

VCIP135 Antibody

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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IP	Reactivity: H Mk	Sensitivity: Endogenous	MW (kDa): 140	Source/Isotype: Rabbit	UniProt ID: #Q96JH7	Entrez-Gene Id: 80124
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Product Usage Information**Application**

Western Blotting
Immunoprecipitation

Dilution

1:1000
1:100

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at -20°C. Do not aliquot the antibody.

Specificity/Sensitivity

VCIP135 Antibody recognizes endogenous levels of total VCIP135 protein.

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala1035 of human VCIP135 protein. Antibodies are purified by protein A and peptide affinity chromatography.

Background

Protein ubiquitination and deubiquitination are reversible processes catalyzed by ubiquitinating enzymes and deubiquitinating enzymes, respectively (1,2). Deubiquitinating enzymes (DUBs) are categorized into five subfamilies based on catalytic domain structure: USP, OTU, MJD, and JAMM. The valosin-containing protein p97/p47 complex-interacting protein 1 (VCIP135, VCPIP1) is a deubiquitinating enzyme that belongs to the A20-like subfamily of ovarian tumor (OTU) DUBs (3). VCIP135 serves as a cofactor for the p97/p47 complex in regulating Golgi membrane fusion and reassembly at the end of mitosis (4-6). Research studies suggest that the phosphorylation status of VCIP135 provides a mechanism to fine-tune the kinetics of Golgi disassembly and reassembly during the cell cycle. For example, these studies demonstrate that VCIP135 undergoes phosphorylation early in mitosis, which blocks its association with the Golgi membrane and p97/VCP, thus inhibiting p97/VCP-mediated Golgi membrane fusion (7,8).

Background References

1. Nijman, S.M. et al. (2005) *Cell* 123, 773-86.
2. Nalepa, G. et al. (2006) *Nat Rev Drug Discov* 5, 596-613.
3. Mevissen, T.E. et al. (2013) *Cell* 154, 169-84.
4. Wang, Y. et al. (2004) *J Cell Biol* 164, 973-8.
5. Uchiyama, K. et al. (2002) *J Cell Biol* 159, 855-66.
6. Totsukawa, G. et al. (2011) *EMBO J* 30, 3581-93.
7. Zhang, X. et al. (2014) *J Cell Sci* 127, 172-81.
8. Totsukawa, G. et al. (2013) *Biochem Biophys Res Commun* 433, 237-42.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween@ 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting **IP:** Immunoprecipitation

Cross-Reactivity Key

H: Human **Mk:** Monkey

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