# SANTA CRUZ BIOTECHNOLOGY, INC.

# ZO-2 (H-110): sc-11448



# BACKGROUND

Tight junctions are complexes of proteins that create intercellular boundaries between the plasma membrane domains of epithelial and endothelial cells. Many of the tight junction-associated proteins are members of the membrane associated guanylate kinase (MAGUK) family and include Occludin, ZO-1, ZO-2 and ZO-3. These proteins are thought to have both structural and signaling roles, and are characteristically defined by three protein-protein interaction modules: the PDZ domain, the SH3 domain and the guanylate kinase (GuK) domain. ZO-1 forms complexes with either ZO-2 or ZO-3. In addition, these proteins can also associate with Claudin, Occludin and F-Actin, at tight junction stands, where they provide a linkage between the actin cytoskeleton and the tight junction. ZO-1 expression is significantly reduced in many breast cancer lines. ZO-2 and ZO-3 are ubiquitously expressed within epithelial tight junctions, and unlike ZO-1, which is also expressed at cell junctions of cardiac myocytes, ZO-2 is not expressed in nonepithelial tissue.

# CHROMOSOMAL LOCATION

Genetic locus: TJP2 (human) mapping to 9q21.11; Tjp2 (mouse) mapping to 19 B.

#### SOURCE

ZO-2 (H-110) is a rabbit polyclonal antibody raised against amino acids 391-500 mapping within an internal region of ZO-2 (Zonula Occludens-2) of human origin.

### PRODUCT

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

# **APPLICATIONS**

ZO-2 (H-110) is recommended for detection of ZO-2 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

ZO-2 (H-110) is also recommended for detection of ZO-2 in additional species, including canine.

Suitable for use as control antibody for ZO-2 siRNA (h): sc-29833, ZO-2 siRNA (m): sc-29926, ZO-2 shRNA Plasmid (h): sc-29833-SH, ZO-2 shRNA Plasmid (m): sc-29926-SH, ZO-2 shRNA (h) Lentiviral Particles: sc-29933-V and ZO-2 shRNA (m) Lentiviral Particles: sc-29926-V.

Molecular Weight of ZO-2: 160 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200, A-431 whole cell lysate: sc-2201 or MDCK cell lysate: sc-2252.

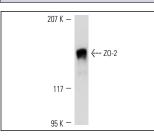
# **STORAGE**

Store at 4° C, \*\*D0 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



ZO-2 (H-110): sc-11448. Western blot analysis of ZO-2 expression in HeLa whole cell lysate.



ZO-2 (H-110): sc-11448. Immunoperoxidase staining of formalin fixed, paraffin-embedded human colorectal cancer tissue showing membrane and cytoplasmic staining of tumor cells at low (A) and high (B) magnification. Kindly provided by The Swedish Human Protein Atlas (HPA) program.

#### SELECT PRODUCT CITATIONS

- 1. Lee, S.W., et al. 2003. SSeCKS regulates angiogenesis and tight junction formation in blood-brain barrier. Nat. Med. 9: 900-906.
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- 3. Han, X., et al. 2004. Increased iNOS activity is essential for hepatic epithelial tight junction dysfunction in endotoxemic mice. Am. J. Physiol. Gastrointest. Liver Physiol. 286: G126-G136.
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- Adachi, M., et al. 2006. Normal establishment of epithelial tight junctions in mice and cultured cells lacking expression of Z0-3, a tight-junction MAGUK protein. Mol. Cell. Biol. 26: 9003-9015.
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- 8. Phua, D.C., et al. 2009. Vimentin regulates scribble activity by protecting it from proteasomal degradation. Mol. Biol. Cell 20: 2841-2855.
- Martin, T.A., et al. 2011. HAVcR-1 reduces the integrity of human endothelial tight junctions. Anticancer Res. 31: 467-473.

# MONOS Satisfation Guaranteed

Try **Z0-2 (E-3): sc-515115** or **Z0-2 (E-5): sc-514557**, our highly recommended monoclonal aternatives to Z0-2 (H-110).