

Basic Information

Product Name	Anti-Calbindin/CALB1 Antibody	
Gene Name	CALB1	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, IF, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human Calbindin recombinant protein (Position: A2-N261). Human Calbindin shares 98.5% amino acid (aa) sequence identity with both mouse and rat Calbindin.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	28 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 Immunofluorescence (IF) : 1:50-400 Flow Cytometry (Fixed): 1:50-200 Enzyme linked immunosorbent assay (ELISA): 1:100-1000 (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.	

Storage

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

Background Information

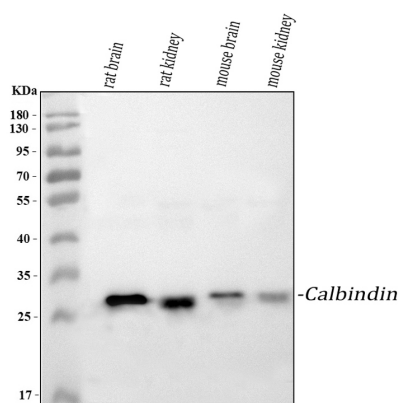
Calbindin is a calcium-binding protein belonging to the troponin C superfamily. And it mapped to 8q21.3. Calretinin is expressed in central and peripheral nervous system and in many normal and pathological tissues. The rat and human calretinin exhibit 98% sequence homology and 91% homology to many other species. Two calcium binding proteins, calbindin and calretinin, have been reported to be expressed in abundance in Purkinje cells and other cell types in the

cerebellum.

Reference

Anti-Calbindin/CALB1 Antibody被引用在1文献中。

Selected Validation Data



Western blot analysis of Calbindin/CALB1 using anti-Calbindin/CALB1 antibody (A03047). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: rat brain tissue lysates,

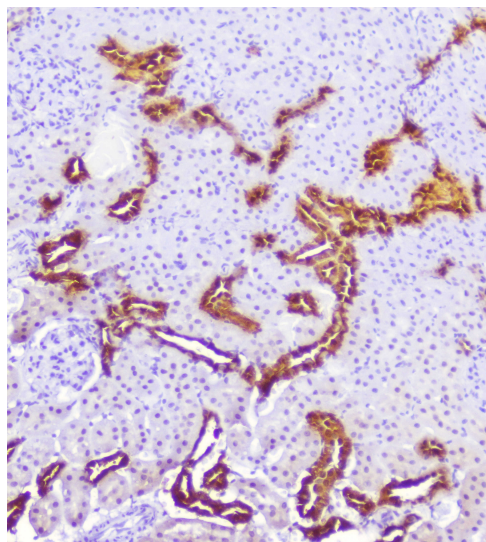
Lane 2: rat kidney tissue lysates,

Lane 3: mouse brain tissue lysates,

Lane 4: mouse kidney tissue lysates.

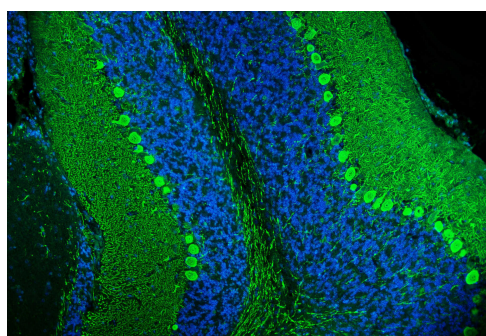
After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-Calbindin/CALB1 antigen affinity purified polyclonal antibody (A03047) at a dilution of 1:1000 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 Calbindin/CALB1 at approximately 28 kDa. The expected band size for Calbindin/CALB1 is at 30 kDa.

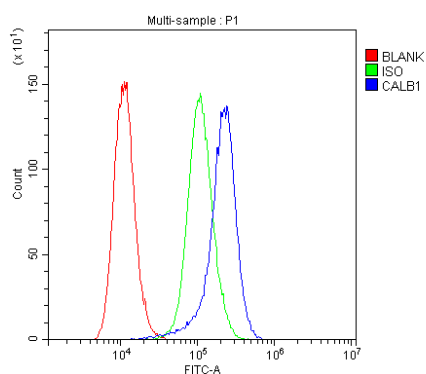


IHC analysis of Calbindin/CALB1 using anti-Calbindin/CALB1 antibody (A03047).

Calbindin/CALB1 was detected in a paraffin-embedded section of rat kidney tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-Calbindin/CALB1 Antibody (A03047) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



IF analysis using anti-Calbindin antibody (A03047), detected in paraffin-embedded section of rat cerebellum tissue. The tissue section were stained using the Fluoro488-conjugated Anti-rabbit IgG Secondary Antibody (green) (Catalog # BA1127) and counterstained with DAPI (blue).



Flow Cytometry analysis of A431 cells using anti-Calbindin/CALB1 antibody (A03047).

Overlay histogram showing A431 cells stained with A03047 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Calbindin/CALB1 Antibody (A03047) 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 (Red line) was also used as a control.