p38α MAPK Antibody



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Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 40	Source/Isotype: Rabbit	UniProt ID: #Q16539	Entrez-Gene Id 1432
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		p38 α MAPK Antibody detects endogenous levels of total p38 α MAPK, regardless of its phosphorylation state. This antibody does not cross-react with other p38 MAPK isoforms such as β , γ or δ .				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy-terminal residues of human p38α MAPK. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		p38 MAP kinase (MAPK), also called RK (1) or CSBP (2), is the mammalian orthologue of the yeast HOG kinase that participates in a signaling cascade controlling cellular responses to cytokines and stress (1-4). Four isoforms of p38 MAPK, p38 α , β , γ (also known as Erk6 or SAPK3), and δ (also known as SAPK4) have been identified. Similar to the SAPK/JNK pathway, p38 MAPK is activated by a variety of cellular stresses, including osmotic shock, inflammatory cytokines, lipopolysaccharide (LPS), UV light, and growth factors (1-5). MKK3, MKK6, and SEK activate p38 MAPK by phosphorylation at Thr180 and Tyr182. Activated p38 MAPK has been shown to phosphorylate and activate MAPKAP kinase 2 (3) and to phosphorylate the transcription factors ATF-2 (5), Max (6), and MEF2 (5-8). SB203580 (4-(4-fluorophenyl)-2-(4-methylsulfinylphenyl)-5-(4-pyridyl)-imidazole) is a selective inhibitor of p38 MAPK. This compound inhibits the activation of MAPKAPK-2 by p38 MAPK and subsequent phosphorylation of HSP27 (9). SB203580 inhibits p38 MAPK catalytic activity by binding to the ATP-binding pocket, but does not inhibit phosphorylation of p38 MAPK by upstream kinases (10).				
Background References		1. Rouse, J. et al. (1994) <i>Cell</i> 78, 1027-37. 2. Han, J. et al. (1994) <i>Science</i> 265, 808-11. 3. Lee, J.C. et al. (1994) <i>Nature</i> 372, 739-46. 4. Freshney, N.W. et al. (1994) <i>Cell</i> 78, 1039-49. 5. Raingeaud, J. et al. (1995) <i>J Biol Chem</i> 270, 7420-6. 6. Zervos, A.S. et al. (1995) <i>Proc Natl Acad Sci U S A</i> 92, 10531-4. 7. Zhao, M. et al. (1999) <i>Mol Cell Biol</i> 19, 21-30. 8. Yang, S.H. et al. (1999) <i>Mol Cell Biol</i> 19, 4028-38. 9. Cuenda, A. et al. (1999) <i>Biochem Biophys Res Commun</i> 263, 825-31.				
Species Reactivi	4	C	stormined by testin	g in at least one approve	ad application (a.g.	atawa blat)

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key

H: Human M: Mouse R: Rat Mk: Monkey

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