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anti-BIK antibody (pThr33)



Overview

# **Images**



Go to Product page

Quantity:	100 μL
Target:	BIK
Binding Specificity:	pThr33
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This BIK antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF),

Immunocytochemistry (ICC)

## **Product Details**

Immunogen:	A synthesized peptide derived from human BIK around the phosphorylation site of Thr33.
Isotype:	IgG
Specificity:	Phospho-BIK (Thr33) Antibody detects endogenous levels of BIK only when phosphorylated at Threonine 33.
Purification:	The antibody is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.

## **Target Details**

Target:	BIK
Alternative Name:	BIK (BIK Products)

## **Target Details**

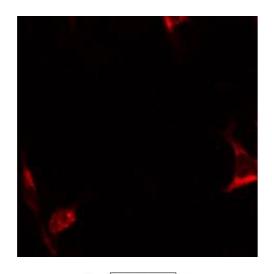
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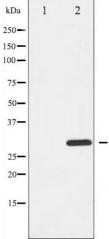
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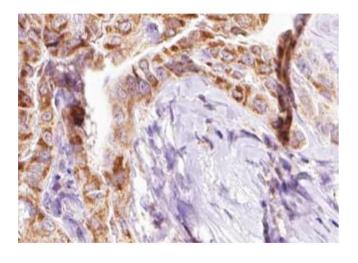
rarget Details	
Background:	Description: Accelerates programmed cell death. Association to the apoptosis repressors Bol-
	X(L), BHRF1, Bcl-2 or its adenovirus homolog E1B 19k protein suppresses this death-promoting
	activity. Does not interact with BAX.
	Gene: BIK
Molecular Weight:	30kDa
Gene ID:	638
UniProt:	Q13323
Pathways:	Positive Regulation of Endopeptidase Activity
Application Details	
Application Notes:	WB 1:500-1:2000, 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 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

Store at -20 °C. Stable for 12 months from date of receipt.

12 months







#### Immunofluorescence (fixed cells)

**Image 1.** ABIN6267637 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.

#### **Western Blotting**

**Image 2.** Western blot analysis of BIK phosphorylation expression in HepG2 whole cell lysates,The lane on the left is treated with the antigen-specific peptide.

### **Immunohistochemistry**

**Image 3.** ABIN6267637 at 1/100 staining human Thyroid cancer tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue 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.