# antibodies -online.com







**Images** 



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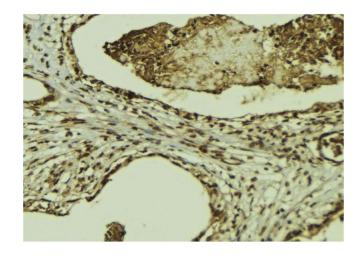
Quantity:	100 μL
Target:	XIAP
Binding Specificity:	N-Term
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This XIAP antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human XIAP, corresponding to a region within N-terminal amino acids.
Isotype:	IgG
Specificity:	XIAP Antibody detects endogenous levels of total XIAP.
Predicted Reactivity:	Pig,Horse,Sheep,Rabbit
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).
Target Details	
Target:	XIAP

Alternative Name:	XIAP (XIAP Products)
Background:	Description: Multi-functional protein which regulates not only caspases and apoptosis, but also
	modulates inflammatory signaling and immunity, copper homeostasis, mitogenic kinase
	signaling, cell proliferation, as well as cell invasion and metastasis. Acts as a direct caspase
	inhibitor. Directly bind to the active site pocket of CASP3 and CASP7 and obstructs substrate
	entry. Inactivates CASP9 by keeping it in a monomeric, inactive state. Acts as an E3 ubiquitin-
	protein ligase regulating NF-kappa-B signaling and the target proteins for its E3 ubiquitin-
	protein ligase activity include: RIPK1, CASP3, CASP7, CASP8, CASP9, MAP3K2/MEKK2,
	DIABLO/SMAC, AIFM1, CCS and BIRC5/survivin. Ubiquitinion of CCS leads to enhancement of
	its chaperone activity toward its physiologic target, SOD1, rather than proteasomal degradation
	Ubiquitinion of MAP3K2/MEKK2 and AIFM1 does not lead to proteasomal degradation. Plays a
	role in copper homeostasis by ubiquitinationg COMMD1 and promoting its proteasomal
	degradation. Can also function as E3 ubiquitin-protein ligase of the NEDD8 conjugation
	pathway, targeting effector caspases for neddylation and inactivation. Regulates the BMP
	signaling pathway and the SMAD and MAP3K7/TAK1 dependent pathways leading to NF-
	kappa-B and JNK activation. Acts as an important regulator of innate immune signaling via
	regulation of Nodlike receptors (NLRs). Protects cells from spontaneous formation of the
	ripoptosome, a large multi-protein complex that has the capability to kill cancer cells in a
	caspase-dependent and caspase-independent manner. Suppresses ripoptosome formation by
	ubiquitinating RIPK1 and CASP8. Acts as a positive regulator of Wnt signaling and ubiquitinates
	TLE1, TLE2, TLE3, TLE4 and AES. Ubiquitination of TLE3 results in inhibition of its interaction
	with TCF7L2/TCF4 thereby allowing efficient recruitment and binding of the transcriptional
	coactivator beta-catenin to TCF7L2/TCF4 that is required to initiate a Wnt-specific
	transcriptional program.
	Gene: XIAP
Molecular Weight:	57kDa
Gene ID:	331
Gene ID: UniProt:	331 P98170
UniProt:	P98170
UniProt: Pathways:	P98170

## Handling

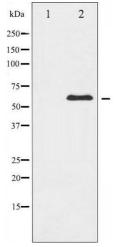
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained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

# **Images**



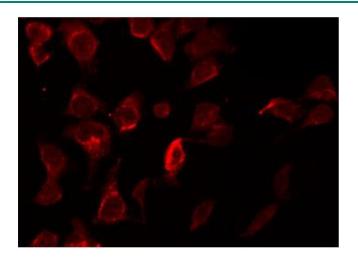
### **Immunohistochemistry**

**Image 1.** ABIN6269302 at 1/100 staining Mouse colon 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°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.



### **Western Blotting**

**Image 2.** Western blot analysis of XIAP expression in 293 whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



# Immunofluorescence (fixed cells)

**Image 3.** ABIN6269302 staining HepG2 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°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.