

Product datasheet for AP15015PU-N

ZAK (MAP3K20) (Center) Rabbit Polyclonal Antibody

Product data:

OriGene Technologies, Inc.

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Product Type:	Primary Antibodies
Applications:	IHC, WB
Recommended Dilution:	ELISA: 1/1,000. Western blotting: 1/100 - 1/500. Immunohistochemistry: 1/50 - 1/100.
Reactivity:	Human
Host:	Rabbit
lsotype:	lg
Clonality:	Polyclonal
Immunogen:	This antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide selected from the center region of human MLKLAK.
Specificity:	This antibody reacts to MLKLAK.
Formulation:	PBS with 0.09% (W/V) sodium azide State: Purified State: Liquid purified Ig
Concentration:	lot specific
Purification:	Protein G column, eluted with high and low pH buffers and neutralized immediately, followed by dialysis against PBS
Conjugation:	Unconjugated
Storage:	Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.
Stability:	Shelf life: one year from despatch.
Gene Name:	mitogen-activated protein kinase kinase kinase 20
Database Link:	<u>Entrez Gene 51776 Human</u> <u>Q9NYL2</u>



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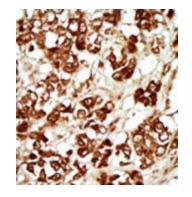
GRIGENE ZAK (MAP3K20) (Center) Rabbit Polyclonal Antibody – AP15015PU-N

Background:MLKLAK is a member of the MAPKKK family of signal transduction molecules. It possesses an
N-terminal kinase catalytic domain, followed by a leucine zipper motif and a sterile-alpha
motif (SAM). This magnesium-binding protein forms homodimers and is located in the
cytoplasm. The protein mediates gamma radiation signaling leading to cell cycle arrest and
activity of this protein plays a role in cell cycle checkpoint regulation in cells. The protein also
has pro-apoptotic activity.

Synonyms: ZAK, HCCS4, HCCS-4, MLKLAK

Product images:

Western blot analysis of anti-MLKLAK-C284 Pab in HepG2 cell line lysate (35ug/lane). MLKLAK-C284 (arrow) was detected using the purified Pab (1:60 dilution). This western blot detected isoform 2 of MLKLAK.



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by AEC staining.

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