

**VAMP3 (D9S6K) XP<sup>®</sup> Rabbit mAb**

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<b>Applications:</b> W, IHC-P	<b>Reactivity:</b> H Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 13	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q15836	<b>Entrez-Gene Id:</b> 9341
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**Product Usage Information****Application**

Western Blotting  
Immunohistochemistry (Paraffin)

**Dilution**

1:1000  
1:800

**Storage**

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

**Specificity/Sensitivity**

VAMP3 (D9S6K) XP<sup>®</sup> Rabbit mAb recognizes endogenous levels of total VAMP3 protein.

**Source / Purification**

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala8 of human VAMP3 protein.

**Background**

Proteins in the soluble N-ethylmaleimide-sensitive factor attachment protein receptor (SNARE) complex are integral membrane proteins involved in vesicle transport and membrane fusion by pairing of vesicular SNAREs (v-SNAREs) with cognate target SNAREs (t-SNAREs) (reviewed in 1,2). Vesicle associated membrane protein 3 (VAMP3), also known as cellubrevin, has a broad tissue distribution and localizes to endosomal compartments (3). VAMP3 interacts with the t-SNAREs syntaxin1, syntaxin4, SNAP23, and SNAP25 (4,5). Research studies indicate that VAMP3 is involved in transferrin receptor recycling to the plasma membrane (6) and in T-cell receptor recycling to immunological synapses (7). Inhibition of VAMP3 with tetanus toxin impairs membrane trafficking during cell migration (8).

**Background References**

1. Jena, B.P. (2011) *Adv Exp Med Biol* 713, 13-32.
2. Kasai, H. et al. (2012) *Physiol Rev* 92, 1915-64.
3. McMahon, H.T. et al. (1993) *Nature* 364, 346-9.
4. Chilcote, T.J. et al. (1995) *J Cell Biol* 129, 219-31.
5. Schraw, T.D. et al. (2003) *Biochem J* 374, 207-17.
6. Galli, T. et al. (1994) *J Cell Biol* 125, 1015-24.
7. Das, V. et al. (2004) *Immunity* 20, 577-88.
8. Tayeb, M.A. et al. (2005) *Exp Cell Res* 305, 63-73.

**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 **IHC-P:** Immunohistochemistry (Paraffin)

**Cross-Reactivity Key**

**H:** Human **Mk:** Monkey

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