# HSP90 (C45G5) Rabbit mAb



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# For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, W-S, IHC-P, IF-IC, FC-FP	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 90	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P08238, #P07900	Entrez-Gene Id 3326, 3320	
Product Usage Information		Application Western Blotting			<b>Dilution</b> 1:1000		
		Simple Western™				- 1:50	
		Immunohistochemist	try (Paraffin)			) - 1:800	
		Immunofluorescence	• •	istrv)	1:50		
		Flow Cytometry (Fixed		<b>,</b>	1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.					
		For a carrier free (BSA	A and azide free) ver	sion of this product se	e product #98099.		
Specificity/Sensitivity		HSP90 (C45G5) Rabbit mAb detects endogenous levels of total HSP90 protein. This antibody does not cross-react with other HSPs.					
Species predictories based on 100% homology		Bovine					
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide surrounding Asn300 of human HSP90.					
Background		maintain protein hom are able to interact w refolding of their sub range of substrates in more limited subset of function collaborative HSP40, Hop, and p23 chaperones or recruit ubiquitin ligase CHIP are subjected to degr beyond their chapero	and HSP90 are molecular chaperones expressed constitutively under normal conditions to a a intain protein homeostasis and are induced upon environmental stress (1). Both HSP70 and HSP90 are able to interact with unfolded proteins to prevent irreversible aggregation and catalyze the folding of their substrates in an ATP- and co-chaperone-dependent manner (1). HSP70 has a broading of substrates including newly synthesized and denatured proteins, while HSP90 tends to have a core limited subset of substrates, most of which are signaling molecules. HSP70 and HSP90 often nection collaboratively in a multi-chaperone system, which requires a minimal set of co-chaperones: FP40, Hop, and p23 (2,3). The co-chaperones either regulate the intrinsic ATPase activity of the aperones or recruit chaperones to specific substrates or subcellular compartments (1,4). When the diquitin ligase CHIP associates with the HSP70/HSP90 complex as a cofactor, the unfolded substrates be subjected to degradation by the proteasome (4). The biological functions of HSP70/HSP90 extend degradation and their chaperone activity. They are essential for the maturation and inactivation of nuclear formones and other signaling molecules (1,3). They also play a role in vesicle formation and protein afficking (2).				
Background Re	ferences	<ol> <li>Nollen, E.A. and Morimoto, R.I. (2002) <i>J. Cell Sci.</i> 115, 2809-2816.</li> <li>Young, J.C. et al. (2003) <i>Trends Biochem. Sci.</i> 28, 541-547.</li> <li>Pratt, W.B. and Toft, D.O. (2003) <i>Exp. Biol. Med.</i> 228, 111-133.</li> <li>Hohfeld, J. et al. (2001) <i>EMBO Rep.</i> 2, 885-890.</li> </ol>					
Species Reactiv	itv	Species reactivity is d	etermined by testin	g in at least one appro	ved application (e.g., w	vestern blot).	

## Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

**Western Blot Buffer** 

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at  $4^{\circ}$ C with gentle shaking, overnight.

# **Applications Key**

W: Western Blotting W-S: Simple Western™ IHC-P: Immunohistochemistry (Paraffin) IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** 

H: Human M: Mouse R: Rat Mk: Monkey

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