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# anti-DOK1 antibody (pTyr362)

3 Images



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Overview
Ouantity:

Quantity:	100 μL
Target:	DOK1
Binding Specificity:	pTyr362
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This DOK1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	
Immunogen:	A synthesized peptide derived from human p62 Dok around the phosphorylation site of Tyr362.
Isotype:	IgG
Specificity:	Phospho-p62 Dok (Tyr362) Antibody detects endogenous levels of p62 Dok only when
	phosphorylated at Tyrosine 362.
Predicted Reactivity:	Pig,Bovine,Horse,Sheep,Rabbit,Dog
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:	DOK1

### **Target Details**

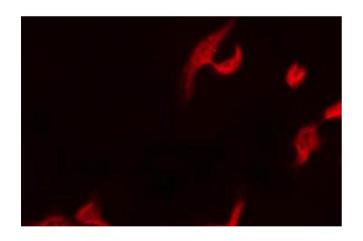
Alternative Name:	DOK1 (DOK1 Products)
Background:	Description: DOK proteins are enzymatically inert adaptor or scaffolding proteins. They provide a docking platform for the assembly of multimolecular signaling complexes. DOK1 appears to be a negative regulator of the insulin signaling pathway. Modulates integrin activation by competing with talin for the same binding site on ITGB3.  Gene: DOK1
Molecular Weight:	62kDa
Gene ID:	1796
UniProt:	Q99704

## **Application Details**

Application Notes:	WB 1:500-1:2000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

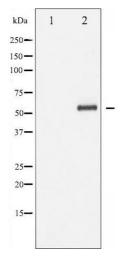
#### Handling

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



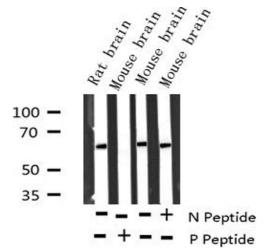
#### Immunofluorescence (fixed cells)

**Image 1.** ABIN6267687 staining Hela 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 p62 Dok phosphorylation expression in Jurkat whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



#### **Western Blotting**

**Image 3.** Western blot analysis of Phospho-p62 Dok (Tyr362) expression in various lysates