

### **Product Data Sheet**

# **Anti-C1QBP Antibody**

Catalog # Source Reactivity Applications

CPA1115 Rabbit H, M, R WB, IH

**Description** Rabbit polyclonal antibody to C1QBP

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

region of human C1QBP. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of C1QBP 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 (1/50 - 1/100)

Gene Symbol C1QBP

Alternative Names GC1QBP; HABP1; SF2P32; Complement component 1 Q subcomponent-binding

protein mitochondrial; ASF/SF2-associated protein p32; Glycoprotein gC1qBP;

C1qBP; Hyaluronan-binding protein 1; Mitochondrial matrix protein p32; gC1q-R

protein; p33

Entrez Gene 708 (Human)

SwissProt Q07021 (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

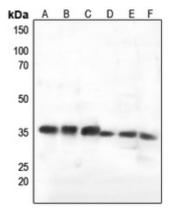
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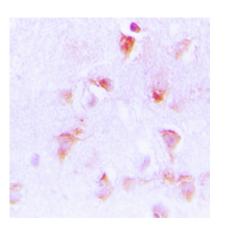




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Western blot analysis of C1QBP expression in HEK293T (A), Hela (B), HepG2 (C), mouse liver (D), mouse brain (E), rat liver (F) whole cell lysates. (Predicted band size: 31 kD; Observed band size: 35 kD)



Immunohistochemical analysis of C1QBP 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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