For Research Use Only

## Phospho-RIPK1 (Ser161) Monoclonal antibody

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Antibodies | ELISA kits | Proteins

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Catalog Number:66854-1-lg

8 Publications

**Basic Information** 

Catalog Number: GenBank Accession Number: Purification Method: 66854-1-lg NM\_003804 Protein A purification

Size: GeneID (NCBI): CloneNo.: 100ul , Concentration: 1500 µg/ml by 8737 1B2G1

Nanodrop;
Full Name: Recommended Dilutions:
Source: receptor (TNFRSF)-interacting serine- WB 1:2000- 1:20000
Mouse threonine kinase 1 IHC 1:250-1:1000
Isotype: Calculated MW: IF 1:200-1:800

IgG1 76 kDa
Observed MW:

Observed MW: 74 kDa, 45 kDa

**Applications** 

Tested Applications: Positive Controls: FC, IF, IHC, WB, EUSA WR: Turket collections

FC, IF, IHC, WB, ELISA

WB: Jurkat cells, Calyculin A treated HEK-293T cells,
Cited Applications:

Raji cells, THP-1 cells

IHC ; human small intestine tissue, human heart tissue

IF : U2OS cells,

Cited Species: human, mouse

Human

Species Specificity:

Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (\*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0

**Background Information** 

RIPK1(Receptor-interacting serine/threonine-protein kinase 1) is primarily involved in mediating TNF-R1-induced cell activation, apoptosis and necroptosis and belongs to a novel class of kinases thatfunction in cell survival and cell death mechanisms(PMID:22685397). It has 2 isoforms produced by alternative splicing. It also can exist as the protein with lower molecular weight about 25kd and 45kd in HAEC and HUVEC(PMID:22685397).

## **Notable Publications**

Author	Pubmed ID	Journal	Application
Ye Wang	36505813	Front Oncol	WB
Chia-Kang Ho	35322540	J Cell Mol Med	WB
Yongchang Yang	35308164	Front Genet	WB

Storage

Storage:

Store at -20°C. Stable for one year after shipment.

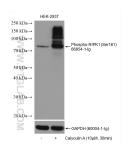
Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

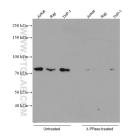
Aliquoting is unnecessary for -20°C storage

\*\*\* 20ul sizes contain 0.1% BSA

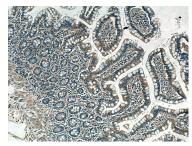
## Selected Validation Data



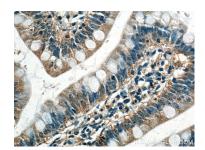
Non-treated HEK-293T and Calyculin A treated HEK-293T cells were subjected to SDS PAGE followed by western blot with 66854-1-lg (Phospho-RIPK1 (Ser161) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



Various lysates were subjected to SDS PAGE followed by western blot with 66854-1-1g (Phospho-RIPKI (Ser161) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was untreated (left) or treated with Lambda Protein Phosphatase (A-PPase, 500U, right) at 37°C for 1 hours before blocking.



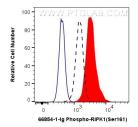
Immunohistochemical analysis of paraffinembedded human small intestine tissue slide using 66854-1-Ig (RIPK1 phospho S161 antibody) at dilution of 1:500 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunohistochemical analysis of paraffinembedded human small intestine tissue slide using 66854-1-Ig (RIPK1 phospho S161 antibody) at dilution of 1:500 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunofluorescent analysis of (-20°C Ethanol) fixed UZOS cells using Phospho-RIPK1 (Ser161) antibody (66854-1-lg, Clone: 1B2G1) at dilution of 1:400 and CoraLite®488-Conjugated Affini Pure Goat Anti-Mouse IgG(H+L).



1X10^6 Calyculin A treated HEK-293T cells were intracellularly stained with 0.13 ug Anti-Human Phospho-RIPK1 (Ser161) (66854-1-1g, Clone:1B2G1) labeled with FlexAble CoraLite® Plus 555 Antibody Labeling Kit for Mouse IgG1 (KFA022). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.