

# pan-cadherin (E-11): sc-515872

## 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. 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. 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.

## REFERENCES

1. Takeichi, M. 1988. The cadherins: cell-cell adhesion molecules controlling animal morphogenesis. *Development* 102: 639-655.
2. Hatta, M., et al. 1991. Genomic organization and chromosomal mapping of the mouse P-cadherin gene. *Nucleic Acids Res.* 19: 4437-4441.

## SOURCE

pan-cadherin (E-11) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 804-829 within a C-terminal cytoplasmic domain of cadherin of human origin.

## PRODUCT

Each vial contains 200  $\mu\text{g}$  IgG<sub>2b</sub> kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

pan-cadherin (E-11) is available conjugated to agarose (sc-515872 AC), 500  $\mu\text{g}$ /0.25 ml agarose in 1 ml, for IP; to HRP (sc-515872 HRP), 200  $\mu\text{g}$ /ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-515872 PE), fluorescein (sc-515872 FITC), Alexa Fluor® 488 (sc-515872 AF488), Alexa Fluor® 546 (sc-515872 AF546), Alexa Fluor® 594 (sc-515872 AF594) or Alexa Fluor® 647 (sc-515872 AF647), 200  $\mu\text{g}$ /ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-515872 AF680) or Alexa Fluor® 790 (sc-515872 AF790), 200  $\mu\text{g}$ /ml, for Near-Infrared (NIR) WB, IF and FCM.

## RESEARCH USE

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

## APPLICATIONS

pan-cadherin (E-11) is recommended for detection of P-cadherin, N-cadherin, E-cadherin, K-cadherin, M-cadherin and R-cadherin of mouse, rat and human origin by Western Blotting (starting dilution 1:100, 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).

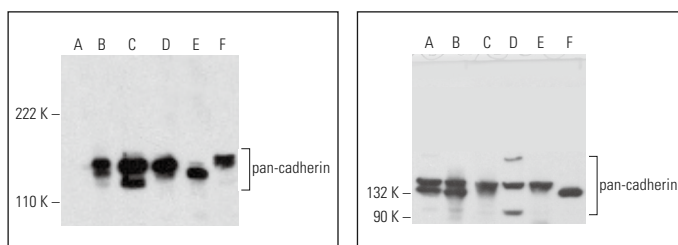
Molecular Weight of pan-cadherin: 120 kDa.

Positive Controls: P-cadherin (h2): 293T Lysate: sc-177672, F9 cell lysate: sc-2245 or mouse brain extract: sc-2253.

## RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

## DATA



pan-cadherin (E-11): sc-515872. Western blot analysis of pan-cadherin expression in non-transfected 293T: sc-117752 (A), human P-cadherin transfected 293T: sc-177672 (B), LNCaP (C) and F9 (D) whole cell lysates and mouse brain (E) and human testis (F) tissue extracts.

pan-cadherin (E-11): sc-515872. Western blot analysis of pan-cadherin expression in F9 (A), A-431 (B), MCF7 (C), Neuro-2A (D) and C6 (E) whole cell lysates and human colon tissue extract (F).

## SELECT PRODUCT CITATIONS

1. Awadia, S., et al. 2019. SGEF forms a complex with Scribble and Dlg1 and regulates epithelial junctions and contractility. *J. Cell Biol.* 218: 2699-2725.
2. Staehleke, S., et al. 2020. ROS dependent Wnt/ $\beta$ -catenin pathway and its regulation on defined micro-pillars-A combined *in vitro* and *in silico* study. *Cells* 9: 1784.
3. Yang, H., et al. 2021. Divergent regulation of OCT and MATE drug transporters by cadmium exposure. *Pharmaceutics* 13: 537.
4. Kim, D.G., et al. 2022. AIMP2-DX2 provides therapeutic interface to control KRAS-driven tumorigenesis. *Nat. Commun.* 13: 2572.
5. Yoon, I., et al. 2023. EPRS1 controls the TGF- $\beta$  signaling pathway via interaction with T $\beta$ RI in hepatic stellate cell. *Mol. Cell. Biol.* 43: 223-240.

## STORAGE

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

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.

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