

## Basic Information

<b>Product Name</b>	Anti-P-Cadherin/CDH3 Antibody	
<b>Gene Name</b>	CDH3	
<b>Source</b>	Rabbit	
<b>Clonality</b>	Polyclonal	
<b>Isotype</b>	IgG	
<b>Species Reactivity</b>	human	
<b>Tested Application</b>	WB, IHC, ICC/IF	
<b>Contents</b>	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
<b>Immunogen</b>	A synthetic peptide corresponding to a sequence in the middle region of human P cadherin, different from the mouse and rat sequences by one amino acid.	
<b>Concentration</b>	500 ug/ml	
<b>Purification</b>	Immunogen affinity purified.	
<b>Observed MW</b>	120 kDa	
<b>Dilution Ratios</b>	Western blot (WB):	1:500-2000
	Immunohistochemistry (IHC):	1:50-400
	Immunocytochemistry/Immunofluorescence (ICC/IF):	1:50-400
	(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

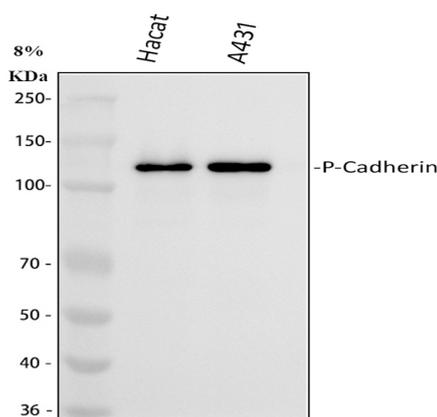
12 months from date of receipt, -20°C as supplied.

## Background Information

Cadherins, such as CDH3, are integral membrane glycoproteins responsible for calcium-dependent cell-cell adhesion. Cadherin-3 is a protein that in humans is encoded by the CDH3 gene. This gene is a classical cadherin from the cadherin superfamily. The encoded protein is a calcium-dependent cell-cell adhesion glycoprotein composed of five extracellular cadherin repeats, a transmembrane region and a highly conserved cytoplasmic tail. This gene is located in a six-cadherin cluster in a region on the long arm of chromosome 16 that is involved in loss of heterozygosity events in breast and prostate cancer. In addition, aberrant expression of this protein is observed in cervical adenocarcinomas.

Mutations in this gene have been associated with congenital hypotrichosis with juvenile macular dystrophy.

## Selected Validation Data



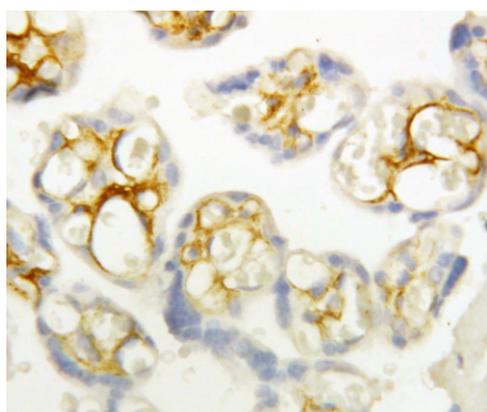
Western blot analysis of P-Cadherin/CDH3 using anti-P-Cadherin/CDH3 antibody (BA0674). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human Hacat whole cell lysates,

Lane 2: human A431 whole cell lysates.

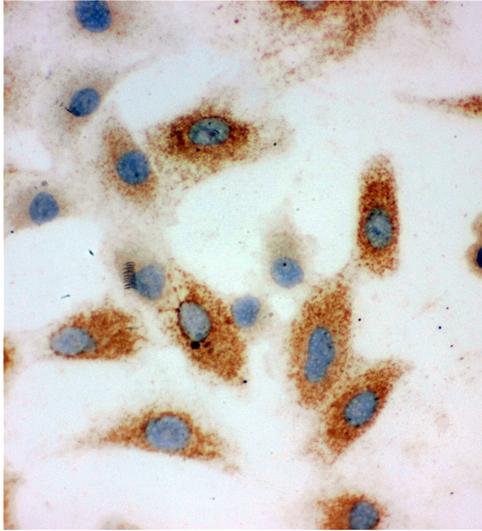
After electrophoresis, proteins were transferred to a membrane.

Then the membrane was incubated with rabbit anti-P-Cadherin/CDH3 antigen affinity purified polyclonal antibody (BA0674) 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 P-Cadherin/CDH3 at approximately 120 kDa. The expected band size for P-Cadherin/CDH3 is at 91 kDa.



IHC analysis of P-Cadherin/CDH3 using anti-P-Cadherin/CDH3 antibody (BA0674).

P-Cadherin/CDH3 was detected in a paraffin-embedded section of human placenta tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-P-Cadherin/CDH3 Antibody (BA0674) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.



ICC analysis of P-Cadherin/CDH3 using anti- P-Cadherin/CDH3 antibody (BA0674).

P-Cadherin/CDH3 was detected in an immunocytochemical section of A549 cells. The section was incubated with rabbit anti-P-Cadherin/CDH3 Antibody (BA0674) at a dilution of 1:100.

Biotinylated goat anti-rabbit IgG was used as secondary antibody.

The section was developed using Streptavidin-Biotin-Complex (SABC)(Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.