#2911 Store at -20C

Phospho-Catenin δ -1 (Tyr228) Antibody



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3 Trask Lane | Danvers | Massachusetts | 01923 | USA

For Research Use Only. Not for Use in Diagnostic Procedures.

WB, IF-IC H Endogenous 95, 100 Rabbit #O60716 1500
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Product Usage
InformationApplicationDilutionWestern Blotting1:1000Immunofluorescence (Immunocytochemistry)1:100

Storage Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at –

20°C. Do not aliquot the antibody.

Specificity / Sensitivity Phospho-Catenin δ -1 (Tyr228) Antibody detects endogenous levels of catenin δ -1 protein only when

phosphorylated at Tyr228. The antibody might cross react with another overexpressed phospho-tyrosine

orotein.

Source / Purification Polyclonal antibodies are produced by immunizing animals with synthetic phosphopeptides corresponding

to residues surrounding Tyr228 of human/mouse catenin δ-1. Antibodies are purified by peptide affinity

chromatography.

Background Catenin δ-1 (p120 catenin) has an amino-terminal coiled-coil domain followed by a regulatory domain

containing multiple phosphorylation sites and a central Armadillo repeat domain of ten linked 42-amino acid repeats. The carboxy-terminal tail has no known function (1). Catenin δ -1 fulfills critical roles in the regulation of cell-cell adhesion as it regulates E-cadherin turnover at the cell surface to determine the level of E-cadherin available for cell-cell adhesion (2). Catenin δ -1 has both positive and negative effects on cadherin-mediated adhesion (3). Actin dynamics are also regulated by catenin δ -1, which modulates RhoA, Rac, and cdc42 proteins (1). Analogous to β -catenin, catenin δ -1 translocates to the nucleus, although its role at this location is unclear. Many studies show that catenin δ -1 is expressed irregularly or is absent in various types of tumor cells, suggesting that catenin δ -1 may function as a tumor suppressor (4). Catenin δ -1 is phosphorylated at multiple tyrosine sites along its sequence both *in vivo* and *in vitro* (5). High levels of catenin δ -1 phosphorylated at Tyr228 are commonly seen in several carcinoma cell lines. EGFR signaling induces catenin δ -1 phosphorylation at Tyr228, with the phosphorylated protein becoming

localized at adherens junctions although phosphorylation is not essential in junction formation (6).

Background References

- 1. Reynolds, A.B. and Roczniak-Ferguson, A. (2004) Oncogene 23, 7947-7956.
- 2. Davis, M. A. et al. (2003) J. Cell Biol. 163, 525-534.
- 3. Thoreson, M.A. and Reynolds, A.B. (2002) Differentiation 70, 583-589.
- 4. Anastasiadis, P.Z. and Reynolds, A.B. (2000) *J. Cell Sci.* 113, 1319-1334.
- 5. Mariner, D.J. et al. (2001) J. Biol. Chem. 276, 28006-28013.
- 6. Mariner, D.J. et al. (2004) J. Cell Sci. 117, 1339-1350.

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 nonfat dry

milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

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

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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Limited Uses

Phospho-Catenin δ-1 (Tyr228) Antibody (#2911) Datasheet Without Images Cell Signaling Technology

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