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anti-PPM1K antibody (Internal Region)

2 Images



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Target:

100 μL
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PPM1K
Internal Region
Human, Mouse
Rabbit
Polyclonal
This PPM1K antibody is un-conjugated
Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunocytochemistry (ICC), Immunofluorescence (IF)
A synthesized peptide derived from human PPM1K, corresponding to a region within the internal amino acids.
IgG
PPM1K Antibody detects endogenous levels of total PPM1K.
Pig,Zebrafish,Bovine,Horse,Sheep,Rabbit,Dog
The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).

PPM1K

Target Details

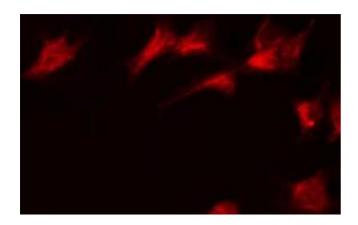
Alternative Name:	PPM1K (PPM1K Products)
Background:	Description: Regulates the mitochondrial permeability transition pore and is essential for cellular survival and development. Gene: PPM1K
Molecular Weight:	45 kDa
Gene ID:	152926
UniProt:	Q8N3J5

Application Details

Application Notes:	WB 1:500-1:1000, IHC: 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

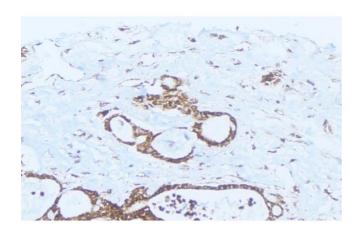
Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit lgG 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



Immunofluorescence (fixed cells)

Image 1. ABIN6275413 staining 293 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_iaC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37_iaC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod



Immunohistochemistry

Image 2. ABIN6275413 at 1/100 staining Human lung cancer 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