

## Basic Information

<b>Product Name</b>	Anti-N-Cadherin/CDH2 Antibody	
<b>Gene Name</b>	CDH2	
<b>Source</b>	Rabbit	
<b>Isotype</b>	IgG	
<b>Species Reactivity</b>	human, mouse, rat	
<b>Tested Application</b>	WB, IHC, IF, FCM	
<b>Contents</b>	500 ug/ml antibody with PBS , 0.02% NaN3 , 1mg BSA	
<b>Immunogen</b>	A synthetic peptide corresponding to a sequence in the middle region of human N Cadherin(701-714aa CQCDSNGDCTDVDR), identical to the related rat and mouse sequences.	
<b>concentration</b>	500 ug/ml	
<b>Purification</b>	Immunogen affinity purified.	
<b>Observed MW</b>	140KD	
<b>Dilution Ratios</b>	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 <sup>6</sup> cells (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10文献中。

## Selected Validation Data

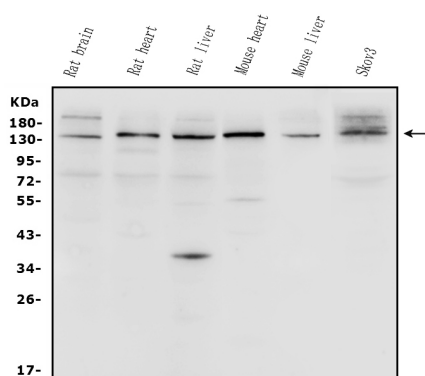


Figure 1. Western blot analysis of anti- CDH2 antibody (BA0673).The sample well of each lane was loaded with 50ug of sample under reducing conditions.Lane 1:Rat brain tissue lysates,Lane 2:Rat heart tissue lysates,Lane 3: Rat liver tissue lysates,Lane 4: Mouse heart tissue lysates,Lane 5: Mouse liver tissue lysates,Lane 6: Human SKOV3 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 140KD. The expected band size for CDH2 is at 100KD.

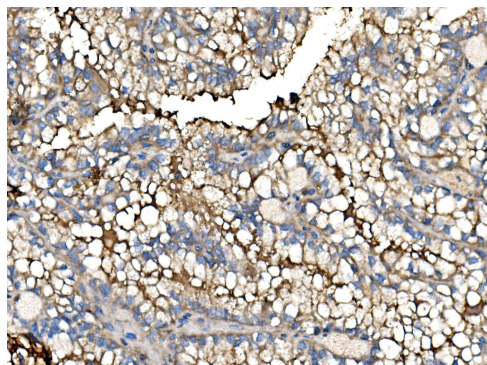


Figure 2.IHC analysis using anti- CDH2 antibody (BA0673).detected in paraffin-embedded section of human tonsil tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

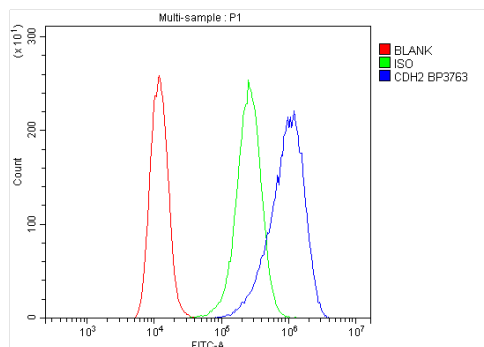


Figure 5.Flow cytometry analysis of HELA cell (1x10<sup>6</sup>) DyLight 488 conjugated goat anti- rabbit IgG(blue) was used as secondary antibody.Isotype control antibody (Green line) was rabbit IgG DyLight 488. Unlabelled sample (Red line).

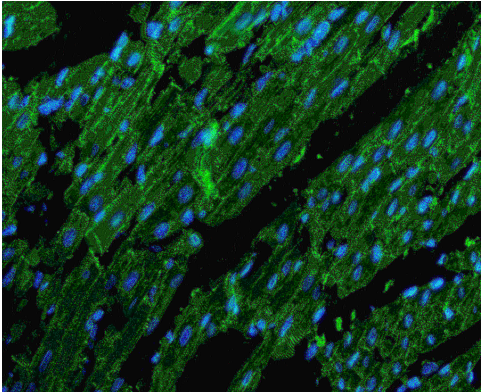


Figure 6. IF analysis using anti- CDH2 antibody (BA0673). detected in paraffin-embedded section of rat heart tissue. The tissue section were stained using the Dylight488-conjugated Anti-rabbit IgG Secondary Antibody (green) (Catalog # BA1127) and counterstained with DAPI (blue).