

# **Product Data Sheet**

# **Anti-hnRNP LL Antibody**

Catalog # Source Reactivity Applications

CPA3500 Rabbit H, M, B WB, IH

**Description** Rabbit polyclonal antibody to hnRNP LL

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

region of human hnRNP LL. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of hnRNP LL 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 HNRPLL

Alternative Names SRRF; Heterogeneous nuclear ribonucleoprotein L-like; hnRNPLL; Stromal

RNA-regulating factor

**Entrez Gene** 92906 (Human); 72692 (Mouse)

SwissProt Q8WVV9 (Human); Q921F4 (Mouse)

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

freeze/thaw cycles.

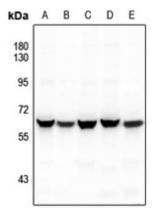
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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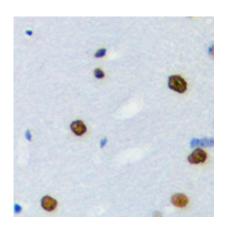
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Western blot analysis of hnRNP LL expression in HCT116 (A), HepG2 (B), MCF7 (C), A549 (D), H1792 (E) whole cell lysates. (Predicted band size: 60 kD; Observed band size: 60 kD)



Immunohistochemical analysis of hnRNP LL 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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