	Pig,Zebrafish,Bovine,Horse,Sheep,Rabbit,Dog,Chicken,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

#### **Target Details**

Target:	PLK3		
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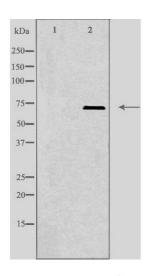
#### **Target Details**

Alternative Name:	PLK3 (PLK3 Products)	
Background:	Description: Serine/threonine-protein kinase involved in cell cycle regulation, response to stress	
	and Golgi disassembly. Polo-like kinases act by binding and phosphorylating proteins are that	
	already phosphorylated on a specific motif recognized by the POLO box domains.	
	Phosphorylates ATF2, BCL2L1, CDC25A, CDC25C, CHEK2, HIF1A, JUN, p53/TP53, p73/TP73,	
	PTEN, TOP2A and VRK1. Involved in cell cycle regulation: required for entry into S phase and	
	cytokinesis. Phosphorylates BCL2L1, leading to regulate the G2 checkpoint and progression to	
	cytokinesis during mitosis. Plays a key role in response to stress: rapidly activated upon stress	
	stimulation, such as ionizing radiation, reactive oxygen species (ROS), hyperosmotic stress, UV	
	irradiation and hypoxia. Involved in DNA damage response and G1/S transition checkpoint by	
	phosphorylating CDC25A, p53/TP53 and p73/TP73. Phosphorylates p53/TP53 in response to	
	reactive oxygen species (ROS), thereby promoting p53/TP53-mediated apoptosis.	
	Phosphorylates CHEK2 in response to DNA damage, promoting the G2/M transition	
	checkpoint. Phosphorylates the transcription factor p73/TP73 in response to DNA damage,	
	leading to inhibit p73/TP73-mediated transcriptional activation and pro-apoptotic functions.	
	Phosphorylates HIF1A and JUN is response to hypoxia. Phosphorylates ATF2 following	
	hyperosmotic stress in corneal epithelium. Also involved in Golgi disassembly during the cell	
	cycle: part of a MEK1/MAP2K1-dependent pathway that induces Golgi fragmentation during	
	mitosis by mediating phosphorylation of VRK1. May participate in endomitotic cell cycle, a form	
	of mitosis in which both karyokinesis and cytokinesis are interrupted and is a hallmark of	
	megakaryocyte differentiation, via its interaction with CIB1.	
	Gene: PLK3	
Molecular Weight:	70 kDa	
Gene ID:	1263	
UniProt:	Q9H4B4	
Pathways:	Regulation of long-term Neuronal Synaptic Plasticity	
Application Details		
Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000	
Restrictions:	For Research Use only	
Handling		

#### Handling

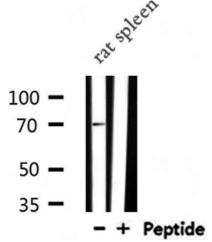
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 $\%$ sodium azide and 50 $\%$ glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.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

#### **Images**



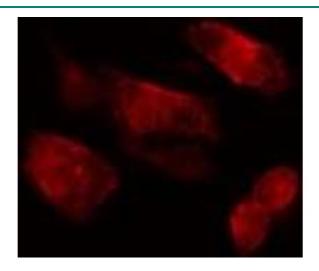
#### **Western Blotting**

**Image 1.** Western blot analysis of extracts from HepG2 cells, using PLK3 antibody. The lane on the left is treated with the antigen-specific peptide.



#### **Western Blotting**

**Image 2.** Western blot analysis of PLK3 expression in Rat spleen lysate



#### Immunofluorescence (fixed cells)

**Image 3.** ABIN6275535 staining HepG2 cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod