

# beta Actin Antibody

Cat.#: AF7018                      Concn.: 1mg/ml                      Mol.Wt.: 43kDa  
 Size: 50ul,100ul,200ul,1ml      Source: Rabbit                      Clonality: Polyclonal

**Application:**                      WB 1:3000-1:20000, IHC 1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000

**Reactivity:**                      Human,Mouse,Rat,Pig,Zebrafish,Bovine,Rabbit,Dog,Monkey, Fish

**Purification:**                      The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).

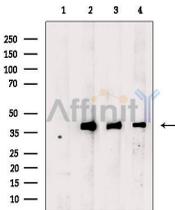
**Specificity:**                      Beta actin antibody detects endogenous levels of total Beta actin.

**Immunogen:**                      A synthesized peptide derived from human Beta actin.

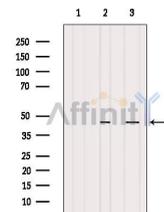
**Uniprot:**                              P60709

**Description:**                      Actin, a ubiquitous eukaryotic protein, is the major component of the cytoskeleton. At least six isoforms are known in mammals. Nonmuscle  $\beta$ - and  $\gamma$ -actin, also known as cytoplasmic actin, are predominantly expressed in nonmuscle cells, controlling cell structure and motility.

**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.



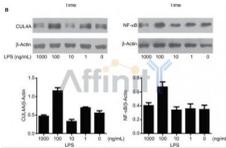
Western blot analysis of extracts from various samples, using beta Actin Antibody.  
 Lane 1: Hela cells, blocked with antigen-specific peptides,  
 Lane 2: Hela cells,  
 Lane 3: Rat liver,  
 Lane 4: MCF7 cells.



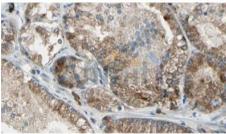
Western blot analysis of extracts from various samples, using beta Actin Antibody.  
 Lane 1: 3T3 treated with blocking peptide;  
 Lane 2: 3T3;  
 Lane 3: COS-7.



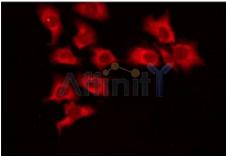
Western blot analysis of extracts from NIH/3T3 (1), Jurkat (2), rat brain (3), rat liver (4), PC12 (5) lysates, using Beta-actin Antibody



LPS enhanced the expression of CUL4A and NF-κB proteins in gastric cancer cells.



AF7018 at 1/100 staining Human prostate tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary antibody.



AF7018 staining HEPG2 cells by ICC/IF. 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 antibody 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) antibody (Red), diluted at 1/600 was used as secondary antibody.



AF7018 staining HEPG2 cells 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 antibody 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) antibody (Red), diluted at 1/600 was used as secondary antibody.

**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.

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