

Basic Information

Product Name	Anti-N-Cadherin/CDH2 Antibody	
Gene Name	CDH2	
Source	Rabbit	
Isotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, IHC, IF, FCM, ELISA	
Contents	500 ug/ml antibody with PBS , 0.02% NaN ₃ , 1mg BSA	
Immunogen	E.coli-derived human N Cadherin/CDH2 recombinant protein (Position: E170-E266).	
concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	140KD	
Dilution Ratios	Western blot(WB): 1:500-2000 Immunohistochemistry in paraffin section IHC 1:50-400 Immunofluorescence (IF) : 1:50-400 Flow cytometry (FCM): 1-3 µg/1x10 ⁶ cells 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. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

N-cadherin (NCAD) is a member of the cadherin cell-cell adhesion receptor family that includes P-, E-, and R-cadherin and liver cell adhesion molecule (L-CAM). N-Cadherin,, also known as Cadherin-2, encodes a 907-amino acid protein that includes a 159-amino acid signal sequence. Human and mouse nucleotide sequences are 96% identical. Mouse Ncad gene consists of 16 exons dispersed over more than 200 kb of genomic DNA. Human N-cadherin gene contains 16 exons and its sequence is highly similar to both the mouse NCAD gene (including the large first and second introns) and other cadherin genes. N-cadherin is mapped to 18q11.2. Cadherin regulates dendritic spine morphogenesis.

Reference

Anti-N-Cadherin/CDH2 Antibody 被引用在3文献中。

Selected Validation Data

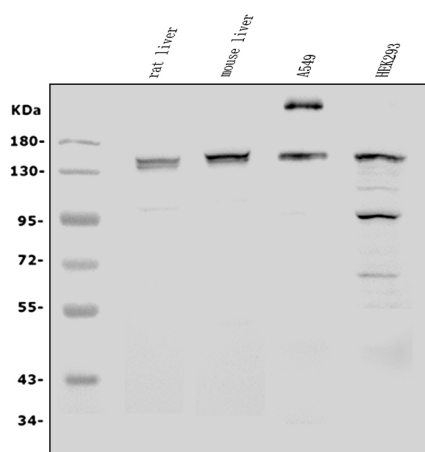


Figure 1. Western blot analysis of anti-CDH2 antibody (A01577-3). The sample well of each lane was loaded with 50ug of sample under reducing conditions. Lane 1: Rat liver tissue lysates, Lane 2: Mouse liver tissue lysates, Lane 3: human A549 whole cell lysates, Lane 4: human HEK293 whole cell lysates. Use rabbit anti-CDH2 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002). A specific band was detected for CDH2 at approximately 140 kDa. The expected band size for CDH2 is at 100 kDa.

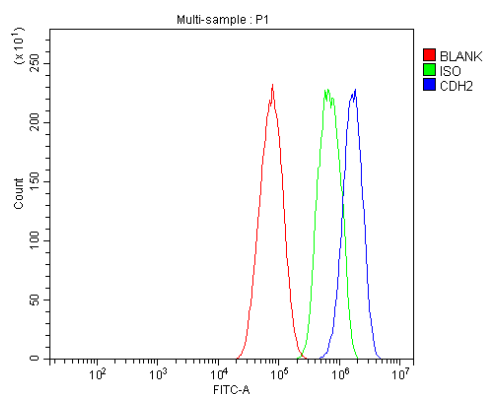


Figure 2. Flow cytometry analysis of HEPG2 cells (1x10⁶). DyLight 488 conjugated goat anti-rabbit IgG (blue line) was used as secondary antibody. Isotype control antibody (Green line) was rabbit IgG DyLight 488. Unlabelled sample (Red line).

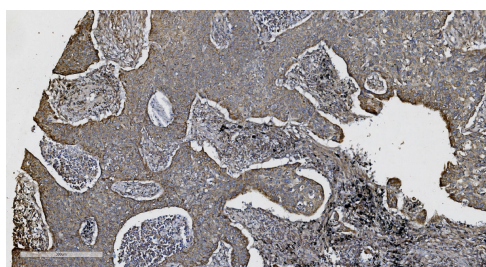


Figure 3. IHC analysis using anti-N-Cadherin/CDH2 antibody (A01577-3) detected in paraffin-embedded section of human liver cancer tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

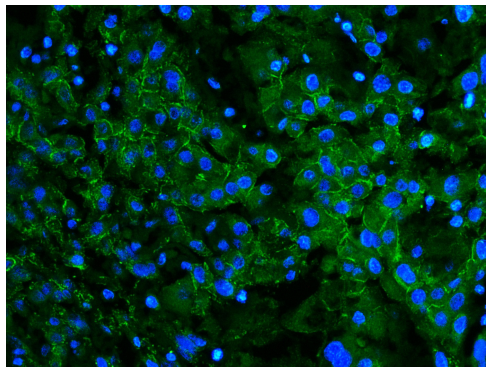


Figure 6. IF analysis using anti- CDH2 antibody (A01577-3). detected in paraffin-embedded section of human liver cancer tissue. The tissue section were stained using the Dylight488-conjugated Anti-rabbit IgG Secondary Antibody (green) (Catalog # BA1127) and counterstained with DAPI (blue).