

Product Data Sheet

Anti-GTF2IRD1 Antibody

Catalog # Source Reactivity Applications

CPA4984 Rabbit H, M WB, IH, IP

Description Rabbit polyclonal antibody to GTF2IRD1

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the N-term

region of human GTF2IRD1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of GTF2IRD1 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), IP (1/10 - 1/100)

Gene Symbol GTF2IRD1

Alternative Names CREAM1; GTF3; MUSTRD1; RBAP2; WBSCR11; WBSCR12; General transcription

factor II-I repeat domain-containing protein 1; GTF2I repeat domain-containing

protein 1; General transcription factor III; MusTRD1/BEN; Muscle TFII-I repeat

domain-containing protein 1; Slow-muscle-fiber enhancer-binding protein; USE

B1-binding protein; Williams-Beuren syndrome chromosomal region 11 protein;

Williams-Beuren syndrome chromosomal region 12 protein

Entrez Gene 9569 (Human); 57080 (Mouse)

SwissProt Q9UHL9 (Human); Q9JI57 (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

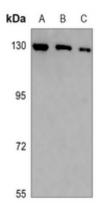
Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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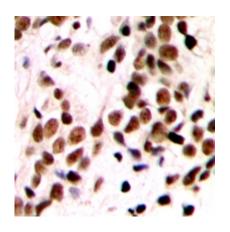
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Western blot analysis of GTF2IRD1 expression in DLD (A), A375 (B), A549 (C) whole cell lysates. (Predicted band size: 106 kD; Observed band size: 130 kD)



Immunohistochemical analysis of GTF2IRD1 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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