

Product Data Sheet

Anti-HSP27 Antibody

| Catalog # | Source | e Reactivity | Applications | |
|-------------------|--------|---|--|--|
| CPA9191 | Mouse | e H | WB, IH | |
| Description | | Mouse monoclonal antiboo | ly to HSP27 | |
| Immunogen | | Recombinant protein corre | sponding to human HSP27. | |
| Purification | | | | |
| Specificity | | Recognizes endogenous lev | els of HSP27 protein. | |
| Clonality | | Monoclonal | | |
| Conjugation | | | | |
| Form | | Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, | | |
| | | and 0.01% sodium azide. | | |
| Dilution | | WB (1/1000 - 1/2000), IH (1/ | (100 - 1/200) | |
| Gene Symbol | | HSPB1 | | |
| Alternative Names | | HSP27; HSP28; Heat shock protein beta-1; HspB1; 28 kDa heat shock protein; | | |
| | | Estrogen-regulated 24 kDa | protein; Heat shock 27 kDa protein; HSP 27; | |
| | | Stress-responsive protein 2 | 7; SRP27 | |
| Entrez Gene | | 3315 (Human) | | |
| SwissProt | | P04792 (Human) | | |
| Storage/Stabi | ility | Shipped at 4°C. Upon delive | ery 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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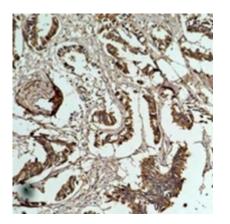
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Western blot analysis of HSP27 expression in Hela (A) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 27 kD)



Immunohistochemical analysis of HSP27 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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