abcam

Product datasheet

Anti-Hsp60 antibody [HSPD1/2206R] - BSA free ab234286

Recombinant

2 Images

Overview

Product name Anti-Hsp60 antibody [HSPD1/2206R] - BSA free

Description Rabbit monoclonal [HSPD1/2206R] to Hsp60 - BSA free

Host species Rabbit

Tested applications
Suitable for: IHC-P
Species reactivity
Reacts with: Human

Predicted to work with: Mouse, Rat, Sheep, Rabbit, Chicken, Hamster, Cow, Dog, Pig,

Xenopus laevis, Drosophila melanogaster, Monkey

Immunogen Recombinant full length protein corresponding to Human Hsp60.

Database link: P10809

Positive control IHC-P: Human liver tissue.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.05% Sodium azide

Constituent: PBS

Purity Protein A/G purified

Purification notes Purified from Bioreactor Concentrate by Protein A/G.

Clonality Monoclonal
Clone number HSPD1/2206R

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab234286 in the following tested applications.

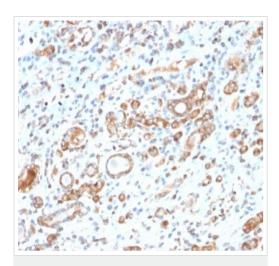
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		Use a concentration of 0.5 - 1 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. (Primary incubation for 10 minutes at room temperature).

Target

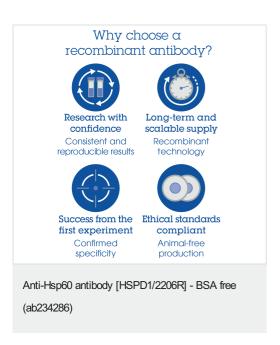
Target		
Function	Implicated in mitochondrial protein import and macromolecular assembly. May facilitate the correct folding of imported proteins. May also prevent misfolding and promote the refolding and proper assembly of unfolded polypeptides generated under stress conditions in the mitochondrial matrix.	
Involvement in disease	Defects in HSPD1 are a cause of spastic paraplegia autosomal dominant type 13 (SPG13) [MIM:605280]. Spastic paraplegia is a degenerative spinal cord disorder characterized by a slow, gradual, progressive weakness and spasticity of the lower limbs. Defects in HSPD1 are the cause of leukodystrophy hypomyelinating type 4 (HLD4) [MIM:612233]; also called mitochondrial HSP60 chaperonopathy or MitCHAP-60 disease. HLD4 is a severe autosomal recessive hypomyelinating leukodystrophy. Clinically characterized by infantile-onset rotary nystagmus, progressive spastic paraplegia, neurologic regression, motor impairment, profound mental retardation. Death usually occurrs within the first two decades of life.	
Sequence similarities	Belongs to the chaperonin (HSP60) family.	
Cellular localization	Mitochondrion matrix.	

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hsp60 antibody
[HSPD1/2206R] - BSA free (ab234286)

Formalin-fixed, paraffin-embedded human liver tissue stained for Hsp60 using ab234286 at 1 μ g/ml in immunohistochemical analysis.



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- · Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.com/abpromise or contact our technical team.

Terms and conditions

• Guarantee only valid for products bought direct from Abcam or one of our authorized distributors