

Product Data Sheet

Anti-HSP47 Antibody

Catalog # Source Reactivity Applications

CQA2228 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to HSP47

Immunogen Recombinant full length protein of human HSP47

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of HSP47 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 SERPINH1

Alternative Names CBP1; CBP2; HSP47; SERPINH2; Serpin H1; 47 kDa heat shock protein;

Arsenic-transactivated protein 3; AsTP3; Cell proliferation-inducing gene 14 protein;

Collagen-binding protein; Colligin; Rheumatoid arthritis-related antigen RA-A47

Entrez Gene 871 (Human); 12406 (Mouse); 29345 (Rat)

SwissProt P50454 (Human); P19324 (Mouse); P29457 (Rat)

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

freeze/thaw cycles.

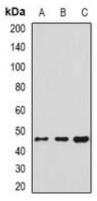
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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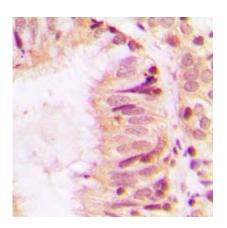
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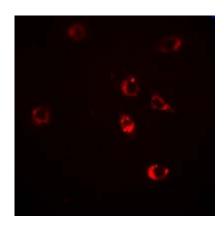
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Western blot analysis of HSP47 expression in Hela (A), U251 (B), THP1 (C) whole cell lysates.



Immunohistochemical analysis of HSP47 staining in human colon 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 HSP47 staining in Hela 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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