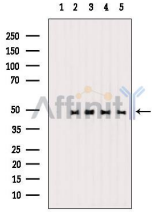
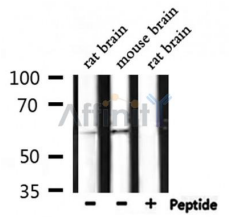
Phospho-HDAC3 (Ser424) Ab

Cat.#: AF3016	Concn.: 1mg/ml	Mol.Wt.: 48kDa
Size: 100ul,200ul	Source: Rabbit	Clonality: Polyclonal

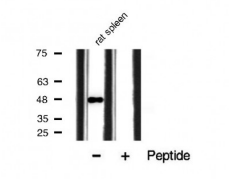
Application:	WB 1:500-1:2000 IHC 1:50-1:200 IF/ICC 1:100-1:500
Reactivity:	Human,Mouse,Rat
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.
Specificity:	Phospho-HDAC3 (Ser424) Ab detects endogenous levels of HDAC3 only when phosphorylated at Serine 424.
Immunogen:	A synthesized peptide derived from human HDAC3 around the phosphorylation site of Serine 424.
Uniprot:	O15379
Description:	Histones play a critical role in transcriptional regulation, cell cycle progression, and developmental events. Histone acetylation/deacetylation alters chromosome structure and affects transcription factor access to DNA.
Subcellular Location:	Nucleus.
Tissue Specificity:	Widely expressed.
Similarity:	Belongs to the histone deacetylase family. HD type 1 subfamily.
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.



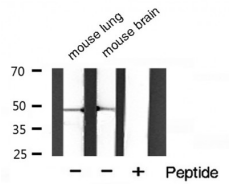
Western blot analysis of extracts from various samples, using Phospho-HDAC3 (Ser424) Ab.
Lane 1: Mouse brain treated with blocking peptide;
Lane 2: Mouse brain;
Lane 3: 3T3;
Lane 4: HepG2;
Lane 5: HeLa.



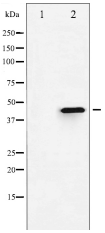
Western blot analysis of extracts from rat brain, mouse brain, using Phospho-HDAC3 (Ser424) Ab.



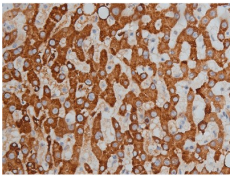
Western blot analysis of extracts of rat spleen tissue lysates,using Phospho-HDAC3 (Ser424) Ab.



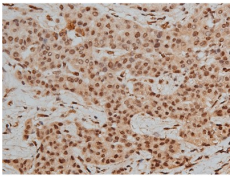
Western blot analysis of Phospho-HDAC3 (Ser424) Ab expression in mouse lung and mouse brain tissues lysates.The lane on the right is treated with the antigen-specific peptide.



Western blot analysis of HDAC3 phosphorylation expression in NIH-3T3 whole cell lysates,The lane on the left is treated with the antigen-specific peptide.

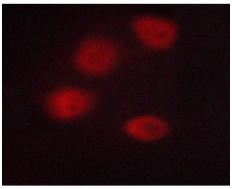


AF3016 at 1/200 staining human liver cancer tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF3016 at 1/50 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 anti-rabbit Ab was used as the secondary.

AF3016 staining NIH-3T3 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.



AF3016 staining A549 cells by ICC/IF. Cells were fixed with PFA and permeabilized in 0.1% saponin prior to blocking in 10% serum for 45 minutes at 37°C. The primary Ab was diluted 1/400 and incubated with the sample for 1 hour at 37°C. A Alexa Fluor® 594 conjugated goat polyclonal to rabbit IgG (H+L), diluted 1/600 was used as 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.