

## Datasheet for ABIN6242350

# anti-E-cadherin antibody (AA 1-392)





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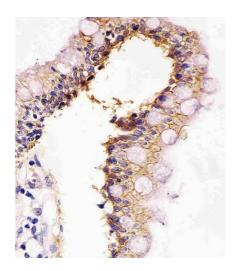
Quantity:	200 μL
Target:	E-cadherin (CDH1)
Binding Specificity:	AA 1-392
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This E-cadherin antibody is un-conjugated
Application:	Western Blotting (WB), Immunofluorescence (IF), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))
Product Details	
Immunogen:	This CADH1 antibody is generated from a mouse immunized with a recombinat protein between 1-392 amino acids from human CADH1.
Clone:	1579CT577-150-80
Isotype:	IgG1 kappa
Purification:	This antibody is purified through a protein G column, followed by dialysis against PBS.
Target Details	
Target:	E-cadherin (CDH1)
Alternative Name:	CADH1 (CDH1 Products)

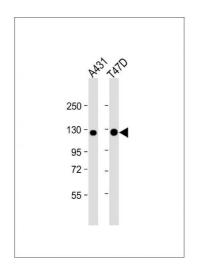
## **Target Details**

Expiry Date:

6 months

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Background:	Cadherins are calcium-dependent cell adhesion proteins. They preferentially interact with themselves in a homophilic manner in connecting cells, cadherins may thus contribute to the sorting of heterogeneous cell types. CDH1 is involved in mechanisms regulating cell-cell adhesions, mobility and proliferation of epithelial cells. Has a potent invasive suppressor role. It is a ligand for integrin alpha-E/beta-7.	
Molecular Weight:	97456	
UniProt:	P12830	
Pathways:	WNT Signaling, Sensory Perception of Sound, Cell-Cell Junction Organization, Tube Formation	
Application Details		
Application Notes:	IF: 1:25. WB: 1:4000. IHC-P: 1:25	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Buffer:	Purified monoclonal antibody supplied in PBS with 0.09 % (W/V) sodium azide.	
Preservative:	Sodium azide	
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.	
Storage:	4 °C,-20 °C	





## Immunohistochemistry (Paraffin-embedded Sections)

Image 1. (ABIN6242350 and ABIN6577163) staining CADH1 in human colon tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with formaldehyde and blocked with 3 % BSA for 0. 5 hour at room temperature, antigen retrieval was by heat mediation with a citrate buffer (pH 6). Samples were incubated with primary antibody (1/25) for 1 hours at 37 °C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.

### **Immunofluorescence**

Image 2. Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized MCF-7 (human breast cancer cell line) cells labeling CADH1 with (ABIN6242350 and ABIN6577163) at 1/25 dilution, followed by Dylight® 488-conjugated goat anti-mouse IgG (NA166821) secondary antibody at 1/200 dilution (green). Immunofluorescence image showing membrane and weak cytoplasm staining on MCF-7 cell line. The nuclear counter stain is DI (blue).

## **Western Blotting**

Image 3. All lanes: Anti-CADH1 Antibody at 1:4000 dilution Lane 1: A431 whole cell lysate Lane 2: T47D whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-mouse IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size: 97 kDa Blocking/Dilution buffer: 5 % NFDM/TBST.