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P38 MAPK(Phospho-Thr180) Antibody

±11252

Catalog Number: 11252-1, 11252-2

Amount: 50µg/50µl, 100µg/100µl

Swiss-Prot No. : Q16539

Form of Antibody: Rabbit IgG in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl,0.02% sodium azide and 50% glycerol.

Storage/Stability: Store at -20°C/1 year

Immunogen: The antiserum was produced against synthesized phosphopeptide derived from human p38 MAPK around the phosphorylation site of threonine 180 (E-M-T^P-G-Y).

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific phosphopeptide. The antibody against non-phosphopeptide was removed by chromatogramphy using non-phosphopeptide corresponding to the phosphorylation site.

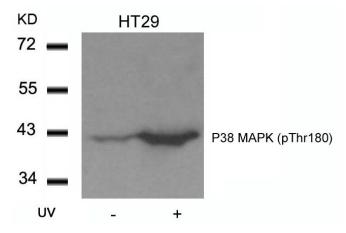
Specificity/Sensitivity: p38 MAPK (phospho-Thr180) antibody detects endogenous levels of p38 MAPK only when phosphorylated at threonine 180.

Reactivity: Human, Mouse, Rat

Applications:

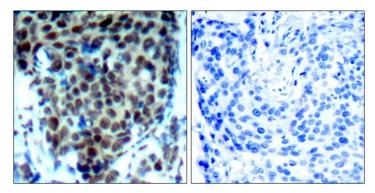
Predicted MW: 43kd

WB: 1:500~1:1000 IHC: 1:50~1:100



Western blot analysis of extracts from HT29 cells untreated or treated with UV using P38 MAPK(Phospho-Thr180) Antibody(#11252).

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Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue using p38 MAPK (phospho-Thr180) antibody (#11252).

Background :

Responds to activation by environmental stress, pro-inflammatory cytokines and lipopolysaccharide (LPS) by phosphorylating a number of transcription factors, such as ELK1 and ATF2 and several downstream kinases, such as MAPKAPK2 and MAPKAPK5. Plays a critical role in the production of some cytokines, for example IL-6. May play a role in stabilization of EPO mRNA during hypoxic stress. 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.

References:

Kim JE, et al. (2005) J Proteome Res Jul-Aug; 4(4): 1339-1346 Meng F, et al. (2005) Am J Physiol Cell Physiol May 25 Jin ZH, et al. (2005)Oncogene Mar 17; 24(12): 1973-1981 Jin ZH, et al. (2005)Oncogene Mar 17; 24(12): 1973-1981