

## 19267

## Phospho-β-Catenin (Ser675) Antibody



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

| <b>Applications:</b><br>W, IP                                    | <b>Reactivity:</b><br>H R | <b>Sensitivity:</b><br>Endogenous  | <b>MW (kDa):</b><br>92 | <b>Source/Isotype:</b><br>Rabbit | UniProt ID:<br>#P35222      | Entrez-Gene Id:<br>1499 |
|--|---------------------------|--|------------------------|----------------------------------|-----------------------------|-------------------------|
| Product Usage<br>Information                                     |                           | <b>Application</b> Western Blotting Immunoprecipitation  |                        |                                  | <b>Dilution</b> 1:1000 1:50 |                         |
| 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- $\beta$ -Catenin (Ser675) Antibody detects endogenous levels of $\beta$ -catenin only when phosphorylated at Ser675.  |                        |                                  |                             |                         |
| Species predicted to react<br>based on 100% sequence<br>homology |                           | Mouse, Chicken, Xenop  | us, Zebrafish          |                                  |                             |                         |
| Source / Purification  |                           | Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser675 of human $\beta$ -catenin. Antibodies are purified by peptide affinity chromatography.   |                        |                                  |                             |                         |
| Background   |                           | $\beta$ -catenin is a key downstream effector in the Wnt signaling pathway (1). It is implicated in two major biological processes in vertebrates: early embryonic development (2) and tumorigenesis (3). CK1 phosphorylates $\beta$ -catenin at Ser45. This phosphorylation event primes $\beta$ -catenin for subsequent phosphorylation by GSK-3 $\beta$ (4-6). GSK-3 $\beta$ destabilizes $\beta$ -catenin by phosphorylating it at Ser33, Ser37, and Thr41 (7). Mutations at these sites result in the stabilization of $\beta$ -catenin protein levels and have been found in many tumor cell lines (8).                        |                        |                                  |                             |                         |
|  |                           | PKA was shown to phosphorylate $\beta$ -catenin at Ser675. Phosphorylation at Ser675 induces $\beta$ -catenin accumulation in the nucleus and increases its transcriptional activity (9,10).   |                        |                                  |                             |                         |
| Background Re  | ferences                  | 1. Cadigan, K.M. and Nusse, R. (1997) <i>Genes Dev</i> 11, 3286-3305. 2. Wodarz, A. and Nusse, R. (1998) <i>Annu Rev Cell Dev Biol</i> 14, 59-88. 3. Polakis, P. (1999) <i>Curr Opin Genet Dev</i> 9, 15-21. 4. Amit, S. et al. (2002) <i>Genes Dev</i> 16, 1066-76. 5. Liu, C. et al. (2002) <i>Cell</i> 108, 837-47. 6. Yanagawa, S. et al. (2002) <i>EMBO J</i> 21, 1733-42. 7. Yost, C. et al. (1996) <i>Genes Dev</i> 10, 1443-54. 8. Morin, P.J. et al. (1997) <i>Science</i> 275, 1787-90. 9. Taurin, S. et al. (2006) <i>J. Biol. Chem.</i> 281, 9971-9976. 10. Hino, S. et al. (2005) <i>Mol. Cell Biol.</i> 25, 9063-9072. |                        |                                  |                             |                         |
| Species Beastive   | :4.,                      | Coories reactivity is det  | arminad by tastin      | r in at least one approve        | . d                         |                         |

**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** 

W: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

H: Human R: Rat

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