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JNK1/2/3 (Phospho Tyr185) Rabbit pAb

CatalogNo: YP0843 Orthogonal Validated 💽

Key Features

Host Species Rabbit 	Reactivity • Human,Mouse,Rat	Applications • IF,WB,IHC,ELISA
MW • 46kD,54kD (Observed)	Isotype • IgG	

Recommended Dilution Ratios

IF 1:50-200 WB 1:500-1:2000 IHC 1:100-1:300 ELISA 1:20000 Not yet tested in other applications

Storage

Storage*-15°C to -25°C/1 year(Do not lower than -25°C)FormulationLiquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Basic Information

Clonality Polyclonal

Immunogen Information

Immunogen The antiserum was produced against synthesized peptide derived from human SAPK/JNK around the phosphorylation site of Tyr185. AA range:151-200

Specificity

Phospho-JNK1/2/3 (Y185) Polyclonal Antibody detects endogenous levels of JNK1/2/3 protein only when phosphorylated at Y185.The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification sequence (lowercase letters are modification sites):MMtPY

Target Information

Gene name	MAPK8/9/10		
Protein Name	Mitogen-activated protein kinase 8/9/10		
	Organism	Gene ID	UniProt ID
	Human	<u>5599; 5601; 5602;</u>	<u>P45983; P45984; P53779;</u>
	Mouse	<u>26419; 26420;</u>	
	Rat	116554; 50658; 25272;	P49185; P49186; P49187;

Cellular Cytoplasm . Nucleus . Cell junction, synapse . In the cortical neurons, predominantly cytoplasmic and associated with the Golgi apparatus and endosomal fraction. Increased neuronal activity increases phosphorylated form at synapses (By similarity). Colocalizes with POU5F1 in the nucleus. .

Tissue specificity Brain, Epithelium, Fetal brain, Lung, Pooled, Testis,

Function Catalytic activity: ATP + a protein = ADP + aphosphoprotein.,cofactor:Magnesium.,Domain:The TXY motif contains the threonine and tyrosine residues whose phosphorylation activates the MAP kinases., enzyme regulation: Activated by threonine and tyrosine phosphorylation by either of two dual specificity kinases, MAP2K4 and MAP2K7. Inhibited by dual specificity phosphatases, such as DUSP1., Function: JNK1 isoforms display different binding patterns: beta-1 preferentially binds to c-Jun, whereas alpha-1, alpha-2, and beta-2 have a similar low level of binding to both c-lun or ATF2. However, there is no correlation between binding and phosphorylation, which is achieved at about the same efficiency by all isoforms., Function: Responds to activation by environmental stress and pro-inflammatory cytokines by phosphorylating a number of transcription factors, primarily components of AP-1 such as JUN, JDP2 and ATF2 and thus regulates AP-1 transcriptional activity. In T-cells, INK1 and INK2 are required for polarized differentiation of T-helper cells into Th1 cells (By similarity). Phosphorylates heat shock factor protein 4 (HSF4).,online information:C-Jun N-terminal kinases entry,PTM:Dually phosphorylated on Thr-183 and Tyr-185, which activates the enzyme., similarity: Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. MAP kinase subfamily., similarity: Contains 1 protein kinase domain., subunit: Binds to at least four scaffolding proteins, MAPK8IP1/IIP-1, MAPK8IP2/IIP-2, MAPK8IP3/IIP-3/ISAP1 and SPAG9/MAPK8IP4/IIP-4. These proteins also bind other components of the INK signaling pathway. Interacts with TP53 and WWOX. Interacts with JAMP. Forms a complex with MAPK8IP1 and RGNEF (By similarity). Interacts with NFATC4.,

Validation Data





Immunofluorescence analysis of human-lung tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

Immunofluorescence analysis of human-stomach tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B



Immunohistochemical analysis of paraffin-embedded Human-uterus-cancer tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Human-colon tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Human-lung tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Human-stomach-cancer tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Rat-liver tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

only.

Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using SAPK/JNK (Phospho-Tyr185) Antibody



1.400

1.200

1.000 0.800 0.600 0.400

0.000

1.260

phosphopeptide

OD 450nm Reading

0.049

non-phosphonentide







Immunohistochemical analysis of paraffin-embedded Mouse-brain tissue.

1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody

Immunohistochemical analysis of paraffin-embedded Rat-lung tissue. 1,JNK1/2/3 (phospho Tyr185) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Western blot analysis of lysates from HepG2 cells treated with nocodazole 1ug/ml 16h, using SAPK/JNK (Phospho-Tyr185) Antibody. The lane on the right is blocked with the phospho peptide.

Contact information

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Please scan the QR code to access additional product information: JNK1/2/3 (Phospho Tyr185) Rabbit pAb

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