

Product Data Sheet

Anti-NRBF2 Antibody

Catalog # Source Reactivity Applications

CQA3180 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to NRBF2

Immunogen Recombinant full length protein of human NRBF2

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of NRBF2 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 NRBF2

Alternative Names COPR; Nuclear receptor-binding factor 2; NRBF-2; Comodulator of PPAR and RXR

Entrez Gene 29982 (Human); 641340 (Mouse); 58839 (Rat)

SwissProt Q96F24 (Human); Q8VCQ3 (Mouse); Q9QYK3 (Rat)

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

freeze/thaw cycles.

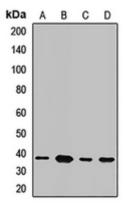
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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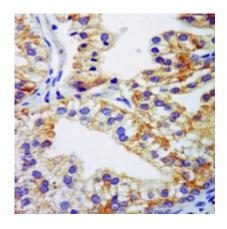
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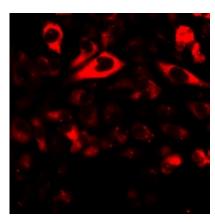
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Western blot analysis of NRBF2 expression in HepG2 (A), Hela (B), mouse heart (C), rat brain (D) whole cell lysates. (Predicted band size: 26; 31; 32 kD; Observed band size: 38 kD)



Immunohistochemical analysis of NRBF2 staining in human liver 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 NRBF2 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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