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Recombinant TSPAN1 Monoclonal Antibody

catalog number: AN300264P

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description

Reactivity Human

Immunogen Recombinant Human TSPAN1 Protein

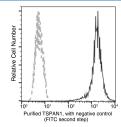
HostRabbitIsotypeIgGCloneB182PurificationProtein A

Buffer 0.2 µm filtered solution in PBS

Applications Recommended Dilution

IHC-P 1:5000-1:20000 **FCM** 1:25-1:100

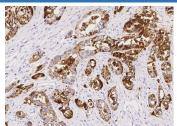
Data



Flow cytometric analysis of Human TSPAN1 expression on SW480 cells. The cells were stained with purified anti-Human TSPAN1, then a FITC-conjugated second step antibody. The fluorescence histograms were derived from gated events with the forward and side light-scatter characteristics of intact cells.



Immunohistochemistry of paraffin-embedded human prostate using TSPAN1 Monoclonal Antibody at dilution of 1:10000.



Immunohistochemistry of paraffin-embedded human rectal cancer using TSPAN1 Monoclonal Antibody at dilution of 1:10000.

Preparation & Storage

Storage This antibody can be stored at 2°C-8°C for one month without detectable loss of

activity. Antibody products are stable for twelve months from date of receipt when stored at -20°C to -80°C. Preservative-Free. Avoid repeated freeze-thaw cycles.

Shipping Ice bag

Background

For Research Use Only

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TSPAN1 belongs to the transmembrane 4 superfamily, also known as the tetraspanin family. Tetraspanins have four hydrophobic domains, intracellular N- and C-termini and two extracellular domains. Tetraspanins act as scaffolding proteins, anchoring multiple proteins to one area of the cell membrane. They also mediate signal transduction events that play a role in the regulation of cell development, activation, growth and motility. TSPAN1 interacts with human thiamine transporter-1 (hTHTR-1). HTHTR-1 contributes to intestinal thiamine uptake, and its function is regulated at both the transcriptional and posttranscriptional levels. TSPAN1 and hTHTR-1 colocalize in human intestinal epithelial HuTu-8 cells. Coexpression of TSPAN1 in these cells led to a significant decrease in the rate of degradation of hTHTR-1 compared with cells expressing the hTHTR-1 alone; in fact the half-life of the TSPAN1 protein was twice longer in the former cell type compared with the latter cell type.

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