#### For Research Use Only

# APC Polyclonal antibody

Catalog Number: 19782-1-AP 9 Publications



**Basic Information** 

Catalog Number:

GenBank Accession Number: 19782-1-AP NM 000038

GeneID (NCBI): Size:

900 μg/ml

**UNIPROT ID:** Source: Rabbit P25054 Isotype: Full Name:

adenomatous polyposis coli

Calculated MW: 312 kDa

**Purification Method:** Antigen affinity purification Recommended Dilutions:

IHC 1:20-1:200

#### **Applications**

**Tested Applications:** 

IHC, ELISA

**Cited Applications:** 

WB

Species Specificity:

human

**Cited Species:** 

human, mouse

Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (\*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0

#### Positive Controls:

IHC: human breast cancer tissue, human colon tissue, human colon cancer tissue, human endometrial cancer

### **Background Information**

APC, also named as DP2.5, belongs to the adenomatous polyposis coli (APC) family. APC is a tumor suppressor that regulates cell division, helps ensure that the number of chromosomes in a cell is correct following cell division, and associates with other proteins involved in cell attachment and signaling. APC promotes rapid degradation of CTNNB1 and participates in Wnt signaling as a negative regulator. It plays a critical role in several cellular  $processes. APC\ regulates\ beta-catenin\ levels\ through\ Wnt-signaling\ and\ is\ involved\ in\ actin\ cytoskeletal\ integrity,$ cell-cell adhesion and cell migration. APC activity is correlated with its phosphorylation state. Defects in APC are a cause of familial adenomatous polyposis (FAP) which includes also Gardner syndrome (GS). Defects in APC are a cause of hereditary desmoid disease (HDD) which also known as familial infiltrative fibromatosis (FIF). Defects in APC are a cause of medulloblastoma (MDB) which is a malignant, invasive embryonal tumor of the cerebellum with a preferential manifestation in children. Defects in APC are a cause of mismatch repair cancer syndrome (MMRCS) which also known as Turcot syndrome or brain tumor-polyposis syndrome 1 (BTPS1).

#### **Notable Publications**

Author	Pubmed ID	Journal	Application
Xiaobo Hu	31637871	Cancer Med	WB
Yang Zhou	31627092	Biomed Pharmacother	WB
Hongting Guo	34786330	J Bone Oncol	WB

#### Storage

Store at -20°C. Stable for one year after shipment.

Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

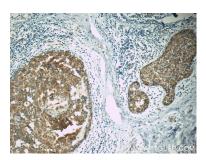
Aliquoting is unnecessary for -20°C storage

For technical support and original validation data for this product please contact:

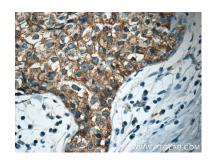
T: 4006900926 E: Proteintech-CN@ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

## Selected Validation Data



Immunohistochemical analysis of paraffinembedded human breast cancer tissue slide using 19782-1-AP (APC Antibody) at dilution of 1:50. Heat mediated antigen retrieved with Citric acid buffer, pH6.0.



Immunohistochemical analysis of paraffinembedded human breast cancer tissue slide using 19782-1-AP (APC Antibody) at dilution of 1:50. Heat mediated antigen retrieved with Citric acid buffer, pH6.0.