

Product Data Sheet

Anti-VAMP1/2/3 Antibody

Catalog # Source Reactivity Applications

CPA4861 Rabbit H, M, R, B WB, IH

Description Rabbit polyclonal antibody to VAMP1/2/3

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human VAMP1/2/3. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of VAMP1/2/3 protein.

Clonality Polyclonal

Conjugation

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

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/100 - 1/200)

Gene Symbol VAMP1; VAMP2; VAMP3

Alternative Names VAMP1; SYB1; Vesicle-associated membrane protein 1; VAMP-1; Synaptobrevin-1;

VAMP2; SYB2; Vesicle-associated membrane protein 2; VAMP-2; Synaptobrevin-2;

VAMP3; SYB3; Vesicle-associated membrane protein 3; VAMP-3; Cellubrevin; CEB;

Synaptobrevin-3

Entrez Gene 6843, 6844, 9341 (Human); 22317, 22318, 22319 (Mouse); 25624, 24803, 29528

(Rat)

SwissProt P23763, P63027, Q15836 (Human); Q62442, P63044, P63024 (Mouse); Q63666,

P63045, P63025 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

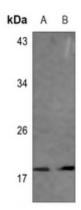
Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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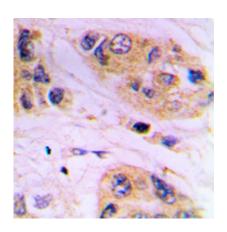
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Western blot analysis of VAMP1/2/3 expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 12; 11 kD; Observed band size: 18 kD)



Immunohistochemical analysis of VAMP1/2/3 staining in human prostate cancer 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.

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