

## **Product Data Sheet**

## **Anti-Nav1.2 Antibody**

Catalog # Source Reactivity Applications

CPA5986 Rabbit H, M, R WB, IH

**Description** Rabbit polyclonal antibody to Nav1.2

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

region of human Nav1.2. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of Nav1.2 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 (1/50 - 1/200)

Gene Symbol SCN2A

Alternative Names NAC2; SCN2A1; SCN2A2; Sodium channel protein type 2 subunit alpha; HBSC II;

Sodium channel protein brain II subunit alpha; Sodium channel protein type II

subunit alpha; Voltage-gated sodium channel subunit alpha Nav1.2

Entrez Gene 6326 (Human); 24766 (Rat)

**SwissProt** Q99250 (Human); P04775 (Rat)

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

freeze/thaw cycles.

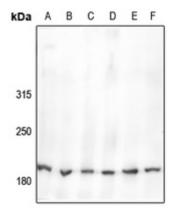
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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Western blot analysis of Nav1.2 expression in C6 (A), BV2 (B), rat brain (C), mouse brain (D), HEK293T (E), LO2 (F) whole cell lysates. (Predicted band size: 227 kD; Observed band size: 200 kD)



Immunohistochemical analysis of Nav1.2 staining in human brain 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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