

**HSP70 Antibody**  
**HSP70 Antibody, Clone C92F3A-5**  
**Catalog # ASM10000**

**Specification**

---

**HSP70 Antibody - Product Information**

Application	WB, IHC, ICC, E, FC, IEM, BL, AM
Primary Accession	<a href="#">P08107</a>
Other Accession	<a href="#">NP_005336.3</a>
Host	Mouse
Isotype	IgG
Reactivity	Human, Mouse, Rat, Rabbit, Hamster, Monkey, Pig, Chicken, Bovine, C.Elegans, Dog, Sheep, Guinea Pig, Drosophila
Clonality	Monoclonal

**Description**  
Mouse Anti-Human HSP70 Monoclonal IgG

**Target/Specificity**  
Detects ~70kDa. Does not cross-react with HSC70 (HSP73).

**Other Names**  
HSP70 1 Antibody, HSP70 2 Antibody, HSP70.1 Antibody, HSP72 Antibody, HSPA1 Antibody, HSPA1A Antibody, HSPA1B Antibody

**Immunogen**  
Human HSP70

**Purification**  
Protein G Purified

**Storage** -20°C  
**Storage Buffer**  
PBS pH7.4, 50% glycerol, 0.1% sodium azide

**Shipping Temperature** Blue Ice or 4°C

**Certificate of Analysis**  
1 µg/ml of SMC-100 was sufficient for detection of HSP70 in 20 µg of heat shocked HeLa cell lysate by colorimetric immunoblot analysis using Goat anti-mouse IgG:HRP as the secondary antibody.

**Cellular Localization**  
Cytoplasm

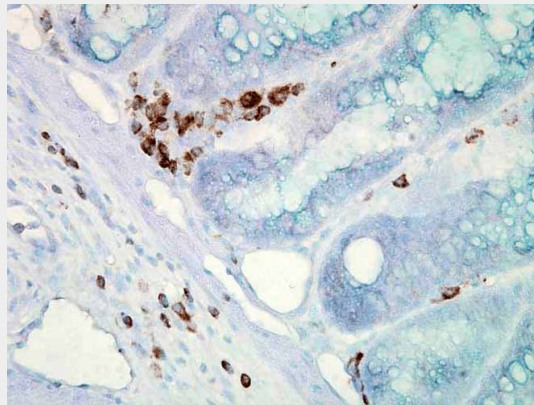
**HSP70 Antibody - Protocols**

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

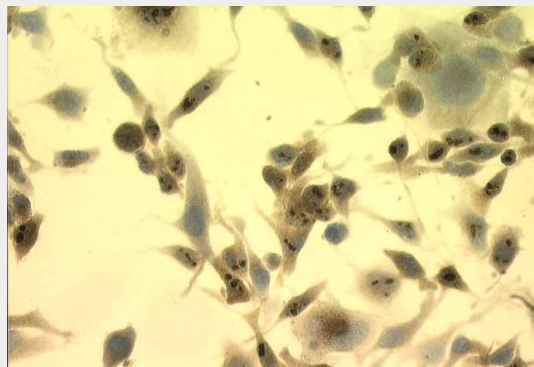
- [Western Blot](#)
- [Blocking Peptides](#)

- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

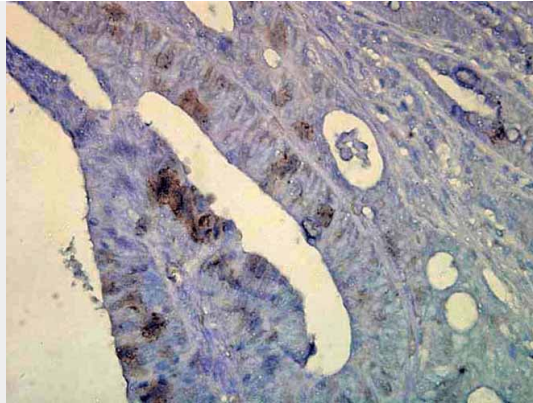
### HSP70 Antibody - Images



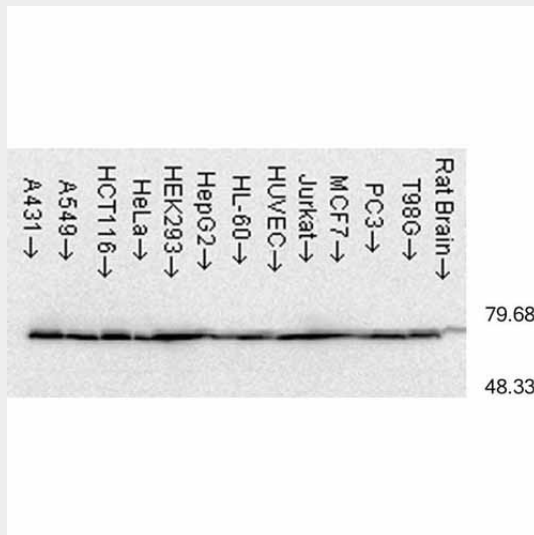
Immunohistochemistry analysis using Mouse Anti-Hsp70 Monoclonal Antibody, Clone C92 (ASM10000). Tissue: colon carcinoma. Species: Mouse. Fixation: Formalin. Primary Antibody: Mouse Anti-Hsp70 Monoclonal Antibody (ASM10000) at 1:10000 for 12 hours at 4°C. Secondary Antibody: Biotin Goat Anti-Mouse at 1:2000 for 1 hour at RT. Counterstain: Mayer Hematoxylin (purple/blue) nuclear stain at 200  $\mu$ l for 2 minutes at RT. Localization: Inflammatory cells. Magnification: 40x.



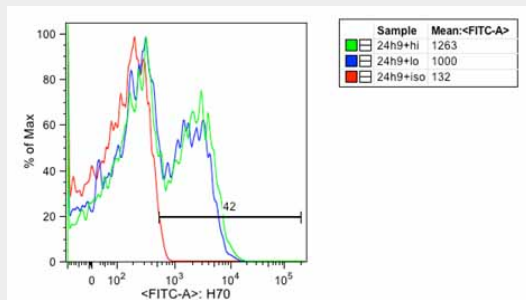
Immunocytochemistry/Immunofluorescence analysis using Mouse Anti-Hsp70 Monoclonal Antibody, Clone C92 (ASM10000). Tissue: Heat Shocked Melanoma cells. Species: Mouse. Fixation: Formalin. Primary Antibody: Mouse Anti-Hsp70 Monoclonal Antibody (ASM10000) at 1:1000 for 16 hours at RT. Secondary Antibody: Biotin Goat Anti-Mouse. Courtesy of: Dr. Ewa Malusecka, Maria Skłodowska-Curie Memorial Cancer Centre and Inst. Of Oncology, Poland.



Immunohistochemistry analysis using Mouse Anti-Hsp70 Monoclonal Antibody, Clone C92 (ASM10000). Tissue: colon carcinoma. Species: Human. Fixation: Formalin. Primary Antibody: Mouse Anti-Hsp70 Monoclonal Antibody (ASM10000) at 1:10000 for 12 hours at 4°C. Secondary Antibody: Biotin Goat Anti-Mouse at 1:2000 for 1 hour at RT. Counterstain: Mayer Hematoxylin (purple/blue) nuclear stain at 200 µl for 2 minutes at RT. Localization: Inflammatory cells. Magnification: 40x.



Western Blot analysis of Human cell lysates from various cell lines showing detection of Hsp70 protein using Mouse Anti-Hsp70 Monoclonal Antibody, Clone C92 (ASM10000). Load: 15 µg. Block: 1.5% BSA for 30 minutes at RT. Primary Antibody: Mouse Anti-Hsp70 Monoclonal Antibody (ASM10000) at 1:1000 for 2 hours at RT. Secondary Antibody: Sheep Anti-Mouse IgG: HRP for 1 hour at RT.



FACS analysis. Anti-Hsp70-FITC staining on heat shock treated CD3+CD8+ T cells

Fluorescence Activated Cell Sorting analysis using Mouse Anti-Hsp70: FITC Monoclonal Antibody, Clone C92 (ASM10000). Tissue: Heat Shocked CD3+ CD8+ T cells . Species: Mouse. Primary Antibody: Mouse Anti-Hsp70: FITC Monoclonal Antibody (ASM10000) at 1:1000. Courtesy of: Cheryl Cameron, Vaccine and Gene Therapy Instit. Florida.

### **HSP70 Antibody - Background**

HSP70 genes encode abundant heat-inducible 70-kDa HSPs (HSP70s). In most eukaryotes HSP70 genes exist as part of a multigene family. They are found in most cellular compartments of eukaryotes including nuclei, mitochondria, chloroplasts, the endoplasmic reticulum and the cytosol, as well as in bacteria. The genes show a high degree of conservation, having at least 50% identity (2). The N-terminal two thirds of HSP70s are more conserved than the C-terminal third. HSP70 binds ATP with high affinity and possesses a weak ATPase activity which can be stimulated by binding to unfolded proteins and synthetic peptides (3). When HSC70 (constitutively expressed) present in mammalian cells was truncated, ATP binding activity was found to reside in an N-terminal fragment of 44 kDa which lacked peptide binding capacity. Polypeptide binding ability therefore resided within the C-terminal half (4). The structure of this ATP binding domain displays multiple features of nucleotide binding proteins (5).

All HSP70s, regardless of location, bind proteins, particularly unfolded ones. The molecular chaperones of the HSP70 family recognize and bind to nascent polypeptide chains as well as partially folded intermediates of proteins preventing their aggregation and misfolding. The binding of ATP triggers a critical conformational change leading to the release of the bound substrate protein (6). The universal ability of HSP70s to undergo cycles of binding to and release from hydrophobic stretches of partially unfolded proteins determines their role in a great variety of vital intracellular functions such as protein synthesis, protein folding and oligomerization and protein transport. For more information visit our HSP70 Scientific Resource Guide at <http://www.HSP70.com>.

### **HSP70 Antibody - References**

1. Welch W.J. and Suhan J.P. (1986) *J Cell Biol.* 103: 2035-2050.
2. Boorstein W. R., Ziegelhoffer T. & Craig E. A. (1993) *J.Mol. Evol.* 38(1): 1-17.
3. Rothman J. (1989) *Cell* 59: 591-601.
4. DeLuca-Flaherty et al. (1990) *Cell* 62: 875-887.
5. Bork P., Sander C. & Valencia A. (1992) *Proc. Natl Acad. Sci. USA* 89: 7290-7294.
6. Fink A.L. (1999) *Physiol. Rev.* 79: 425-449.
7. Galan A., et al. (2000) *J. Biol. Chem.* 275: 11418-11424.
8. Kondo T., et al. (2000) *J. Biol. Chem.* 275: 8872-8879.
9. Misaki T., et al. (1994) *Clin. Exp. Immun.* 98: 234-239.
10. Pockley A.G., et al. (1998) *Immunol. Invest.* 27: 367-377.
11. Moon I.S., et al. (2001) *Cereb Cortex* 11(3): 238-248.
12. Dressel et al. (2000) *J. Immunol.* 164: 2362-2371.
13. Verma A.K., et al. (2007) *Fish and Shellfish Immunology.* 22(5): 547-555.
14. Banduseela V.C., et al. (2009) *Physiol Genomics.* 39(3): 141-159.