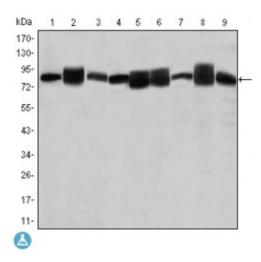


Anti-HSP9 beta antibody



Description Mouse monoclonal to HSP90beta.

Model STJ98144

Host Mouse

Reactivity Human, Mouse, Rat, Simian

Applications ELISA, FC, IF, IHC, WB

Immunogen Purified recombinant fragment of human HSP90beta expressed in E. Coli.

Gene ID 3326

Gene Symbol HSP90AB1

Dilution range WB 1:500-1:2000IHC 1:200-1:1000IF 1:200-1:1000FC 1:200-1:400ELISA

1:10000

Specificity HSP90beta Monoclonal Antibody detects endogenous levels of HSP90beta

protein.

Purification Affinity purification

Clone ID 5G4

Note For Research Use Only (RUO).

Protein Name Heat shock protein HSP 90-beta HSP 90 Heat shock 84 kDa HSP 84 HSP84

Clonality Monoclonal

Conjugation Unconjugated

Isotype IgG1

Formulation Ascitic fluid containing 0.03% sodium azide.

Storage Instruction Store at -20°C, and avoid repeat freeze-thaw cycles.

Database Links <u>HGNC:5258OMIM:140572</u>

Alternative Names Heat shock protein HSP 90-beta HSP 90 Heat shock 84 kDa HSP 84 HSP84

Function

Molecular chaperone that promotes the maturation, structural maintenance and proper regulation of specific target proteins involved for instance in cell cycle control and signal transduction. Undergoes a functional cycle that is linked to its ATPase activity. This cycle probably induces conformational changes in the client proteins, thereby causing their activation. Interacts dynamically with various co-chaperones that modulate its substrate recognition, ATPase cycle and chaperone function. Engages with a range of client protein classes via its interaction with various co-chaperone proteins or complexes, that act as adapters, simultaneously able to interact with the specific client and the central chaperone itself. Recruitment of ATP and co-chaperone followed by client protein forms a functional chaperone. After the completion of the chaperoning process, properly folded client protein and co-chaperone leave HSP90 in an ADP-bound partially open conformation and finally, ADP is released from HSP90 which acquires an open conformation for the next cycle . Apart from its chaperone activity, it also plays a role in the regulation of the transcription machinery. HSP90 and its co-chaperones modulate transcription at least at three different levels. In the first place, they alter the steady-state levels of certain transcription factors in response to various physiological cues. Second, they modulate the activity of certain epigenetic modifiers, such as histone deacetylases or DNA methyl transferases, and thereby respond to the change in the environment. Third, they participate in the eviction of histones from the promoter region of certain genes and thereby turn on gene expression. Antagonizes STUB1-mediated inhibition of TGF-beta signaling via inhibition of STUB1-mediated SMAD3 ubiquitination and degradation. Promotes cell differentiation by chaperoning BIRC2 and thereby protecting from auto-ubiquitination and degradation by the proteasomal machinery. Main chaperone that is involved in the phosphorylation/activation of the STAT1 by chaperoning both JAK2 and PRKCE under heat shock and in turn, activates its own transcription.

Sequence and Domain Family

The TPR repeat-binding motif mediates interaction with TPR repeatcontaining proteins.

Cellular Localization

Cytoplasm Melanosome Nucleus Secreted Cell membrane. Identified by mass spectrometry in melanosome fractions from stage I to stage IV . Translocates with BIRC2 from the nucleus to the cytoplasm during differentiation . Secreted when associated with TGFB1 processed form (LAP) .

Post-translational Modifications Ubiquitinated in the presence of STUB1-UBE2D1 complex (in vitro). ISGylated. S-nitrosylated; negatively regulates the ATPase activity. Phosphorylation at Tyr-301 by SRC is induced by lipopolysaccharide. Phosphorylation at Ser-226 and Ser-255 inhibits AHR interaction. Methylated by SMYD2; facilitates dimerization and chaperone complex formation; promotes cancer cell proliferation. Cleaved following oxidative stress resulting in HSP90AB1 protein radicals formation; disrupts the chaperoning function and the degradation of its client proteins.

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