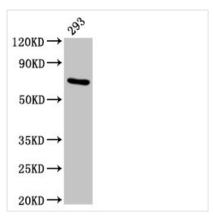




HSPA1A Antibody

Product Code	CSB-PA10899A0Rb
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P0DMV8
Immunogen	Recombinant Human Heat shock 70 kDa protein 1A protein (418-512AA)
Raised In	Rabbit
Species Reactivity	Human
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:200-1:500, IF:1:50-1:200
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Storage Buffer Purification Method	
	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified
Purification Method	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified IgG
Purification Method Isotype Clonality	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified IgG Polyclonal Heat shock 70 kDa protein 1A (Heat shock 70 kDa protein 1) (HSP70-1)
Purification Method Isotype Clonality Alias	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified IgG Polyclonal Heat shock 70 kDa protein 1A (Heat shock 70 kDa protein 1) (HSP70-1) (HSP70.1), HSPA1A, HSP72 HSPA1 HSX70
Purification Method Isotype Clonality Alias Immunogen Species	Constituents: 50% Glycerol, 0.01M PBS, pH 7.4 >95%, Protein G purified IgG Polyclonal Heat shock 70 kDa protein 1A (Heat shock 70 kDa protein 1) (HSP70-1) (HSP70.1), HSPA1A, HSP72 HSPA1 HSX70 Homo sapiens (Human)





Positive WB detected in: 293 whole cell lysate All lanes: HSPA1A antibody at 4.9µg/ml

Goat polyclonal to rabbit IgG at 1/50000 dilution

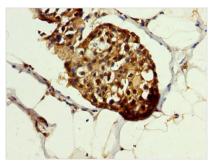
Predicted band size: 71, 64 kDa Observed band size: 71 kDa



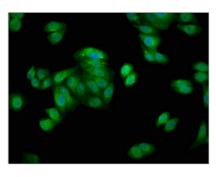








IHC image of CSB-PA10899A0Rb diluted at 1:200 and staining in paraffin-embedded human breast cancer performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescence staining of HepG2 cells with CSB-PA10899A0Rb at 1:66, counterstained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).