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Quantity:	100 μg
Target:	HSPB8
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This HSPB8 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

#### **Product Details**

Immunogen:	His-tagged human recombinant HSP22
Clone:	3C12-H11
Isotype:	IgG1 kappa
Specificity:	Detects ~22 kDa. Detects endogenous and exogenous HSP22 in monomeric, dimeric and tetrameric forms in WB. Does not cross react with alpha crystallin.
Cross-Reactivity:	Human, Mouse, Rat
Purification:	Protein G Purified

## **Target Details**

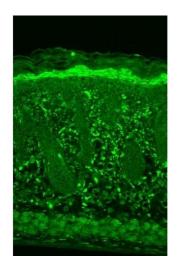
Target:	HSPB8

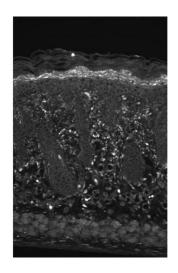
Alternative Name:	HSP22 (HSPB8 Products)
Background:	HSP27s belong to an abundant and ubiquitous family of small heat shock proteins (sHSP). It is
	an important HSP found in both normal human cells and cancer cells. The basic structure of
	most sHSPs is a homologous and highly conserved amino acid sequence, with an $\alpha\mbox{-crystallin}$
	domain at the C-terminus and the WD/EPF domain at the less conserved N-terminus. This N-
	terminus is essential for the development of high molecular oligomers (1, 2). HSP27-oligomers
	consist of stable dimers formed by as many as 8-40 HSP27 protein monomers (3). The
	oligomerization status is connected with the chaperone activity: aggregates of large oligomers
	have high chaperone activity, whereas dimers have no chaperone activity (4). HSP27 is
	localized to the cytoplasm of unstressed cells but can redistribute to the nucleus in response to
	stress, where it may function to stabilize DNA and/or the nuclear membrane. Other functions
	include chaperone activity (as mentioned above), thermo tolerance in vivo, inhibition of
	apoptosis, and signal transduction. Specifically, in vitro, it acts as an ATP independent
	chaperone by inhibiting protein aggregation and by stabilizing partially denatured proteins,
	which ensures refolding of the HSP70 complex. HSP27 is also involved in the apoptotic
	signaling pathway because it interferes with the activation of cytochrome c/Apaf-1/dATP
	complex, thereby inhibiting the activation of procaspase-9. It is also hypothesized that HSP27
	may serve some role in cross-bridge formation between actin and myosin (5). And finally,
	HSP27 is also thought to be involved in the process of cell differentiation. The up-regulation of
	HSP27 correlates with the rate of phosphorylation and with an increase of large oligomers. It is
	possible that HSP27 may play a crucial role in termination of growth (6).
Gene ID:	26353
NCBI Accession:	NP_055180
UniProt:	Q9UJY1
Application Details	
Application Notes:	• WB (1:2000)
	• IHC (1:100)
	optimal dilutions for assays should be determined by the user.
Comment:	1 μg/ml of ABIN863102 was sufficient for detection of HSP22 in 20 μg of whole rat tissue
	extract by ECL immunoblot analysis using Goat anti-mouse IgG:HRP as the secondary
	antibody.
Restrictions:	For Research Use only

#### Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	PBS pH 7.4, 50 % glycerol, 0.09 % sodium azide, Storage buffer may change when conjugated
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	-20°C

# **Images**



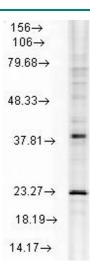


### **Immunohistochemistry**

Image 1. Immunohistochemistry analysis using Mouse Anti-Hsp22 Monoclonal Antibody, Clone 3C12-H11 (ABIN863102 and ABIN863103). Tissue: backskin. Species: Mouse. Fixation: Bouin's Fixative and paraffin-embedded. Primary Antibody: Mouse Anti-Hsp22 Monoclonal Antibody (ABIN863102 and ABIN863103) at 1:100 for 1 hour at RT. Secondary Antibody: FITC Goat Anti-Mouse (green) at 1:50 for 1 hour at RT. Localization: Granular layer of the epidermis. Some dermal staining.

#### **Immunohistochemistry**

Image 2. HSP22 (3C12 H11), Mouse backskin.



#### **Western Blotting**

**Image 3.** Rat tissue Mix WB 1 in 1000 Hsp22 monomer and dimer.