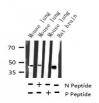


Phospho-MEK1/2 (Ser221) Ab

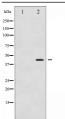
Cat.#: AF3385 Size: 100ul,200ul	Concn.: 1mg/ml Source: Rabbit	Mol.Wt.: 45kDa Clonality: Polyclonal
Application:	WB 1:500-1:2000 IHC 1:50-1:200	
Reactivity:	Human,Mouse,Rat	
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.	
Specificity:	Phospho-MEK1/2 (Ser221) Ab detects endogenous levels of MEK1/2 only when phosphorylated at Serine 221.	
Immunogen:	A synthesized peptide derived from human MEK1/2 around the phosphorylation site of Serine 221.	
Uniprot:	Q02750/P36507	
Description:	The protein encoded by this gene is a member of the dual specificity protein kinase family, which acts as a mitogen- activated protein (MAP) kinase kinase. MAP kinases, also known as extracellular signal-regulated kinases (ERKs), act as an integration point for multiple biochemical signals.	
Subcellular Location:	Nucleus;Cytoskeleton;	
Tissue Specificity:	Widely expressed, with extremely low levels in brain.	
Similarity:	The proline-rich region localized between residues 270 and 307 is important for binding to RAF1 and activation of MAP2K1/MEK1.Belongs to the protein kinase superfamily. STE Ser/Thr protein kinase family. MAP kinase kinase subfamily.	
Storage Condition and Buffer:	Rabbit lgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Store at -20 °C.Stable for 12 months from date of receipt.	



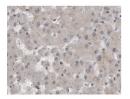
Western blot analysis of Phospho-MEK1/2 (Ser221) expression in various lysates



Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com



Western blot analysis of MEK1/2 phosphorylation expression in UV treated Jurkat whole cell lysates,The lane on the left is treated with the antigen-specific peptide.



AF3385 at 1/200 staining human liver cancer tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat antirabbit Ab was used as the secondary.

<code>IMPORTANT:</code> 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.

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