

AP17828PU-N**Polyclonal Antibody to Villin-1 (N-term)****Alternate names:**

VIL1

Quantity:

0.4 ml

Concentration:

0.5 mg/ml (Lot specific)

Background:

Villin-1 is a member of a family of calcium-regulated actin-binding proteins. This protein represents a dominant part of the brush border cytoskeleton which functions in the capping, severing, and bundling of actin filaments.

Uniprot ID:[P09327](#)**NCBI:**[NP_009058](#)**GeneID:**[7429](#)**Host:**

Rabbit

Immunogen:

KLH conjugated synthetic peptide between 180-207 amino acids from the N-terminal region of Human Villin-1.

Format:**State:** Liquid purified Ig fraction**Purification:** Purified through a protein A column, followed by peptide affinity purification.**Buffer System:** PBS**Preservatives:** 0.09% (W/V) Sodium Azide**Applications:****ELISA:** 1/1,000.**Western blotting:** 1/1,000.**Flow Cytometry:** 1/10-1/50.**Immunohistochemistry on Paraffin Sections:** 1/50-1/100.

Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.

Specificity:

This antibody recognizes Villin-1.

Species Reactivity:**Tested:** Human, Mouse.**Expected from sequence similarity:** Bovine, Pig.**Storage:**

Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Shelf life: one year from despatch.

General Readings:

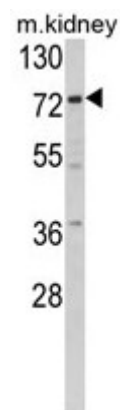
1. Yamamichi, N., et al., Exp. Cell Res. 315 (10), 1779-1789 (2009)

Pictures:

Formalin-fixed and paraffin-embedded human colon carcinoma reacted with Villin-1 Antibody (N-term), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining.



Western blot analysis of Villin-1 Antibody (N-term) (AP17828PU-N) in mouse kidney tissue lysates (35ug/lane). VIL1 (arrow) was detected using the purified Pab.



Flow Cytometric analysis of WiDr cells using Villin-1 Antibody (N-term) (AP17828PU-N) (bottom histogram) compared to a negative control cell (top histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

