

# E-cadherin (N-20): sc-1500

## BACKGROUND

Cadherins comprise a family of  $\text{Ca}^{2+}$ -dependent adhesion molecules that function to mediate cell-cell binding critical to the maintenance of tissue structure and morphogenesis. Members of this family of adhesion proteins include rat cadherin-K (and its human homolog, cadherin-6), R-cadherin, B-cadherin, E/P-cadherin and cadherin-5. The classical cadherins, E-, N- and P-cadherin, consist of large extracellular domains characterized by a series of five homologous  $\text{NH}_2$ -terminal repeats. The most distal of these cadherins is thought to be responsible for binding specificity, transmembrane domains and carboxy-terminal intracellular domains. The relatively short intracellular domains interact with a variety of cytoplasmic proteins, such as  $\beta$ -catenin, to regulate cadherin function.

## CHROMOSOMAL LOCATION

Genetic locus: CDH1 (human) mapping to 16q22.1; Cdh1 (mouse) mapping to 8 D3.

## SOURCE

E-cadherin (N-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an N-terminal extracellular domain of E-cadherin of human origin.

## PRODUCT

Each vial contains 200  $\mu\text{g}$  IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-1500 P, (100  $\mu\text{g}$  peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

## APPLICATIONS

E-cadherin (N-20) is recommended for detection of E-cadherin of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu\text{g}$  per 100-500  $\mu\text{g}$  of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

E-cadherin (N-20) is also recommended for detection of E-cadherin in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for E-cadherin siRNA (h): sc-35242, E-cadherin siRNA (m): sc-35243, E-cadherin shRNA Plasmid (h): sc-35242-SH, E-cadherin shRNA Plasmid (m): sc-35243-SH, E-cadherin shRNA (h) Lentiviral Particles: sc-35242-V and E-cadherin shRNA (m) Lentiviral Particles: sc-35243-V.

Molecular Weight of E-cadherin precursor: 135 kDa.

Molecular Weight of mature E-cadherin: 120/80 kDa.

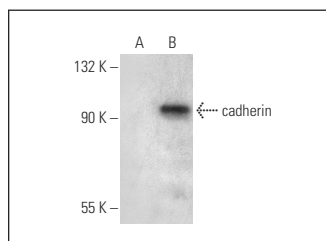
## STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## DATA



E-cadherin (N-20): sc-1500. Western blot analysis of cadherin expression in non-transfected: sc-117752 (A) and human cadherin transfected: sc-112608 (B) 293T whole cell lysates.

## SELECT PRODUCT CITATIONS

1. Hiscox, S., et al. 1999. Hepatocyte growth factor/scatter factor disrupts epithelial tumour cell-cell adhesion: involvement of  $\beta$ -catenin. *Anticancer Res.* 19: 509-518.
2. Bokobza, S.M., et al. 2010. Growth and differentiation factor 9 (GDF-9) induces epithelial-mesenchymal transition in prostate cancer cells. *Mol. Cell. Biochem.* 349: 33-40.
3. Pakala, S.B., et al. 2011. TGF- $\beta$ 1 signaling targets metastasis-associated protein 1, a new effector in epithelial cells. *Oncogene* 30: 2230-2241.
4. Cuccioloni, M., et al. 2012. *Sanguisorba minor* extract suppresses plasmin-mediated mechanisms of cancer cell migration. *Biochim. Biophys. Acta* 1820: 1027-1034.
5. Calaf, G.M., et al. 2013. Differential expression of cell adhesion molecules in an ionizing radiation-induced breast cancer model system. *Oncol. Rep.* 30: 285-291.
6. Li, G., et al. 2013. Lyn mitigates mouse airway remodeling by downregulating the TGF- $\beta$ 3 isoform in house dust mite models. *J. Immunol.* 191: 5359-5370.
7. Wang, D., et al. 2014. Oct-4 and Nanog promote the epithelial-mesenchymal transition of breast cancer stem cells and are associated with poor prognosis in breast cancer patients. *Oncotarget* 5: 10803-11085.
8. Sun, L., et al. 2015. Hypoxia promotes HO-8910PM ovarian cancer cell invasion via Snail-mediated MT1-MMP upregulation. *Exp. Biol. Med.* 240: 1434-1445.

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