

# Hsc70 Rabbit mAb

**Catalog # AP75575** 

# **Specification**

#### Hsc70 Rabbit mAb - Product Information

Application
Primary Accession
Reactivity
Host
Clonality
Calculated MW

WB, IHC-P, IP
P11142
Human, Mouse, Rat, Hamster
Rabbit
Monoclonal Antibody
70898

#### Hsc70 Rabbit mAb - Additional Information

**Gene ID 3312** 

Other Names HSPA8

**Dilution**WB~~1/500-1/1000
IHC-P~~N/A
IP~~1/20

# **Format**

50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40%Glycerol, 0.01% sodium azide and 0.05% BSA.

#### Storage

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.

## Hsc70 Rabbit mAb - Protein Information

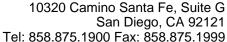
# Name HSPA8 (HGNC:5241)

### **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 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">2799391</a>, PubMed:<a href="http://www.uniprot.org/citations/36586411" target="\_blank">2799391</a>, PubMed:<a href="http://www.uniprot.org/citations/2799391" target="\_blank">2799391</a>, PubMed:<a href="http://www.uniprot.org/citations/2799391" target="\_blank">2799391</a>, PubMed:<a href="http://www.uniprot.org/citations/2799391" target="\_blank">2799391</a>, PubMed:<a hre



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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
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
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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..

# **Hsc70 Rabbit mAb - 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 Cytomety
- Cell Culture

# Hsc70 Rabbit mAb - Images

