Technical Data Sheet

FITC Mouse Anti- E-Cadherin

Product Information

Material Number: 612130

Alternate Name: CD324; CDH1; CADH1; Cadherin-1; ECAD; CDHE; Arc-1; LCAM; UVO; Uvomorulin

 Size:
 50 μg

 Concentration:
 250 μg/ml

 Clone:
 36/E-Cadherin

Immunogen: Human E-Cadherin C-terminal Recombinant Protein

Isotype:Mouse IgG2a, κ Reactivity:QC Testing: Human

Tested in Development: Mouse, Rat, Dog

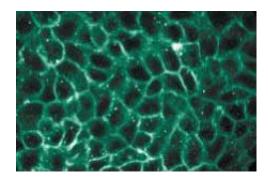
Storage Buffer: Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium

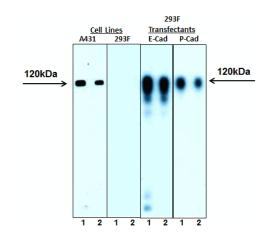
azide.

Description

E-Cadherin is a 120-kDa transmembrane glycoprotein that is localized in the adherens junctions of epithelial cells. There it interacts with the cytoskeleton through the associated cytoplasmic catenin proteins. In addition to being a calcium-dependent adhesion molecule, E-Cadherin is also a critical regulator of epithelial junction formation. Its association with catenins is necessary for cell-cell adhesion. These E-cadherin/catenin complexes associate with cortical actin bundles at both the zonula adherens and the lateral adhesion plaques. Tyrosine phosphorylation can disrupt these complexes, leading to changes in cell adhesion properties. E-Cadherin expression is often down-regulated in highly invasive, poorly differentiated carcinomas. Increased expression of E-Cadherin in these cells reduces invasiveness. Thus, loss of expression or function of E-Cadherin appears to be an important step in tumorigenic progression. The 36/E-Cadherin monoclonal antibody recognizes the cytoplasmic domain of E-Cadherin, regardless of phosphorylation status. The peptide immunogen was generated from human E-Cadherin aa. 735-883.

Note: Investigators are advised that this antibody has some degree of cross-reactivity to P-Cadherin.





LEFT IMAGE: Immunofluorescence staining of A431 cells.

RIGHT IMAGE: Western blot analysis of E-Cadherin using Purified Mouse Anti-E-Cadherin (Cat. No. 610181/610182). E-Cadherin is observable at 120kDa. Left Panel: A431 lysate (ATCC CRL-1555; Human epithelial carcinoma) was blotted at 1:10000 & 1:20000 (Lanes 1 & 2 respectively; 30 second exposure). Middle Left Panel: 293F control lysate was blotted at 1:250 & 1:500 (Lanes 1 & 2 respectively; 30 second exposure). Middle Right Panel: 293F cells transfected with human E-Cadherin (CDH1) was blotted at 1:2500 & 1:5000 (Lanes 1 & 2 respectively; 5 second exposure). Right Panel: 293 cells transfected with human P-Cadherin (CDH3) was blotted using Purified Mouse Anti-E-Cadherin (Cat. No. 610181/610182) at 1:2500 & 1:5000 (Lanes 1 & 2 respectively; 5 second exposure).

Preparation and Storage

Store undiluted at -20°C.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. The antibody was conjugated with FITC under optimum conditions, and unreacted FITC was removed.

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Page 1 of 2

612130 Rev. 2

Application Notes

Application

Immunofluorescence	Routinely Tested	
Immunoprecipitation	Not Recommended	
Immunohistochemistry	Not Recommended	

Suggested Companion Products

Catalog Number	Name	Size	Clone
610181	Purified Mouse Anti-E-Cadherin	50 μg	36/E-Cadherin
612131	FITC Mouse Anti- E-Cadherin	150 μg	36/E-Cadherin
558050	Perm Buffer III	125 mL	(none)

Product Notices

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.

References

Behrens J, Vakaet L, Friis R, et al. Loss of epithelial differentiation and gain of invasiveness correlates with tyrosine phosphorylation of the E-cadherin/beta-catenin complex in cells transformed with a temperature-sensitive v-SRC gene. J Cell Biol. 1993; 120(3):757-766. (Biology)

Jaksits S, Kriehuber E, Charbonnier AS, Rappersberger K, Stingl G, Maurer D. CD34+ cell-derived CD14+ precursor cells develop into Langerhans cells in a TGF-beta 1-dependent manner. J Immunol. 1999; 163(9):4869-4877. (Clone-specific: Flow cytometry)

Miyoshi K, Shillingford JM, Smith GH, et al. Signal transducer and activator of transcription (Stat) 5 controls the proliferation and differentiation of mammary alveolar epithelium. J Cell Biol. 2001; 155(4):531-542. (Clone-specific: Immunohistochemistry)

Sheibani N, Sorenson CM, Frazier WA. Differential modulation of cadherin-mediated cell-cell adhesion by platelet endothelial cell adhesion molecule-1 isoforms through activation of extracellular regulated kinases. Mol Biol Cell. 2000; 11(8):2793-2802. (Clone-specific: Immunofluorescence, Western blot) Takeichi M. The cadherins: cell-cell adhesion molecules controlling animal morphogenesis. Development. 1988; 102(4):639-655. (Biology)

Tamm I, Cardinale I, Kikuchi T, Krueger JG. E-cadherin distribution in interleukin 6-induced cell-cell separation of ductal breast carcinoma cells. Proc Natl Acad Sci U S A. 1994; 91(10):4338-4342. (Biology)

Weng Z, Xin M, Pablo L, et al. Protection against anoikis and down-regulation of cadherin expression by a regulatable beta-catenin protein. J Biol Chem. 2002; 277(21):18677-18686. (Clone-specific: Immunofluorescence, Immunoprecipitation, Western blot)

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612130 Rev. 2 Page 2 of 2