

Anti-HSP20 (pS16) Antibody

Catalog # AP53928

Product Information

Application WB, IF Primary Accession 014558

Reactivity Human, Mouse, Rat

HostRabbitClonalityPolyclonalCalculated MW17136

Additional Information

Gene ID 126393

Other Names Heat shock protein beta-6; HspB6; Heat shock 20 kDa-like protein p20

Target/Specificity KLH-conjugated synthetic peptide encompassing a sequence within the

N-term region of human HSP20. The exact sequence is proprietary.

Dilution WB~~1/500 - 1/2000 IF~~1/50 - 1/100

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name HSPB6

Function Small heat shock protein which functions as a molecular chaperone

probably maintaining denatured proteins in a folding- competent state. Seems to have versatile functions in various biological processes. Plays a role in regulating muscle function such as smooth muscle vasorelaxation and cardiac myocyte contractility. May regulate myocardial angiogenesis

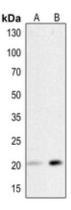
implicating KDR. Overexpression mediates cardioprotection and angiogenesis after induced damage. Stabilizes monomeric YWHAZ thereby supporting

YWHAZ chaperone-like activity.

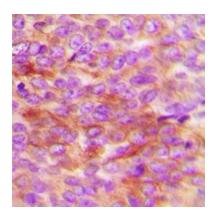
Cellular Location Cytoplasm. Nucleus. Secreted Note=Translocates to nuclear foci during heat

shock

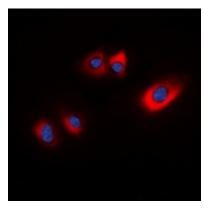
Background



Western blot analysis of HSP20 (pS16) expression in H460 (A), rat heart (B) whole cell lysates.



Immunohistochemical analysis of HSP20 (pS16) 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.



Immunofluorescent analysis of HSP20 (pS16) staining in HEK293T 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.