

## **Product datasheet for TA389230**

## **VASP Mouse Antibody [Clone ID: M277]**

**Product data:** 

**Product Type:** Primary Antibodies

Clone Name: M277

Applications: ICC, WB

Recommended Dilution: WB: 1:1000

**ICC**: 1:50

Reactivity: Human
Host: Mouse

Isotype: lgG1

**Immunogen:** Clone (M277) was generated from a recombinant protein that includes amino acids from the

C-terminal region of human VASP.

**Specificity:** This antibody detects 46 and 50 kDa\* proteins corresponding to the apparent molecular

mass of VASP on SDS-PAGE immunoblots of human A431, HeLa, and HUVEC.

Formulation: PBS + 1 mg/ml BSA, 0.05% NaN3 and 50% glycerol

**Concentration:** lot specific

**Purification:** Protein A Purified

**Conjugation:** Unconjugated

Storage: Storage at -20°C is recommended, as aliquots may be taken without freeze/thawing due to

presence of 50% glycerol. Stable for at least 1 year at -20°C.

**Stability:** After date of receipt, stable for at least 1 year at -20°C.

Predicted Protein Size: 46/50

Database Link: P50552



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Background:

Actin filament tethering and bundling are important mechanisms involved in actin superstructure assembly. The ENA/VASP family includes VASP, mena, and Ena-Vasp-like (EVL). These multidomain proteins localize to the leading edge of filopodia where they associate with AFs, interact with profilin, and compete with capping proteins at the barbed end of AFs. Artificial relocalization of VASP from the plasma membrane to mitochondrial membranes inhibits filopodial formation and axon branching, while deletion of all three ENA/VASP proteins produces defects in cortical axon-tract formation. Regulation of VASP protein activity occurs through phosphorylation at Ser-157, Ser-239, and Thr-278. AMPK phosphorylates Thr-278, leading to impaired actin stress fiber assembly and changes in cell morphology.

Note:

Protein G purified tissue culture supernatant.