

### Goat Anti-HSPA8 (Isoform 1) Antibody

Peptide-affinity purified goat antibody Catalog # AF1544a

### **Specification**

### Goat Anti-HSPA8 (Isoform 1) Antibody - Product Information

Application WB
Primary Accession P11142

Other Accession NP 006588, 3312, 15481 (mouse), 24468 (rat)

Reactivity Human

Predicted Mouse, Rat, Dog

Host Goat
Clonality Polyclonal
Concentration 100ug/200ul

Isotype IgG Calculated MW 70898

# Goat Anti-HSPA8 (Isoform 1) Antibody - Additional Information

### **Gene ID 3312**

### **Other Names**

Heat shock cognate 71 kDa protein, Heat shock 70 kDa protein 8, Lipopolysaccharide-associated protein 1, LAP-1, LPS-associated protein 1, HSPA8, HSC70, HSP73, HSPA10

#### **Format**

0.5~mg lgG/ml in Tris saline (20mM Tris pH7.3, 150mM NaCl), 0.02% sodium azide, with 0.5% bovine serum albumin

### **Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

### **Precautions**

Goat Anti-HSPA8 (Isoform 1) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

## Goat Anti-HSPA8 (Isoform 1) Antibody - Protein Information

#### Name HSPA8 (HGNC:5241)

#### **Function**

Molecular chaperone implicated in a wide variety of cellular processes, including protection of the proteome from stress, folding and transport of newly synthesized polypeptides, chaperone-mediated autophagy, activation of proteolysis of misfolded proteins, formation and dissociation of protein complexes, and antigen presentation. Plays a pivotal role in the protein quality control system, ensuring the correct folding of proteins, the re-folding of misfolded proteins



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and controlling the targeting of proteins for subsequent degradation (PubMed:<a
href="http://www.uniprot.org/citations/21148293" target=" blank">21148293</a>, PubMed:<a
href="http://www.uniprot.org/citations/21150129" target="blank">21150129</a>, PubMed:<a
href="http://www.uniprot.org/citations/23018488" target="_blank">23018488</a>, PubMed:<a
href="http://www.uniprot.org/citations/24732912" target="blank">24732912</a>, PubMed:<a
href="http://www.uniprot.org/citations/27916661" target="blank">27916661</a>, PubMed:<a
href="http://www.uniprot.org/citations/2799391" target=" blank">2799391</a>, PubMed:<a
href="http://www.uniprot.org/citations/36586411" target=" blank">36586411</a>). This is
achieved through cycles of ATP binding, ATP hydrolysis and ADP release, mediated by
co-chaperones (PubMed: <a href="http://www.uniprot.org/citations/12526792"
target=" blank">12526792</a>, PubMed:<a href="http://www.uniprot.org/citations/21148293"
target="blank">21148293</a>, PubMed:<a href="http://www.uniprot.org/citations/21150129"
target="blank">21150129</a>, PubMed:<a href="http://www.uniprot.org/citations/23018488"
target="blank">23018488</a>, PubMed:<a href="http://www.uniprot.org/citations/24732912"
target="blank">24732912</a>, PubMed:<a href="http://www.uniprot.org/citations/27916661"
target="blank">27916661</a>). The co-chaperones have been shown to not only regulate
different steps of the ATPase cycle of HSP70, but they also have an individual specificity such that
one co-chaperone may promote folding of a substrate while another may promote degradation
(PubMed:<a href="http://www.uniprot.org/citations/12526792" target=" blank">12526792</a>,
PubMed:<a href="http://www.uniprot.org/citations/21148293" target=" blank">21148293</a>,
PubMed:<a href="http://www.uniprot.org/citations/21150129" target=" blank">21150129</a>,
PubMed: <a href="http://www.uniprot.org/citations/23018488" target="blank">23018488</a>,
PubMed:<a href="http://www.uniprot.org/citations/24732912" target="_blank">24732912</a>,
PubMed: <a href="http://www.uniprot.org/citations/27916661" target="blank">27916661</a>).
The affinity of HSP70 for polypeptides is regulated by its nucleotide bound state. In the ATP-bound
form, it has a low affinity for substrate proteins. However, upon hydrolysis of the ATP to ADP, it
undergoes a conformational change that increases its affinity for substrate proteins. HSP70 goes
through repeated cycles of ATP hydrolysis and nucleotide exchange, which permits cycles of
substrate binding and release. The HSP70-associated co-chaperones are of three types: J- domain
co-chaperones HSP40s (stimulate ATPase hydrolysis by HSP70), the nucleotide exchange factors
(NEF) such as BAG1/2/3 (facilitate conversion of HSP70 from the ADP-bound to the ATP-bound
state thereby promoting substrate release), and the TPR domain chaperones such as HOPX and
STUB1 (PubMed: <a href="http://www.uniprot.org/citations/24121476"
target=" blank">24121476</a>, PubMed:<a href="http://www.uniprot.org/citations/24318877"
target="blank">24318877</a>, PubMed:<a href="http://www.uniprot.org/citations/26865365"
target=" blank">26865365</a>, PubMed:<a href="http://www.uniprot.org/citations/27474739"
target="_blank">27474739</a>). Plays a critical role in mitochondrial import, delivers preproteins
to the mitochondrial import receptor TOMM70 (PubMed: <a
href="http://www.uniprot.org/citations/12526792" target="_blank">12526792</a>). Acts as a
repressor of transcriptional activation. Inhibits the transcriptional coactivator activity of CITED1 on
Smad- mediated transcription. Component of the PRP19-CDC5L complex that forms an integral
part of the spliceosome and is required for activating pre- mRNA splicing. May have a scaffolding
role in the spliceosome assembly as it contacts all other components of the core complex. Binds
bacterial lipopolysaccharide (LPS) and mediates LPS-induced inflammatory response, including
TNF secretion by monocytes (PubMed: <a href="http://www.uniprot.org/citations/10722728"
target=" blank">10722728</a>, PubMed:<a href="http://www.uniprot.org/citations/11276205"
target=" blank">11276205</a>). Substrate recognition component in chaperone-mediated
autophagy (CMA), a selective protein degradation process that mediates degradation of proteins
with a -KFERQ motif: HSPA8/HSC70 specifically recognizes and binds cytosolic proteins bearing a
-KFERQ motif and promotes their recruitment to the surface of the lysosome where they bind to
lysosomal protein LAMP2 (PubMed: <a href="http://www.uniprot.org/citations/11559757"
target=" blank">11559757</a>, PubMed:<a href="http://www.uniprot.org/citations/2799391"
target="blank">2799391</a>, PubMed:<a href="http://www.uniprot.org/citations/36586411"
target=" blank">36586411</a>). KFERQ motif- containing proteins are eventually transported
into the lysosomal lumen where they are degraded (PubMed:<a
href="http://www.uniprot.org/citations/11559757" target=" blank">11559757</a>, PubMed:<a
href="http://www.uniprot.org/citations/2799391" target="_blank">2799391</a>, PubMed:<a
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href="http://www.uniprot.org/citations/36586411" target="\_blank">36586411</a>). In conjunction with LAMP2, facilitates MHC class II presentation of cytoplasmic antigens by guiding antigens to the lysosomal membrane for interaction with LAMP2 which then elicits MHC class II presentation of peptides to the cell membrane (PubMed:<a

href="http://www.uniprot.org/citations/15894275" target="\_blank">15894275</a>). Participates in the ER-associated degradation (ERAD) quality control pathway in conjunction with J domain-containing co- chaperones and the E3 ligase STUB1 (PubMed:<a

href="http://www.uniprot.org/citations/23990462" target="\_blank">23990462</a>). It is recruited to clathrin-coated vesicles through its interaction with DNAJC6 leading to activation of HSPA8/HSC70 ATPase activity and therefore uncoating of clathrin-coated vesicles (By similarity).

#### **Cellular Location**

Cytoplasm. Melanosome. Nucleus, nucleolus. Cell membrane. Lysosome membrane; Peripheral membrane protein; Cytoplasmic side. Note=Localized in cytoplasmic mRNP granules containing untranslated mRNAs (PubMed:17289661). Translocates rapidly from the cytoplasm to the nuclei, and especially to the nucleoli, upon heat shock (PubMed:1586970)

**Tissue Location** Ubiquitous..

### Goat Anti-HSPA8 (Isoform 1) Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

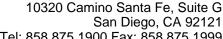
- Western Blot
- Blocking Peptides
- Dot Blot
- <u>Immunohistochemistry</u>
- Immunofluorescence
- Immunoprecipitation
- Flow Cvtometv
- Cell Culture

## Goat Anti-HSPA8 (Isoform 1) Antibody - Images



AF1544a (2  $\mu$ g/ml) staining of A431 cell lysate (35  $\mu$ g protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

## Goat Anti-HSPA8 (Isoform 1) Antibody - Background





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The product encoded by this gene belongs to the heat shock protein 70 family which contains both heat-inducible and constitutively expressed members. The latter are called heat-shock cognate proteins. This gene encodes a heat-shock cognate protein. This protein binds to nascent polypeptides to facilitate correct folding. It also functions as an ATPase in the disassembly of clathrin-coated vesicles during transport of membrane components through the cell. Two alternatively spliced variants have been characterized to date.

## Goat Anti-HSPA8 (Isoform 1) Antibody - References

Variation at the NFATC2 Locus Increases the Risk of Thiazolinedinedione-Induced Edema in the Diabetes REduction Assessment with ramipril and rosiglitazone Medication (DREAM) Study. Bailey SD, et al. Diabetes Care, 2010 Jul 13. PMID 20628086.

Proteome analysis of the thalamus and cerebrospinal fluid reveals glycolysis dysfunction and potential biomarkers candidates for schizophrenia. Martins-de-Souza D, et al. J Psychiatr Res, 2010 May 14. PMID 20471030.

Nucleolar targeting of the chaperone hsc70 is regulated by stress, cell signaling, and a composite targeting signal which is controlled by autoinhibition. Ba?ski P, et al. J Biol Chem, 2010 Jul 9. PMID 20457599.

Genetic variations in HSPA8 gene associated with coronary heart disease risk in a Chinese population. He M, et al. PLoS One, 2010 Mar 16. PMID 20300519.

Heat shock cognate protein 70 is essential for Akt signaling in endothelial function. Shiota M, et al. Arterioscler Thromb Vasc Biol, 2010 Mar. PMID 20018937.