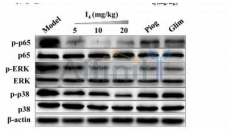
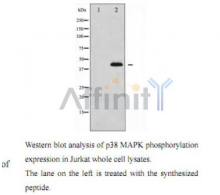
Phospho-p38 MAPK (Thr180/Tyr182) Ab

Cat.#: AF4001	Concn.: 1mg/ml	Mol.Wt.: 43 kDa
Size: 50ul,100ul,200ul	Source: Rabbit	Clonality: Polyclonal

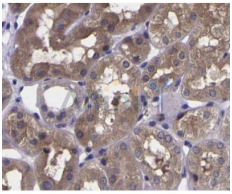
Application:	WB 1:500-1:2000 IHC 1:50-1:200 IF 1:200 IP
Reactivity:	Human,Mouse,Rat
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.
Specificity:	Phospho-p38 MAPK (Thr180/Tyr182) Ab detects endogenous levels of p38 MAPK only when phosphorylated at Thr180 and Tyr182 .
Immunogen:	A synthesized peptide derived from human p38 MAPK around the phosphorylation site of Thr180/Tyr182 .
Uniprot:	Q16539
Description:	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 environmental stresses and proinflammatory cytokines.
Subcellular Location:	Cytoplasm. Nucleus.
Tissue Specificity:	Brain, heart, placenta, pancreas and skeletal muscle. Expressed to a lesser extent in lung, liver and kidney.
Similarity:	The TXY motif contains the threonine and tyrosine residues whose phosphorylation activates the MAP kinases.Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. MAP kinase subfamily.
Storage Condition and Buffer:	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.



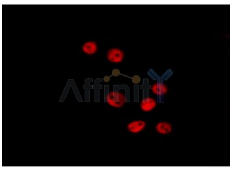
This image is a courtesy of anonymous review.http://www.sciencedirect.com/science/article/pii/S0303720716304154



Western blot analysis of p38 MAPK phosphorylation expression in Jurkat whole cell lysates.



AF4001 at 1/200 staining human kidney tissue sections by IHC-P.



AF4001 staining 293T treated with UV 30min by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

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