For research purposes only, not for human use

# Product Data Sheet

## Anti-hnRNP R Antibody

| Catalog # | Source | Reactivity | Applications |
|---|---|---|---|
| CQA2007 | Rabbit | H, M, R | WB, IH, IF/IC |

| | |
|---|---|
| **Description** | Rabbit polyclonal antibody to hnRNP R |
| **Immunogen** | KLH-conjugated synthetic peptide of human hnRNP R |
| **Purification** | The antibody was purified by immunogen affinity chromatography. |
| **Specificity** | Recognizes endogenous levels of hnRNP R 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 - 1/2000), IH (1/50 - 1/200), IF/IC (1/50 - 1/200) |
| **Gene Symbol** | HNRNPR |
| **Alternative Names** | HNRPR; Heterogeneous nuclear ribonucleoprotein R; hnRNP R |
| **Entrez Gene** | 10236 (Human) |
| **SwissProt** | O43390 (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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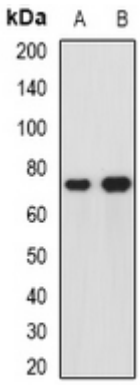
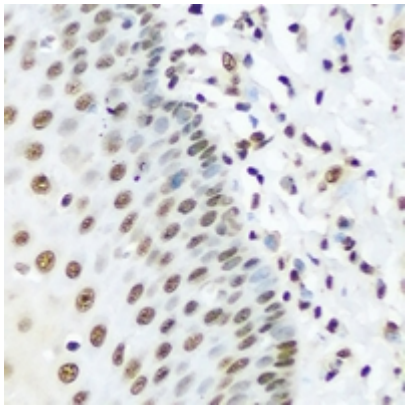
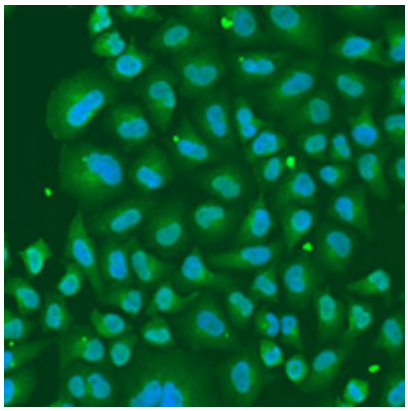



Western blot analysis of hnRNP R expression in Hela (A), HepG2 (B) whole cell lysates. (Predicted band size: 59; 66; 70; 71 kD; Observed band size: 71 kD)



Immunohistochemical analysis of hnRNP R staining in human esophageal 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.



Immunofluorescent analysis of hnRNP R staining in U2OS cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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