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## [KD Validated] HSP27/HSPB1 Rabbit mAb

Catalog No.: A27821 Recombinant

## **Basic Information**

## **Observed MW**

27kDa

## **Calculated MW**

23kDa

## Category

Primary antibody

## **Applications**

WB,IP,IF/ICC,IHC-P,ELISA

#### **Cross-Reactivity**

Human

#### CloneNo number

ARC75566

## **Background**

This gene encodes a member of the small heat shock protein (HSP20) family of proteins. In response to environmental stress, the encoded protein translocates from the cytoplasm to the nucleus and functions as a molecular chaperone that promotes the correct folding of other proteins. This protein plays an important role in the differentiation of a wide variety of cell types. Expression of this gene is correlated with poor clinical outcome in multiple human cancers, and the encoded protein may promote cancer cell proliferation and metastasis, while protecting cancer cells from apoptosis. Mutations in this gene have been identified in human patients with Charcot-Marie-Tooth disease and distal hereditary motor neuropathy.

## **Recommended Dilutions**

**WB** 1:10000 - 1:40000

IP 0.5μg-4μg antibody for 200μg-400μg extracts of

whole cells

IF/ICC 1:100 - 1:200

**IHC-P** 1:500 - 1:2000

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

## **Contact**

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## Immunogen Information

**Gene ID**3315

Swiss Prot
P04792

#### **Immunogen**

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

## Synonyms

CMT2F; HMN2B; HSP27; HSP28; Hsp25; SRP27; HS.76067; HEL-S-102

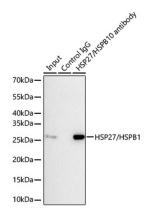
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

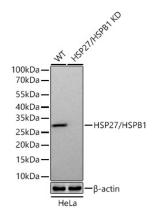
#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Immunoprecipitation of HSP27/HSPB1 from 300  $\mu$ g extracts of HeLa cells was performed using 2  $\mu$ g of [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821). Rabbit IgG isotype control (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1X reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821) at a dilution of 1:5000.

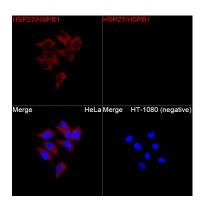


Western blot analysis of lysates from wild type (WT) and HSP27/HSPB1 knockdown (KD) HeLa cells using [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821) at 1:13000 dilution incubated overnight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25  $\mu$ g per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

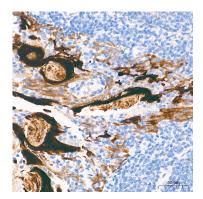
Exposure time: 1s.



Immunofluorescence analysis of HeLa cells and HT-1080 (negative) cells using [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821) at a dilution of 1:200 (40x lens). Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L)(AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.



Immunohistochemistry analysis of paraffinembedded Human cervix tissue using [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821) at a dilution of 1:600 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Human tonsil tissue using [KD Validated] HSP27/HSPB1 Rabbit mAb (A27821) at a dilution of 1:600 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.