

Product datasheet

Anti-Hsp27 antibody ab26942

1 References 1 图像

概述

产品名称	Anti-Hsp27抗体
描述	鸡多克隆抗体to Hsp27
宿主	Chicken
经测试应用	适用于: WB
种属反应性	与反应: Mouse, Cow, Human
免疫原	Fusion protein: PPGVDPTQVS SSSLSPGTLT VEAPMPKLAT QSNEITIPVT FESRAQLGGP K, corresponding to amino acids 145-205 of Human Hsp27 Run BLAST with ExPASy Run BLAST with NCBI
阳性对照	Oral cancer epithelium.

性能

形式	Liquid
存放说明	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term.
存储溶液	Preservative: None Constituents: PBS
纯度	Protein L purified
克隆	多克隆
同种型	IgY

应用

Our [Abpromise guarantee](#) covers the use of **ab26942** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应用	Ab评论	说明
WB		1/3000. Predicted molecular weight: 23 kDa.

靶标

功能

Involved in stress resistance and actin organization.

组织特异性

Detected in all tissues tested: skeletal muscle, heart, aorta, large intestine, small intestine, stomach, esophagus, bladder, adrenal gland, thyroid, pancreas, testis, adipose tissue, kidney, liver, spleen, cerebral cortex, blood serum and cerebrospinal fluid. Highest levels are found in the heart and in tissues composed of striated and smooth muscle.

疾病相关

Defects in HSPB1 are the cause of Charcot-Marie-Tooth disease type 2F (CMT2F) [MIM:606595]. CMT2F is a form of Charcot-Marie-Tooth disease, the most common inherited disorder of the peripheral nervous system. Charcot-Marie-Tooth disease is classified in two main groups on the basis of electrophysiologic properties and histopathology: primary peripheral demyelinating neuropathy or CMT1, and primary peripheral axonal neuropathy or CMT2. Neuropathies of the CMT2 group are characterized by signs of axonal regeneration in the absence of obvious myelin alterations, normal or slightly reduced nerve conduction velocities, and progressive distal muscle weakness and atrophy. Nerve conduction velocities are normal or slightly reduced. CMT2F onset is between 15 and 25 years with muscle weakness and atrophy usually beginning in feet and legs (peroneal distribution). Upper limb involvement occurs later. CMT2F inheritance is autosomal dominant.

Defects in HSPB1 are a cause of distal hereditary motor neuronopathy type 2B (HMN2B) [MIM:608634]. Distal hereditary motor neuronopathies constitute a heterogeneous group of neuromuscular disorders caused by selective impairment of motor neurons in the anterior horn of the spinal cord, without sensory deficit in the posterior horn. The overall clinical picture consists of a classical distal muscular atrophy syndrome in the legs without clinical sensory loss. The disease starts with weakness and wasting of distal muscles of the anterior tibial and peroneal compartments of the legs. Later on, weakness and atrophy may expand to the proximal muscles of the lower limbs and/or to the distal upper limbs.

序列相似性

Belongs to the small heat shock protein (HSP20) family.

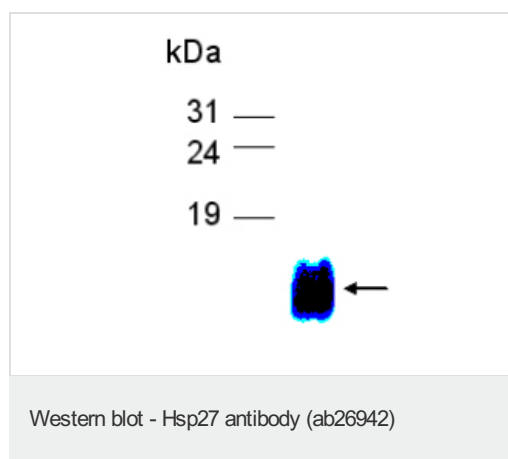
翻译后修饰

Phosphorylated in MCF-7 cells on exposure to protein kinase C activators and heat shock.

细胞定位

Cytoplasm. Nucleus. Cytoplasm > cytoskeleton > spindle. Cytoplasmic in interphase cells. Colocalizes with mitotic spindles in mitotic cells. Translocates to the nucleus during heat shock and resides in sub-nuclear structures known as SC35 speckles or nuclear splicing speckles.

图片



Anti-Hsp27 antibody (ab26942) at 1/3000 dilution + Hsp27 fusion protein

Predicted band size: 23 kDa

Western blot - Hsp27 antibody (ab26942)

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