

### **Product Data Sheet**

# **Anti-NBEAL1 Antibody**

Catalog # Source Reactivity Applications

CPA7218 Rabbit H WB, IH

**Description** Rabbit polyclonal antibody to NBEAL1

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

region of human NBEAL1. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of NBEAL1 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 NBEAL1

Alternative Names ALS2CR16; ALS2CR17; Neurobeachin-like protein 1; Amyotrophic lateral sclerosis 2

chromosomal region candidate gene 16 protein; Amyotrophic lateral sclerosis 2

chromosomal region candidate gene 17 protein

Entrez Gene 65065 (Human)

SwissProt Q6ZS30 (Human)

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

freeze/thaw cycles.

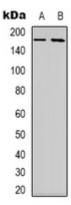
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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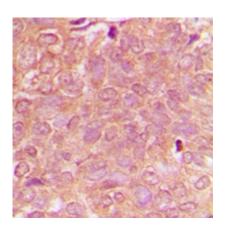
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Western blot analysis of NBEAL1 expression in HEK293T (A), HepG2 (B) whole cell lysates. (Predicted band size: 307 kD; Observed band size: 160 kD)



Immunohistochemical analysis of NBEAL1 staining in human breast 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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