

## Basic Information

Product Name	Anti-GAPDH Antibody	
Gene Name	GAPDH	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat, chicken, monkey, zebrafish	
Tested Application	WB, IHC, ICC/IF, FCM	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Observed MW	36 kDa	
Dilution Ratios	Western blot (WB): Immunohistochemistry (IHC): Immunocytochemistry/Immunofluorescence (ICC/IF): Flow Cytometry (Fixed): (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or pH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	1:20000-200000 1:50-400 1:50-400 1:50-200

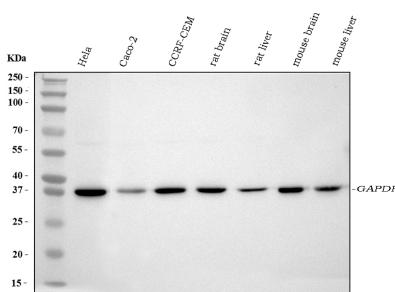
## Storage

12 months from date of receipt, -20°C as supplied.

## Reference

Anti-GAPDH Antibody被引用在125文献中。

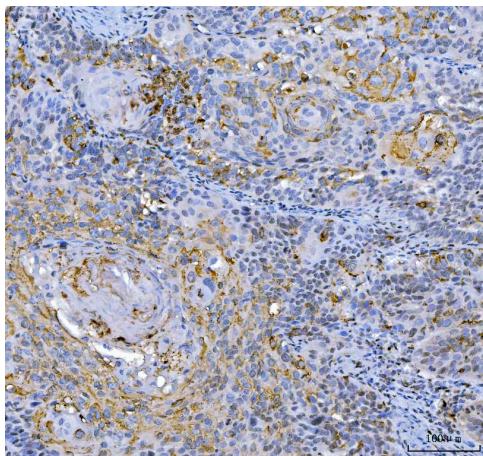
## Selected Validation Data



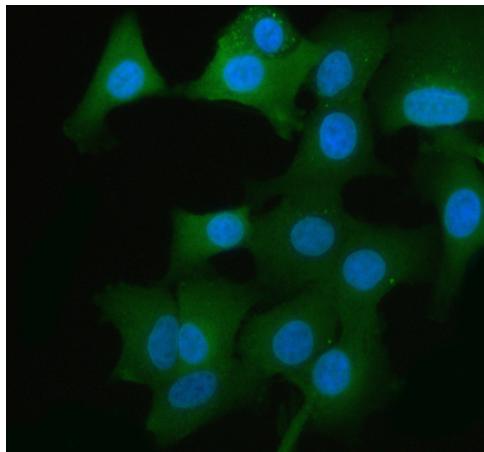
Western blot analysis of GAPDH using anti-GAPDH antibody (A00227-1). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: Hela whole cell lysates,  
Lane 2: Caco-2 whole cell lysates,  
Lane 3: CCRF-CEM whole cell lysates,  
Lane 4: rat brain tissue lysates,  
Lane 5: rat liver tissue lysates,  
Lane 6: mouse brain tissue lysates,  
Lane 7: mouse liver tissue lysates.

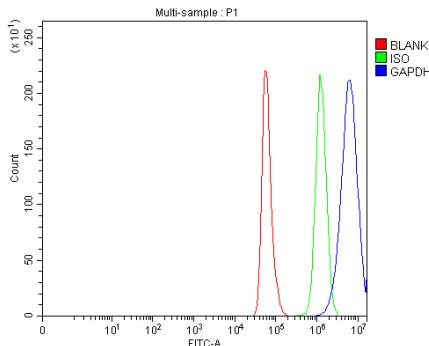
After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-GAPDH antigen affinity purified polyclonal antibody (A00227-1) at a dilution of 1:50000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for GAPDH at approximately 36 kDa. The expected band size for GAPDH is at 36 kDa.



IHC analysis of GAPDH using anti-GAPDH antibody (A00227-1). GAPDH was detected in a paraffin-embedded section of human Laryngeal squamous cell carcinoma tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-GAPDH Antibody (A00227-1) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



ICC/IF analysis of GAPDH using anti-GAPDH antibody (A00227-1). GAPDH was detected in an immunocytochemical section of A549 cells. The section was incubated with rabbit anti-GAPDH Antibody (A00227-1) at a dilution of 1:100. Fluoro488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of HeLa cells using anti-GAPDH antibody (A00227-1).

Overlay histogram showing HeLa cells stained with A00227-1 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-GAPDH Antibody (A00227-1) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.