

TCTN2 Antibody

Rabbit Polyclonal Antibody Catalog # ABV10207

Specification

TCTN2 Antibody - Product Information

Application
Primary Accession
Reactivity

Host Clonality Isotype

Calculated MW

WB

Q3B7D3

Human, Mouse, Rat

Rabbit Polyclonal Rabbit IgG 77301

TCTN2 Antibody - Additional Information

Gene ID 689779

Positive Control

Application & Usage

Jurkat cell lysate

Western Blot analysis (0.5-4 µg/ml). However, the optimal concentrations should be determined individually. Blocking peptide is available separately.

Other Names Tectonic-2

Target/Specificity

TCTN2

Antibody Form

Liquid

Appearance

Colorless liquid

Formulation

 $100~\mu g$ (0.5 mg/ml) affinity purified rabbit anti- TCTN2 polyclonal antibody in phosphate buffered saline (PBS), pH 7.2, containing 30% glycerol, 0.5% BSA, 5 mM EDTA and 0.01% thimerosal

Handling

The antibody solution should be gently mixed before use.

Reconstitution & Storage

-20 °C

Background Descriptions

Precautions

TCTN2 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.



TCTN2 Antibody - Protein Information

Name Tctn2

Synonyms Tect2

Function

Component of the tectonic-like complex, a complex localized at the transition zone of primary cilia and acting as a barrier that prevents diffusion of transmembrane proteins between the cilia and plasma membranes. Required for hedgehog signaling transduction (By similarity).

Cellular Location

Membrane; Single-pass type I membrane protein. Cytoplasm, cytoskeleton, cilium basal body. Note=Localizes at the transition zone, a region between the basal body and the ciliary axoneme.

TCTN2 Antibody - Protocols

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

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

TCTN2 Antibody - Images

TCTN2 Antibody - Background

TCTN2 is a member of the Tectonic protein family. Tectonic proteins are transmembrane proteins that regulate the Hedgehog (Hh)-signaling pathway. Defects in TCTN2 can lead to Meckel syndrome type 8. This disorder is characterized by a combination of renal cysts and variable associated features which includes developmental anomalies of the central nervous system, hepatic ductal dysplasia and cysts, and polydactyly.