# SANTA CRUZ BIOTECHNOLOGY, INC.

# E-cadherin (5F133): sc-71007



# BACKGROUND

Cadherins comprise a family of Ca<sup>2+</sup>-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 NH<sub>2</sub>-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.

#### REFERENCES

- Hirsch, H.A., et al. 1978. Surgical therapy of breast cancer. Gynakol. Rundsch. 18: 132-141.
- Takeichi, M. 1988. The cadherins: cell-cell adhesion molecules controlling animal morphogenesis. Development 102: 639-655.

#### CHROMOSOMAL LOCATION

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

# SOURCE

E-cadherin (5F133) is a mouse monoclonal antibody raised against affinity purified 80 kDa extracellular fragments of E-cadherin derived from tryptic digestion of A-431 vulva carcinoma cells of human origin.

#### PRODUCT

Each vial contains 50  $\mu g~lg G_1$  in 0.5 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

# **APPLICATIONS**

E-cadherin (5F133) 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  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500).

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.

Positive Controls: ZR-75-1 cell lysate: sc-2241, LNCaP cell lysate: sc-2231 or MCF7 whole cell lysate: sc-2206.

## STORAGE

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

# DATA



E-cadherin (5F133): sc-71007. Western blot analysis of E-cadherin expression in LNCaP whole cell lysate.

# SELECT PRODUCT CITATIONS

- 1. Yu, M., et al. 2017. Interference with Tim-3 protein expression attenuates the invasion of clear cell renal cell carcinoma and aggravates anoikis. Mol. Med. Rep. 15: 1103-1108.
- Fang, L., et al. 2018. TGF-β1 stimulates epithelial-mesenchymal transition mediated by ADAM33. Exp. Ther. Med. 15: 985-992.
- Chen, S., et al. 2018. Conversion of epithelial-to-mesenchymal transition to mesenchymal-to-epithelial transition is mediated by oxygen concentration in pancreatic cancer cells. Oncol. Lett. 15: 7144-7152.
- 4. Feng, X., et al. 2018. miR-495 enhances the efficacy of radiotherapy by targeting GRP78 to regulate EMT in nasopharyngeal carcinoma cells. Oncol. Rep. 40: 1223-1232.
- Jin, X., et al. 2019. MicroRNA-105 promotes epithelial-mesenchymal transition of nonsmall lung cancer cells through upregulating Mcl-1. J. Cell. Biochem. 120: 5880-5888.
- 6. Liu, Z., et al. 2019. Nuclear factor I/B promotes colorectal cancer cell proliferation, epithelial-mesenchymal transition and 5-fluorouracil resistance. Cancer Sci. 110: 86-98.
- 7. Li, N., et al. 2019. The role of Zeb1 in the pathogenesis of morbidly adherent placenta. Mol. Med. Rep. 20: 2812-2822.
- Hang, C., et al. 2019. MicroRNA-9 inhibits gastric cancer cell proliferation and migration by targeting neuropilin-1. Exp. Ther. Med. 18: 2524-2530.
- Kline, K.T., et al. 2020. Neonatal injury increases gut permeability by epigenetically suppressing E-cadherin in adulthood. J. Immunol. 204: 980-989.
- Li, S., et al. 2020. Effect of DEC1 on the proliferation, adhesion, invasion and epithelial-mesenchymal transition of osteosarcoma cells. Exp. Ther. Med. 19: 2360-2366.

## **RESEARCH USE**

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