

JNK1/2/3 (phospho Thr183) Polyclonal Antibody

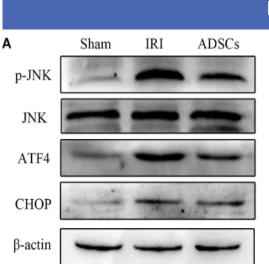
Catalog No :	YP0156
Reactivity :	Human;Mouse;Rat;Chicken
neactivity .	
Applications :	WB;IHC;IF;ELISA
Target :	JNK1/2/3
Fields :	>>Endocrine resistance;>>MAPK signaling pathway;>>ErbB signaling pathway;>>Ras signaling pathway;>>cAMP signaling pathway;>>FoxO signaling pathway;>>Sphingolipid signaling pathway;>>Mitophagy - animal;>>Autophagy - animal;>>Protein processing in endoplasmic reticulum;>>Apoptosis;>>Apoptosis - multiple species;>>Necroptosis;>>Wnt signaling pathway;>>Osteoclast differentiation;>>Focal adhesion;>>Tight junction;>>Toll-like receptor signaling pathway;>>NOD-like receptor signaling pathway;>>RIG-I-like receptor signaling pathway;>>C-type lectin receptor signaling pathway;>>IL-17 signaling pathway;>>Th1 and Th2 cell differentiation;>>Th17 cell differentiation;>>T cell receptor signaling pathway;>>Retrograde endocannabinoid signaling;>>Dopaminergic synapse;>>Inflammatory mediator regulation of TRP channels;>>Insulin signaling pathway;>>Snr
Gene Name :	MAPK8/9/10
Protein Name :	Mitogen-activated protein kinase 8/9/10
Human Gene Id :	5599/5601/5602
Human Swiss Prot	P45983/P45984/P53779
No : Mouse Gene Id :	26419/26420
Rat Gene Id :	116554/50658/25272
Rat Swiss Prot No :	P49185/P49186/P49187
Immunogen :	The antiserum was produced against synthesized peptide derived from human SAPK/JNK around the phosphorylation site of Thr183. AA range:151-200



Best Tools for immunology Research		
Specificity :	Phospho-JNK1/2/3 (T183) Polyclonal Antibody detects endogenous levels of JNK1/2/3 protein only when phosphorylated at T183.	
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.	
Source :	Polyclonal, Rabbit,IgG	
Dilution :	WB 1:500 - 1:2000. IHC 1:100 - 1:300. IF 1:200 - 1:1000. ELISA: 1:5000. Not yet tested in other applications.	
Purification :	The antibody was affinity-purified from rabbit antiserum by affinity- chromatography using epitope-specific immunogen.	
Concentration :	1 mg/ml	
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)	
Observed Band :	46kD,54kD	
Cell Pathway :	Toll_Like; Cell Growth; Stem cell pathway; Insulin Receptor; MAPK_ERK_Growth;MAPK_G_Protein; ErbB/HER; B Cell Receptor; SAPK_JNK; WNT;WNT-T CELL;β-Catenin	
Background :	The protein encoded by this gene is a member of the MAP kinase family. MAP kinases act as an integration point for multiple biochemical signals, and are involved in a wide variety of cellular processes such as proliferation, differentiation, transcription regulation and development. This kinase is activated by various cell stimuli, and targets specific transcription factors, and thus mediates immediate-early gene expression in response to cell stimuli. The activation of this kinase by tumor-necrosis factor alpha (TNF-alpha) is found to be required for TNF-alpha induced apoptosis. This kinase is also involved in UV radiation induced apoptosis, which is thought to be related to cytochrom c-mediated cell death pathway. Studies of the mouse counterpart of this gene suggested that this kinase play a key role in T cell proliferation, apoptosis and differentiation. Several alternatively spl	
Function :	catalytic activity:ATP + a protein = ADP + a phosphoprotein.,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-Jun 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	



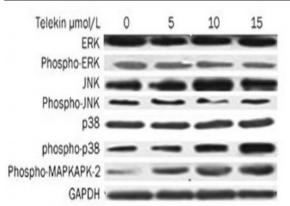
	absorber detion or such as of the provinting foretage, where the province of AD 4
	phosphorylating a number of transcription factors, primarily components of AP-1
	such as JUN, JDP
Subcellular	Cytoplasm . Nucleus . Cell junction, synapse . In the cortical neurons,
Location :	predominantly cytoplasmic and associated with the Golgi apparatus and
Location .	endosomal fraction. Increased neuronal activity increases phosphorylated form at
	synapses (By similarity). Colocalizes with POU5F1 in the nucleus.
	synapses (by similarity). Colocalizes with FOOSFT in the hucleus.
Expression :	Brain,Epithelium,Fetal brain,Lung,Pooled,Testis,
Tag:	orthogonal
	5
Contr	204
Sort :	204
No3 :	ab59196
No4 :	1
	- <u>·</u>
Host :	Rabbit
Modifications :	Phospho



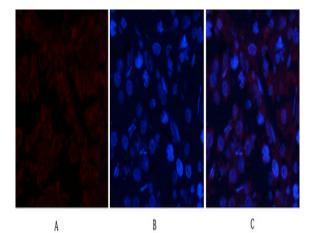
Products Images

Jiao, Zhihui, et al. "Adipose-derived stem cells protect ischemiareperfusion and partial hepatectomy by attenuating endoplasmic reticulum stress." Frontiers in cell and developmental biology 8 (2020): 177.

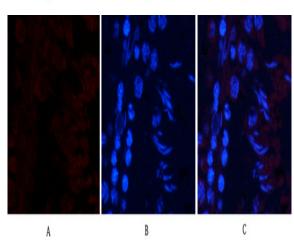




Li, Lin, et al. "Telekin suppresses human hepatocellular carcinoma cells in vitro by inducing G 2/M phase arrest via the p38 MAPK signaling pathway." Acta Pharmacologica Sinica 35.10 (2014): 1311.

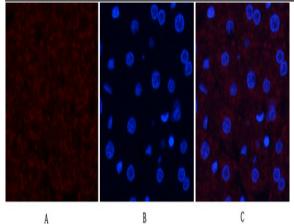


Immunofluorescence analysis of rat-testis tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

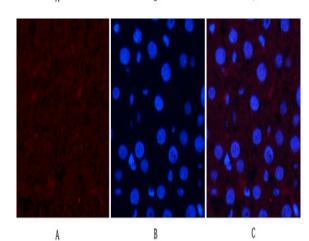


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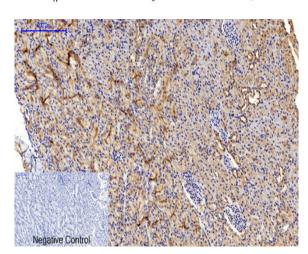




Immunofluorescence analysis of rat-liver tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

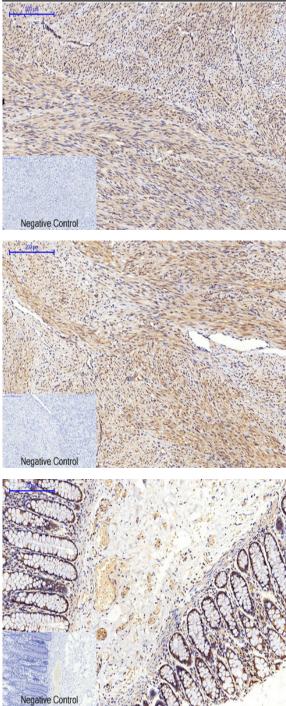


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Immunohistochemical analysis of paraffin-embedded Rat-kidney tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



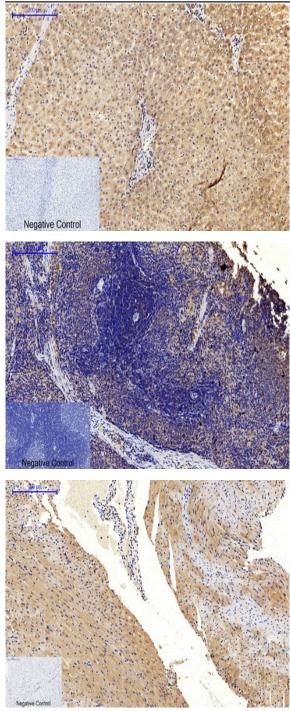


Immunohistochemical analysis of paraffin-embedded Humanuterus tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

Immunohistochemical analysis of paraffin-embedded Humanuterus-cancer tissue. 1,JNK1/2/3 (phospho Thr183) 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 Humancolon tissue. 1,JNK1/2/3 (phospho Thr183) 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.



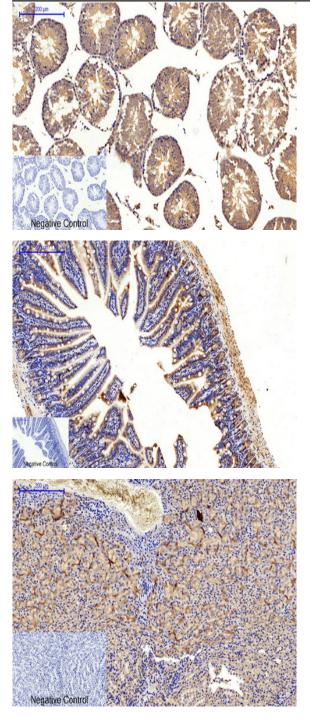


Immunohistochemical analysis of paraffin-embedded Human-liver tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

Immunohistochemical analysis of paraffin-embedded Rat-spleen tissue. 1,JNK1/2/3 (phospho Thr183) 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 Mouseheart tissue. 1,JNK1/2/3 (phospho Thr183) 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.



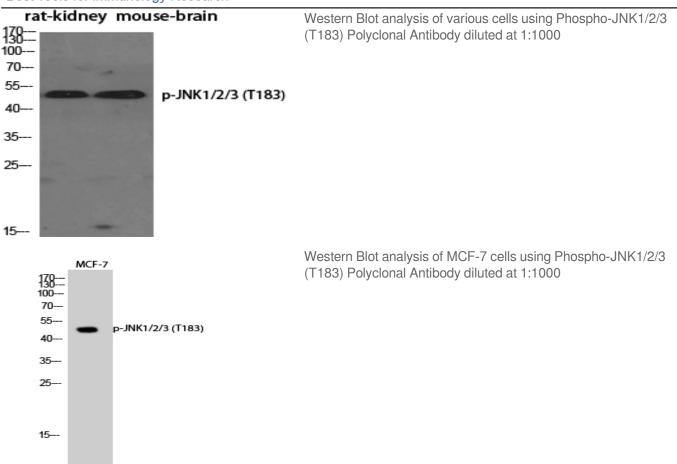


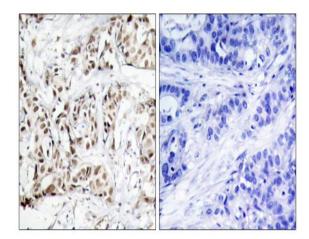
Immunohistochemical analysis of paraffin-embedded Mousetestis tissue. 1,JNK1/2/3 (phospho Thr183) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.

Immunohistochemical analysis of paraffin-embedded Mousecolon tissue. 1,JNK1/2/3 (phospho Thr183) 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 Mousekidney tissue. 1,JNK1/2/3 (phospho Thr183) 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.







Immunohistochemistry analysis of paraffin-embedded human breast carcinoma, using SAPK/JNK (Phospho-Thr183) Antibody. The picture on the right is blocked with the phospho peptide.



	117
	85
JNK2/3/SAPK	48
	34
	26
	19 (kD)

Western blot analysis of lysates from HeLa cells treated with
Anisomycin 200ng/ml 10', using SAPK/JNK (Phospho-Thr183)
Antibody. The lane on the right is blocked with the phospho peptide.