

## **Product Data Sheet**

## **Anti-GABRA1 Antibody**

Catalog # Source Reactivity Applications

CPA3225 Rabbit H, M, R, B, C, Mk WB, IH

**Description** Rabbit polyclonal antibody to GABRA1

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

region of human GABRA1. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of GABRA1 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 - 1/100)

Gene Symbol GABRA1

Alternative Names Gamma-aminobutyric acid receptor subunit alpha-1; GABA(A) receptor subunit

alpha-1

**Entrez Gene** 2554 (Human); 14394 (Mouse); 29705 (Rat)

SwissProt P14867 (Human); P62812 (Mouse); P62813 (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

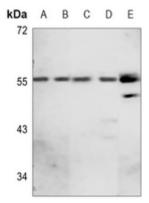
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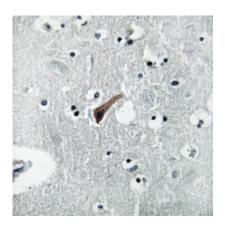




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Western blot analysis of GABRA1 expression in HEK293T (A), HepG2 (B), mouse kidney (C), mouse liver (D), rat liver (E) whole cell lysates. (Predicted band size: 51 kD; Observed band size: 55 kD)



Immunohistochemical analysis of GABRA1 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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