

## Rabbit Anti-phospho-P38 MAPK (Thr180) antibody

SL5476R

Product Name:	phospho-P38 MAPK (Thr180)
Chinese Name:	磷酸化p38MAPK抗体
Alias:	<ul> <li>p38 (phospho T180); p-p38 (phospho T180); MAPK14(phospho T180); CSAID Binding Protein 1; CSAID binding protein; CSAID-binding protein; Csaids binding protein; CSBP 1; CSBP 2; CSBP; CSBP1; CSBP2; CSPB 1; CSPB1; Cytokine suppressive anti inflammatory drug binding protein; Cytokine suppressive anti-inflammatory drug- binding protein; EXIP; MAP kinase 14; MAP kinase MXI2; MAP kinase p38 alpha; MAPK 14; MAPK14; MAX interacting protein 2; MAX-interacting protein 2; Mitogen Activated Protein Kinase 14; Mitogen activated protein kinase p38 alpha; MK14_HUMAN; Mxi 2; Mxi2; p38 ALPHA; p38; p38 MAP kinase; p38 MAPK; p38 mitogen activated protein kinase; p38ALPHA; p38alpha Exip; PRKM14; PRKM15; RK; SAPK 2A; SAPK2A; Stress Activated Protein Kinase 2A.</li> </ul>
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,Chicken,Dog,Pig,Horse,Rabbit,
Applications:	WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow- Cyt=1µg/TestIF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	41kDa
Cellular localization:	The nucleuscytoplasmic
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated Synthesised phosphopeptide derived from human MAPK14 around the phosphorylation site of Thr180:EM(p-T)G
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.

	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized
Storage:	antibody is stable at room temperature for at least one month and for greater than a year
	when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of
	antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
	p38 is a 38 kDa Stress Activated Protein Kinase/Map Kinase (SAPK/MAPK) that is
	fully activated by dual phosphorylation on threonine 180 and tyrosine 182, within the
	activation loop. p38 MAPK plays a critical role in the initiation of G2 delay after
	ultraviolet radiation; gene knock out studies have also revealed a critical role for p38 in
	cardiac remodeling. Downstream targets of p38 include the transcription factors ELK1
	and ATF2 and the kinases MAPKAPK2 and MAPKAPK5. p38 MAPK plays a role in
	the production of IL6 and is thought to stabilize erythropoietin production during
	hypoxic stress. It is activated by environmental stress, many proinflammatory cytokines
	and lipopolysaccharide. Dual phosphorylation by MAP2K3 and MAP2K6 is required for
	activation of p38 MAPK. It interacts with MAX, Cdc25B, Cdc25C and binds to the
	kinase interaction domain in the protein tyrosine phosphatase PTPRR; this interaction
	retains p38 MAPK in the cytoplasm.
	Function:
	Serine/threonine kinase which acts as an essential component of the MAP kinase signal
	transduction pathway. MAPK14 is one of the four p38 MAPKs which play an important
	role in the cascades of cellular responses evoked by extracellular stimuli such as
	proinflammatory cytokines or physical stress leading to direct activation of transcription
	factors. Accordingly, p38 MAPKs phosphorylate a broad range of proteins and it has
Draduat Dataile	the targets are downstroom lineage which are activated through phospharylation and
Product Detail:	further phoenhorylate additional targets DDS6V A 5/MSV1 and DDS6V A 4/MSV2 con
	directly phosphorylate and activate transcription factors such as CREB1 ATE1 the NE-
	kanna-B isoform RELA/NEKB3_STAT1 and STAT3_but can also phosphorylate
	histone H3 and the nucleosomal protein HMGN1 RPS6KA5/MSK1 and
	RPS6KA4/MSK2 play important roles in the rapid induction of immediate-early genes
	in response to stress or mitogenic stimuli either by inducing chromatin remodeling or by
	recruiting the transcription machinery. On the other hand, two other kinase targets.
	MAPKAPK2/MK2 and MAPKAPK3/MK3, participate in the control of gene expression
	mostly at the post-transcriptional level, by phosphorylating ZFP36 (tristetraprolin) and
	ELAVL1, and by regulating EEF2K, which is important for the elongation of mRNA
	during translation. MKNK1/MNK1 and MKNK2/MNK2, two other kinases activated by
	p38 MAPKs, regulate protein synthesis by phosphorylating the initiation factor EIF4E2.
	MAPK14 interacts also with casein kinase II, leading to its activation through
	autophosphorylation and further phosphorylation of TP53/p53. In the cytoplasm, the p38
	MAPK pathway is an important regulator of protein turnover. For example, CFLAR is
	an inhibitor of TNF-induced apoptosis whose proteasome-mediated degradation is
	regulated by p38 MAPK phosphorylation. In a similar way, MAPK14 phosphorylates
	the ubiquitin ligase SIAH2, regulating its activity towards EGLN3. MAPK14 may also
	inhibit the lysosomal degradation pathway of autophagy by interfering with the
	intracellular trafficking of the transmembrane protein ATG9. Another function of

MAPK14 is to regulate the endocytosis of membrane receptors by different mechanisms that impinge on the small GTPase RAB5A. In addition, clathrin-mediated EGFR internalization induced by inflammatory cytokines and UV irradiation depends on MAPK14-mediated phosphorylation of EGFR itself as well as of RAB5A effectors. Ectodomain shedding of transmembrane proteins is regulated by p38 MAPKs as well. In response to inflammatory stimuli, p38 MAPKs phosphorylate the membrane-associated metalloprotease ADAM17. Such phosphorylation is required for ADAM17-mediated ectodomain shedding of TGF-alpha family ligands, which results in the activation of EGFR signaling and cell proliferation. Another p38 MAPK substrate is FGFR1. FGFR1 can be translocated from the extracellular space into the cytosol and nucleus of target cells, and regulates processes such as rRNA synthesis and cell growth. FGFR1 translocation requires p38 MAPK activation. In the nucleus, many transcription factors are phosphorylated and activated by p38 MAPKs in response to different stimuli. Classical examples include ATF1, ATF2, ATF6, ELK1, PTPRH, DDIT3, TP53/p53 and MEF2C and MEF2A. The p38 MAPKs are emerging as important modulators of gene expression by regulating chromatin modifiers and remodelers. The promoters of several genes involved in the inflammatory response, such as IL6, IL8 and IL12B, display a p38 MAPK-dependent enrichment of histone H3 phosphorylation on 'Ser-10' (H3S10ph) in LPS-stimulated myeloid cells. This phosphorylation enhances the accessibility of the cryptic NF-kappa-B-binding sites marking promoters for increased NF-kappa-B recruitment. Phosphorylates CDC25B and CDC25C which is required for binding to 14-3-3 proteins and leads to initiation of a G2 delay after ultraviolet radiation. Phosphorylates TIAR following DNA damage, releasing TIAR from GADD45A mRNA and preventing mRNA degradation. The p38 MAPKs may also have kinase-independent roles, which are thought to be due to the binding to targets in the absence of phosphorylation. Protein O-Glc-N-acylation catalyzed by the OGT is regulated by MAPK14, and, although OGT does not seem to be phosphorylated by MAPK14, their interaction increases upon MAPK14 activation induced by glucose deprivation. This interaction may regulate OGT activity by recruiting it to specific targets such as neurofilament H, stimulating its O-Glc-N-acylation. Required in mid-fetal development for the growth of embryo-derived blood vessels in the labyrinth layer of the placenta. Also plays an essential role in developmental and stress-induced erythropoiesis, through regulation of EPO gene expression. Isoform MXI2 activation is stimulated by mitogens and oxidative stress and only poorly phosphorylates ELK1 and ATF2. Isoform EXIP may play a role in the early onset of apoptosis. Phosphorylates S100A9 at 'Thr-113'.

## Subunit:

Binds to a kinase interaction motif within the protein tyrosine phosphatase, PTPRR (By similarity). This interaction retains MAPK14 in the cytoplasm and prevents nuclear accumulation. Interacts with SPAG9 and GADD45A. Interacts with CDC25B, CDC25C, DUSP1, DUSP10, DUSP16, NP60, FAM48A and TAB1. Interacts with casein kinase II subunits CSNK2A1 and CSNK2B.

Subcellular Location: Cytoplasm. Nucleus.

**Tissue Specificity:** Brain, heart, placenta, pancreas and skeletal muscle. Expressed to a lesser extent in lung, liver and kidney. Post-translational modifications: Dually phosphorylated on Thr-180 and Tyr-182 by the MAP2Ks MAP2K3/MKK3, MAP2K4/MKK4 and MAP2K6/MKK6 in response to inflammatory citokines, environmental stress or growth factors, which a ctivates the enzyme. Dual phosphorylation can also be mediated by TAB1-mediated autophosphorylation. TCR engagement in T-cells also leads to Tyr-323 phosphorylation by ZAP70. Dephosphorylated and inactivated by DUPS1, DUSP10 and DUSP16. Acetvlated at Lys-53 and Lys-152 by KAT2B and EP300. Acetvlation at Lys-53 increases the affinity for ATP and enhances kinase activity. Lys-53 and Lys-152 are deacetylated by HDAC3. Ubiquitinated. Ubiquitination leads to degradation by the proteasome pathway. Similarity: Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. MAP kinase subfamily. Contains 1 protein kinase domain. SWISS: Q16539 Gene ID: 1432 Database links: Entrez Gene: 1432 Human Entrez Gene: 26416 Mouse Entrez Gene: 81649 Rat Entrez Gene: 403856 Dog GenBank: NM 001315 Human GenBank: NM 139012 Human GenBank: NM 011951 Mouse GenBank: NM 031020 Rat Omim: 600289 Human

	SwissProt: 002812 Dog
	SwissProt: Q16539 Human
	SwissProt: P47811 Mouse
	SwissProt: P70618 Rat
	Unigene: 485233 Human
	Unigene: 311337 Mouse
	Unigene: 88085 Rat
	Important Note:
	I his product as supplied is intended for research use only, not for use in human,
	incrapeutic of diagnostic applications.
	丝裂原活化蛋白激酶p38(p38
	MAPK、磷酸化pERK)参与细胞生长、增殖、分化、死亡及细胞间的功能同步等多种
	生理过程。P-
	p38MAPK是丝裂原活化蛋白激酶家族中的成员之一,大量研究显示p38在能量代谢
	中具有广泛的作用。p38参与脂肪组织、骨骼肌、胰岛细胞和肝脏等组织、器官的能
	量代谢。p38
	MAPK:作为细胞信号传递系统的交汇点,细胞内普遍存在的一条Signal
	transduction通路。细胞外的物理应激因子,如高渗透压、热休克、紫外线以及cell
	factor、内毒素脂多糖(LPS)等都能激活该途径,诱导细胞内蛋白质合成与分泌、Cel
	l differentiation及周亡等生物效应。p38
	MAPK
•	
	WIATK一旦饭放沽石,可以使一些转求凶于如CKEB、转求沽化凶于-I(activating
	[lactor-1, A1F-1)、A1F-2及沾化蛋白-1(AF- 1)等的好氨酸和苏氨酸位占磷酸化 活化过此结基因子 从而调节日的其田的素法





Paraformaldehyde-fixed, paraffin embedded (Rat stomach); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (p-P38 MAPK (Thr180)) Polyclonal Antibody, Unconjugated (SL5476R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (p-P38 MAPK (Thr180)) Polyclonal Antibody, Unconjugated (SL5476R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.

