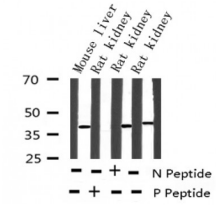
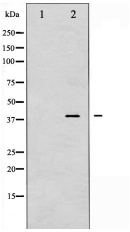
Phospho-AMPK beta-1(Ser181) Ab

Cat.#: AF3494	Concn.: 1mg/ml	Mol.Wt.: 38kDa
Size: 100ul,200ul	Source: Rabbit	Clonality: Polyclonal

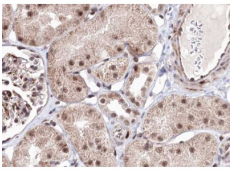
Application:	WB 1:500-1:2000 IHC 1:50-1:200, IF/ICC 1:100-1:500
Reactivity:	Human,Mouse,Rat
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.
Specificity:	Phospho-AMPK β1(Ser181) Ab detects endogenous levels of AMPK β1 only when phosphorylated at Serine 181.
Immunogen:	A synthesized peptide derived from human AMPK β1 around the phosphorylation site of Serine 181.
Uniprot:	Q9Y478
Description:	AMPKβ1 is a member of the SNF1 kinase family. It is a heterotrimeric protein comprising of an alpha (catalytic), beta (non-catalytic) and gamma (non-catalytic) subunits, 63, 38 and 38kDa respectively.AMPKβ1 regulates fatty acid and sterol synthesis by phosphorylation of acteyl-coA.
Similarity:	The glycogen-binding domain may target AMPK to glycogen so that other factors like glycogen-bound debranching enzyme or protein phosphatases can directly affect AMPK activity.Belongs to the 5'-AMP-activated protein kinase beta subunit family.
Storage Condition and Buffer:	Rabbit IgG 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-AMPK β1(Ser181) expression in various lysates



Western blot analysis of AMPK β1 phosphorylation expression in Jurkat whole cell lysates,The lane on the left is treated with the antigen-specific peptide.



AF3494 at 1/100 staining human kindey carcinoma 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 anti-rabbit Ab was used as the secondary.

AF3494 staining Hela 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.