

Anti-PDCD12 Antibody

Rabbit polyclonal antibody to PDCD12 Catalog # AP60849

Product Information

ApplicationWB, IF/IC, IHCPrimary AccessionQ9NQS1

Reactivity Human, Mouse, Rat, Monkey

HostRabbitClonalityPolyclonalCalculated MW38506

Additional Information

Gene ID 57099

Other Names Cell death regulator Aven

Target/Specificity KLH-conjugated synthetic peptide encompassing a sequence within the

C-term region of human PDCD12. The exact sequence is proprietary.

Dilution WB~~WB (1/500 - 1/2000), IHC (1/50 - 1/200), IF/IC (1/50 - 1/100) IF/IC~~N/A

IHC~~WB (1/500 - 1/2000), IHC (1/50 - 1/200), IF/IC (1/50 - 1/100)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name AVEN

Function Protects against apoptosis mediated by Apaf-1.

Cellular Location Endomembrane system; Peripheral membrane protein. Note=Associated with

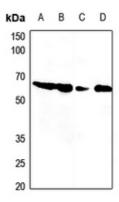
intracellular membranes

Tissue Location Highly expressed in testis, ovary, thymus, prostate, spleen, small intestine,

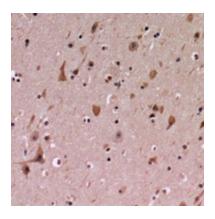
colon, heart, skeletal muscle, liver, kidney and pancreas

Background

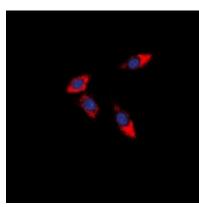
KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human PDCD12. The exact sequence is proprietary.



Western blot analysis of PDCD12 expression in HEK293T (A), Hela (B), mouse muscle (C), rat muscle (D) whole cell lysates.



Immunohistochemical analysis of PDCD12 staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of PDCD12 staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.