

Rabbit Anti-Catenin Beta [EP35]: RM0008, RM0008RTU7

Intended Use: For Research Use Only

Description: Catenin Beta is a 92 kD protein normally found in the cytoplasm of the cell in the submembranous location. This protein is associated with E-Cadherin and may be essential for the function of E-Cadherin. It is a key regulatory protein involved in cell adhesion and signal transduction through the Wnt pathway, and plays important roles in development, cellular proliferation, and differentiation. Mutations in the Beta-Catenin gene CTNNB1 leading to stabilization of Beta-Catenin in the cytoplasm and translocation to the nucleus have been implicated in various forms of tumor including familial adenomatous polyposis, fibromatosis, solitary fibrous tumors and endometrial carcinoma. A nuclear accumulation of Beta-Catenin in fibromatosis (desmoid tumor) in various locations including breast and mesentery is useful in the differentiation of this tumor from other fibroblast like lesions. Nuclear accumulation of Beta-Catenin has also been demonstrated in colorectal carcinoma.

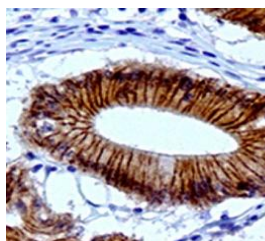
Specifications:

Clone: EP35
Source: Rabbit
Isotype: IgG
Reactivity: Human
Localization: Cytoplasm, membrane, nucleus
Formulation: Antibody in PBS pH7.2, containing < 0.2% BSA and < 0.09% sodium azide (NaN₃).
Storage: Store at 2°- 8°C.
Applications: IHC
Package:

Description	Catalog No.	Size
Catenin Beta Concentrated	RM0008	1 ml
Catenin Beta Prediluted	RM0008RTU7	7 ml

IHC Procedure*:

Positive Control Tissue: Fibromatosis of breast or abdomen
Concentrated Dilution: 50-200
Pretreatment: Citrate pH6.0, 15 minutes using Pressure Cooker, or 30-60 minutes using water bath at 95°-99°C
Incubation Time and Temp: 30-60 minutes @ RT
Detection: Refer to the detection system manual
* Result should be confirmed by an established diagnostic procedure.



FFPE human colon carcinoma stained with anti-Catenin Beta using DAB

References:

1. The Overexpression of IQGAP1 and β -Catenin Is Associated with Tumor Progression in Hepatocellular Carcinoma In Vitro and In Vivo. Jin X, et al. PLoS One 10:e0133770, 2015.
2. The essential role of TNIK gene amplification in gastric cancer growth. Yu DH, et al. Oncogenesis 2:e89, 2014.
3. Wnt signaling as a possible promoting factor of cell differentiation in pleomorphic adenomas. Okuda Y, et al. Int J Med Sci 11:971-8, 2014.

Doc. 100-RM0008
Rev. B