

## **Product Data Sheet**

# Anti-TAF15 Antibody

Catalog # Source Reactivity Applications

CPA4982 Rabbit H WB, IH

**Description** Rabbit polyclonal antibody to TAF15

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

region of human TAF15. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of TAF15 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 TAF15

Alternative Names RBP56; TAF2N; TATA-binding protein-associated factor 2N; 68 kDa TATA-binding

protein-associated factor; TAF(II)68; TAFII68; RNA-binding protein 56

Entrez Gene 8148 (Human)

SwissProt Q92804 (Human)

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

freeze/thaw cycles.

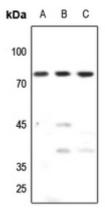
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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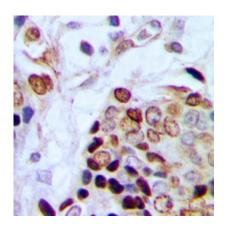
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Western blot analysis of TAF15 expression in HEK293T (A), Hela (B), HGC27 (C) whole cell lysates. (Predicted band size: 61 kD; Observed band size: 75 kD)



Immunohistochemical analysis of TAF15 staining in human prostate 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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