

For Research Use Only

# Phospho-MCL1 (Thr163) Polyclonal antibody



Catalog Number: 29560-1-AP

## Basic Information

Catalog Number:

29560-1-AP

Size:

150 µg/ml

Source:

Rabbit

Isotype:

IgG

GenBank Accession Number:

BC017197

GeneID (NCBI):

4170

UNIPROT ID:

Q07820

Full Name:

myeloid cell leukemia sequence 1  
(BCL2-related)

Calculated MW:

350 aa, 37 kDa

Observed MW:

40 kDa

Purification Method:

Antigen affinity purification

Recommended Dilutions:

WB 1:500-1:2000

## Applications

Tested Applications:

WB, ELISA

Species Specificity:

Human

Positive Controls:

WB : MG132 treated HeLa cells,

## Background Information

MCL1 is an anti-apoptotic member of the BCL-2 family originally isolated from the ML-1 human myeloid leukemia cell line. Similar to BCL2 and BCL2L1, MCL1 can interact with BAX and/or BAK1 to inhibit mitochondria-mediated apoptosis. Recent studies show that MCL1 is upregulated in numerous hematological and solid tumor malignancies. Therefore, MCL1 has been suggested as a potential new therapeutic target. MCL1 can be phosphorylated by several protein kinases which enables the recognition of MCL1 by its E3 ubiquitin-ligases TrCP or FBW7 (PMID: 33308268). MCL1 shows higher stability when phosphorylated on threonine 163 (PMID: 16543145).

## 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

For technical support and original validation data for this product please contact:

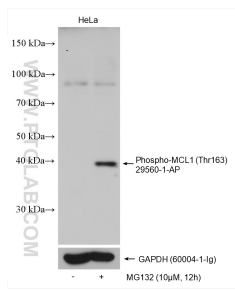
T: 4006900926

E: [Proteintech-CN@ptglab.com](mailto:Proteintech-CN@ptglab.com)

W: [ptgcn.com](http://ptgcn.com)

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

## Selected Validation Data



Non-treated and MG132 treated HeLa cells were subjected to SDS PAGE followed by western blot with 29560-1-AP (Phospho-MCL1 (Thr163) antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as the loading control.