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# **Hsc70 Protein (His tag)**



# **Publications**



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Quantity:	100 μg	
Target:	Hsc70 (HSPA8)	
Origin:	Human	
Source:	Escherichia coli (E. coli)	
Protein Type:	Recombinant	
Biological Activity:	Active	
Purification tag / Conjugate:	This Hsc70 protein is labelled with His tag.	
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA, Functional Studies (Func), Activity Assay (AcA)	
Product Details		
Charificity		
Specificity:	~70 kDa	
Specificity:  Characteristics:	~/0 kDa  The protein has ATPase activity at the time of manufacture of 3.2 μM phosphate liberated/hr/μ	
	The protein has ATPase activity at the time of manufacture of 3.2 μM phosphate liberated/hr/μ	
	The protein has ATPase activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ L reaction at 37 °C (pH 8) in the presence of 20 $\mu$ L of 1 mM ATP using a	
Characteristics:	The protein has ATPase activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ L reaction at 37 °C (pH 8) in the presence of 20 $\mu$ L of 1 mM ATP using a Malachite Green assay.	
Characteristics:  Purification:	The protein has ATPase activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ L reaction at 37 °C (pH 8) in the presence of 20 $\mu$ L of 1 mM ATP using a Malachite Green assay.  Affinity Purified	
Characteristics:  Purification:  Purity:	The protein has ATPase activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ L reaction at 37 °C (pH 8) in the presence of 20 $\mu$ L of 1 mM ATP using a Malachite Green assay.  Affinity Purified  >90%	
Characteristics:  Purification:  Purity:  Biological Activity Comment:	The protein has ATPase activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ L reaction at 37 °C (pH 8) in the presence of 20 $\mu$ L of 1 mM ATP using a Malachite Green assay.  Affinity Purified  >90%	

### Target Details

Alternative Name:	Hsc70 (HSPA8 Products)	
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). When cells	
	are subjected to metabolic stress (e.g., heat shock) a member of the HSP 70 family, HSP 70	
	(HSP72), is expressed, HSP 70 is highly related to HSC70 (>90 % sequence identity).	
	Constitutively expressed HSC70 rapidly forms a stable complex with the highly inducible HSP70	
	in cells following heat shock. The interaction of HSC70 with HSP 70 is regulated by ATP. These	
	two heat shock proteins move together in the cell experiencing stress. Furthermore, research	
	on HSC70 has implicates it with a role in facilitating the recovery of centrosomal structure and	
	function after heat shock (6).	
Molecular Weight:	approx. 70 kDa	
Gene ID:	3312	
UniProt:	P11142	
Application Details		
Application Notes:	Optimal working dilution should be determined by the investigator.	
Comment:	This product has been certified >90% pure using SDS-PAGE analysis. The protein has ATPase	
	activity at the time of manufacture of 3.2 $\mu$ M phosphate liberated/hr/ $\mu$ g protein in a 200 $\mu$ l	
	reaction at 37°C (pH 8) in the presence of 20ul of 1mM ATP using a Malachite Green assay.	
Restrictions:	For Research Use only	
Handling		

### Handling

Buffer:	50 mM Tris/HCl, pH 8, 0.3M NaCl	
Storage:	-20 °C	
Publications		
Product cited in:	Kaiser, Steptoe, Thompson, Henderson: "Monocyte cytokine synthesis in response to extracellular cell stress proteins suggests these proteins exhibit network behaviour." in: <b>Cell stress &amp; chaperones</b> , Vol. 19, Issue 1, pp. 135-44, (2013) (PubMed).	

There are more publications referencing this product on: Product page