

## HSPB8 Ab

[Images\(3\)](#)

|               |                 |                       |
|---------------|-----------------|-----------------------|
| Cat.#: DF2481 | Concn.: ~1mg/ml | Mol.Wt.: 22 kDa       |
| Size:         | Source: Rabbit  | Clonality: Polyclonal |

**Application:** WB 1:500-1:2000, IF/ICC 1:100-1:500  
\*The optimal dilutions should be determined by the end user.

**Reactivity:** Human, Mouse, Rat

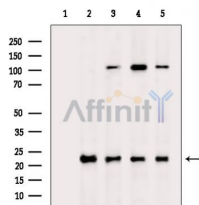
**Storage:** Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20 °C. Stable for 12 months from date of receipt.

**Purification:** The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).

**Immunogen:** A synthesized peptide derived from human HSPB8, corresponding to a region within C-terminal amino acids.

**Uniprot:** Q9UJY1

**Description:** Displays temperature-dependent chaperone activity.



Western blot analysis of extracts from various samples, using HSPB8 Ab.

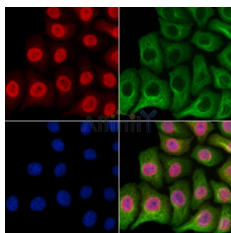
Lane 1: Rat liver, blocked with antigen-specific peptides.

Lane 2: Rat liver.

Lane 3: Mouse brain.

Lane 4: Hepg2 cells (heat-shock treatment).

Lane 5: Hela cells (heat-shock treatment).



DF2481 staining Hela cells by IF/ICC. The samples were fixed with PFA and permeabilized in 0.1% Triton X-100, then blocked in 10% serum for 45 minutes at 25°C. Samples were then incubated with primary Ab (DF2481) and mouse anti-beta tubulin Ab (T0023) for 1 hour at 37°C. An AlexaFluor594 conjugated goat anti-rabbit IgG (H+L) Ab (Red) and an AlexaFluor488 conjugated goat anti-mouse IgG (H+L) Ab (Green) were used as the secondary Ab.

The nuclear counter stain is DAPI (blue).

**IMPORTANT:** For western blot, incubate membrane with diluted primary Ab in 5% w/v milk, 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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