## 48120 Store at -200

## Claudin-2 (E1H9O) Rabbit mAb



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Applications: WB, IP	Reactivity: H M Mk	Sensitivity: Endogenous	<b>MW (kDa):</b> 20	Source/Isotype: Rabbit IgG	<b>UniProt ID:</b> #P57739	Entrez-Gene Id 9075	
Product Usage Information	Ар	Application			Dilution		
	We	Western Blotting			1:1000		
	lmı	munoprecipitation		1:200			
Storage	•	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at $-20$ °C. Do not aliquot the antibody.					
Specificity / Sen	sitivity Cla	Claudin-2 (E1H9O) Rabbit mAb recognizes endogenous levels of total Claudin-2 protein.					
Source / Purifica		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp196 of human Claudin-2 protein.					
Background	end bloc	Tight junctions, or zonula occludens, form a continuous barrier to fluids across the epithelium and endothelium. They function in regulation of paracellular permeability and in the maintenance of cell polarity, blocking the movement of transmembrane proteins between the apical and the basolateral cell surfaces.					

Tight junctions are composed of claudin and occludin proteins, which join the junctions to the cytoskeleton (1,2). The claudin family is composed of 23 integral membrane proteins, and their expression, which varies among tissue types, may determine both the strength and properties of the epithelial barrier. Alteration in claudin protein expression pattern is associated with several types of cancer (2,3). Claudin-1 is expressed primarily in keratinocytes (4) and normal mammary epithelial cells, but is absent or reduced in breast carcinomas and breast cancer cell lines (5,6).

Claudin-2 is expressed primarily in the proximal tubule of the normal mammalian kidney, where it regulates transepithelial ion (e.g., Na+, Cl-) reabsorption (7). Increased expression of Claudin-2 has been reported in some cancer cell lines (8), including A549 lung adenocarcinoma cells, where its nuclear distribution was positively associated with enhanced proliferation (9).

## **Background References**

- 1. Shin, K. et al. (2006) Annu Rev Cell Dev Biol 22, 207-35.
- 2. Oliveira, S.S. and Morgado-Díaz, J.A. (2007) Cell Mol Life Sci 64, 17-28.
- 3. Hewitt, K.J. et al. (2006) BMC Cancer 6, 186.
- 4. Brandner, J.M. et al. (2002) Eur J Cell Biol 81, 253-63.
- 5. Krämer, F. et al. (2000) Hum Genet 107, 249-56.
- 6. Swisshelm, K. et al. (1999) Gene 226, 285-95.
- 7. Muto, S. et al. (2010) Proc Natl Acad Sci U S A 107, 8011-6.
- 8. Ikari, A. et al. (2012) Biochim Biophys Acta 1823, 1110-8.
- 9. Ikari, A. et al. (2014) Biochim Biophys Acta 1843, 2079-88.

**Species Reactivity** Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS,

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** WB: Western Blotting IP: Immunoprecipitation

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster **Cross-Reactivity Key** 

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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