

CTNNB1 Antibody (N-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AX10009

Specification

CTNNB1 Antibody (N-term) - Product Information

Application	IF, WB, IHC-P, FC,E
Primary Accession	P35222
Other Accession	P26233 , Q9WU82 , Q02248 , Q0VCX4 , NP_001091679.1 , F1QGH7
Reactivity	Human
Predicted	Bovine, Zebrafish, Mouse, Rat, Xenopus
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	85497
Antigen Region	78-106

CTNNB1 Antibody (N-term) - Additional Information

Gene ID 1499

Other Names

Catenin beta-1, Beta-catenin, CTNNB1, CTNNB

Target/Specificity

This CTNNB1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 78-106 amino acids from the N-terminal region of human CTNNB1.

Dilution

IF~~1:50

WB~~1:2000

IHC-P~~1:50

FC~~1:50

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

CTNNB1 Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

CTNNB1 Antibody (N-term) - Protein Information

Name CTNNB1 ([HGNC:2514](#))

Synonyms CTNNB

Function Key downstream component of the canonical Wnt signaling pathway (PubMed:[17524503](#), PubMed:[18077326](#), PubMed:[18086858](#), PubMed:[18957423](#), PubMed:[21262353](#), PubMed:[22155184](#), PubMed:[22647378](#), PubMed:[22699938](#)). In the absence of Wnt, forms a complex with AXIN1, AXIN2, APC, CSNK1A1 and GSK3B that promotes phosphorylation on N-terminal Ser and Thr residues and ubiquitination of CTNNB1 via BTRC and its subsequent degradation by the proteasome (PubMed:[17524503](#), PubMed:[18077326](#), PubMed:[18086858](#), PubMed:[18957423](#), PubMed:[21262353](#), PubMed:[22155184](#), PubMed:[22647378](#), PubMed:[22699938](#)). In the presence of Wnt ligand, CTNNB1 is not ubiquitinated and accumulates in the nucleus, where it acts as a coactivator for transcription factors of the TCF/LEF family, leading to activate Wnt responsive genes (PubMed:[17524503](#), PubMed:[18077326](#), PubMed:[18086858](#), PubMed:[18957423](#), PubMed:[21262353](#), PubMed:[22155184](#), PubMed:[22647378](#), PubMed:[22699938](#)). Involved in the regulation of cell adhesion, as component of an E-cadherin:catenin adhesion complex (By similarity). Acts as a negative regulator of centrosome cohesion (PubMed:[18086858](#)). Involved in the CDK2/PTPN6/CTNNB1/CEACAM1 pathway of insulin internalization (PubMed:[21262353](#)). Blocks anoikis of malignant kidney and intestinal epithelial cells and promotes their anchorage-independent growth by down-regulating DAPK2 (PubMed:[18957423](#)). Disrupts PML function and PML-NB formation by inhibiting RANBP2-mediated sumoylation of PML (PubMed:[22155184](#)). Promotes neurogenesis by maintaining sympathetic neuroblasts within the cell cycle (By similarity). Involved in chondrocyte differentiation via interaction with SOX9: SOX9-binding competes with the binding sites of TCF/LEF within CTNNB1, thereby inhibiting the Wnt signaling (By similarity). Acts as a positive regulator of odontoblast differentiation during mesenchymal tooth germ formation, via promoting the transcription of differentiation factors such as LEF1, BMP2 and BMP4 (By similarity). Activity is repressed in a MSX1-mediated manner at the bell stage of mesenchymal tooth germ formation which prevents premature differentiation of odontoblasts (By similarity).

Cellular Location

Cytoplasm. Nucleus. Cytoplasm, cytoskeleton {ECO:0000250|UniProtKB:B6V8E6}. Cell junction, adherens junction Cell junction {ECO:0000250|UniProtKB:B6V8E6}. Cell membrane. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome. Cytoplasm, cytoskeleton, spindle pole. Synapse {ECO:0000250|UniProtKB:Q02248} Cytoplasm, cytoskeleton, cilium basal body {ECO:0000250|UniProtKB:Q02248}. Note=Colocalized with RAPGEF2 and TJP1 at cell-cell contacts (By similarity). Cytoplasmic when it is un-stable (highly phosphorylated) or bound to CDH1. Translocates to the nucleus when it is stabilized (low level of phosphorylation). Interaction with GLIS2 and MUC1 promotes nuclear translocation. Interaction with EMD inhibits nuclear localization. The majority of CTNNB1 is localized to the cell membrane. In interphase, colocalizes with CROCC between CEP250 puncta at the proximal end of centrioles, and this localization is dependent on CROCC and CEP250. In mitosis, when NEK2 activity increases, it localizes to centrosomes at spindle poles independent of CROCC. Colocalizes with CDK5 in the cell-cell contacts and plasma membrane of undifferentiated and differentiated neuroblastoma cells Interaction with FAM53B promotes translocation to the nucleus (PubMed:25183871). Translocates to the nucleus in the presence of SNAIL1 (By similarity). Ca(2+)-mediated localization to the cell membrane in dental epithelial cells is inhibited via WNT3A (By similarity). Localizes to cell-cell contacts as keratinocyte differentiation progresses (By similarity) {ECO:0000250|UniProtKB:B6V8E6, ECO:0000250|UniProtKB:Q02248, ECO:0000269|PubMed:25183871}

Tissue Location

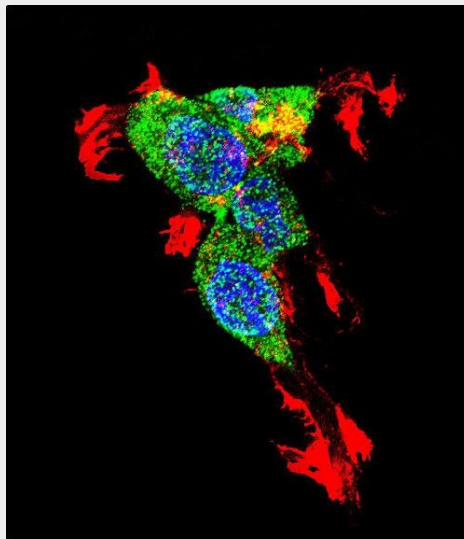
Expressed in several hair follicle cell types: basal and peripheral matrix cells, and cells of the outer and inner root sheaths. Expressed in colon. Present in cortical neurons (at protein level). Expressed in breast cancer tissues (at protein level) (PubMed:29367600).

CTNNB1 Antibody (N-term) - Protocols

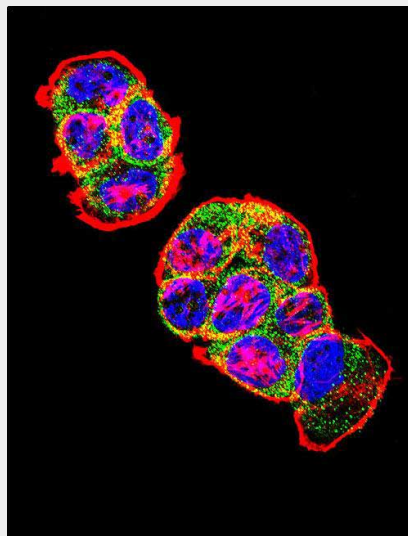
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

CTNNB1 Antibody (N-term) - Images

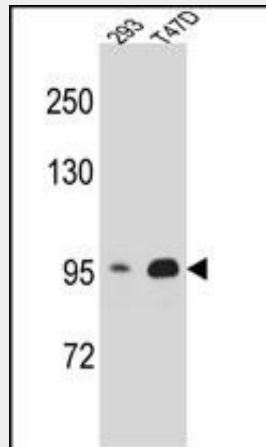


Confocal immunofluorescent analysis of CTNNB1 Antibody (N-term) with 293 cell followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). Actin filaments have been labeled with Alexa Fluor 555 phalloidin (red). DAPI was used to stain the cell nuclear (blue).

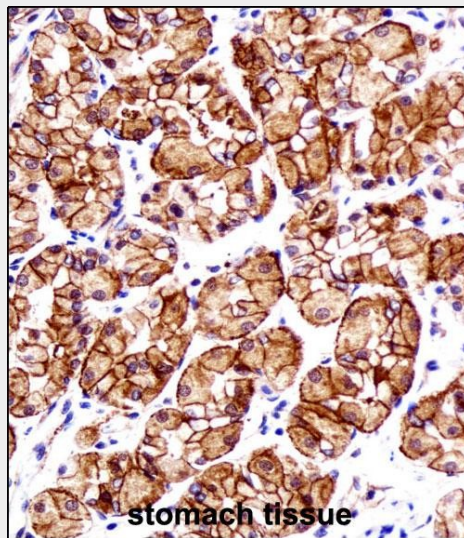


Confocal immunofluorescent analysis of CTNNB1 Antibody (N-term) with T47D cell followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). Actin filaments have been labeled with

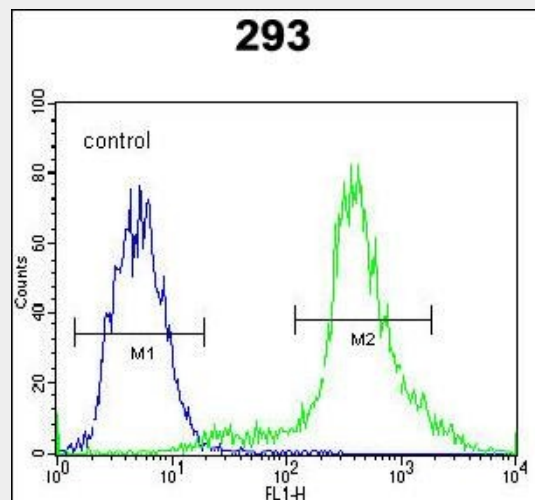
Alexa Fluor 555 phalloidin (red). DAPI was used to stain the cell nuclear (blue).



CTNNB1 Antibody (N-term) (Cat. #AX10009) western blot analysis in 293,T47D cell line lysates (35ug/lane).This demonstrates the CTNNB1 antibody detected the CTNNB1 protein (arrow).



CTNNB1 Antibody (N-term) immunohistochemistry analysis in formalin fixed and paraffin embedded human stomach tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.This data demonstrates the use of CTNNB1 Antibody (N-term) for immunohistochemistry. Clinical relevance has not been evaluated.



CTNNB1 Antibody (N-term) flow cytometric analysis of 293 cells (right histogram) compared to a negative control cell (left histogram). Alexa Fluor 488-conjugated donkey anti-rabbit IgG secondary antibodies were used for the analysis.

CTNNB1 Antibody (N-term) - Background

The protein encoded by this gene is part of a complex of proteins that constitute adherens junctions (AJs). AJs are necessary for the creation and maintenance of epithelial cell layers by regulating cell growth and adhesion between cells. The encoded protein also anchors the actin cytoskeleton and may be responsible for transmitting the contact inhibition signal that causes cells to stop dividing once the epithelial sheet is complete. Finally, this protein binds to the product of the APC gene, which is mutated in adenomatous polyposis of the colon. Mutations in this gene are a cause of colorectal cancer (CRC), pilomatixoma (PTR), medulloblastoma (MDB), and ovarian cancer. Three transcript variants encoding the same protein have been found for this gene.

CTNNB1 Antibody (N-term) - References

Huang, W., et al. Mol. Cell. Biol. 30(19):4575-4594(2010)
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Mirza, M.K., et al. J. Exp. Med. 207(8):1675-1685(2010)
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