abcam

ab303517

Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free

Recombinant | RabMAb | KO Validated

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	pH: 7.2 - 7.4 Constituents: PBS
Form	Liquid
Clonality	Monoclonal
Immunogen	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	EPR27270-11
Purification technique	Affinity purification Protein A
Concentration	1.009 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

WB

Tested

Species	Human, Mouse, Rat
Dilution info	-

Notes	-

IHC-P

Tested

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

IHC-Fr

Not recommended

Species	Mouse, Rat, Human
Dilution info	-
Notes	-

ICC/IF

Not recommended

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

Flow Cyt (Intra)

Tested

Species	Human
Dilution info	-
Notes	-

Expected

Species	Mouse, Rat
Dilution info	Use at an assay dependent concentration.

Notes	-

IP

Not recommended

Species	Mouse, Rat, Human
Dilution info	-
Notes	-

Target data

See full target information HSP90AA1

Function

Molecular chaperone that promotes the maturation, structural maintenance and proper regulation of specific target proteins involved for instance in cell cycle control and signal transduction. Undergoes a functional cycle that is linked to its ATPase activity which is essential for its chaperone activity. This cycle probably induces conformational changes in the client proteins, thereby causing their activation. Interacts dynamically with various co-chaperones that modulate its substrate recognition, ATPase cycle and chaperone function (PubMed:11274138, PubMed:12526792, PubMed:15577939, PubMed:15937123, PubMed:27353360, PubMed:29127155). Engages with a range of client protein classes via its interaction with various co-chaperone proteins or complexes, that act as adapters, simultaneously able to interact with the specific client and the central chaperone itself (PubMed:29127155). Recruitment of ATP and co-chaperone followed by client protein forms a functional chaperone. After the completion of the chaperoning process, properly folded client protein and co-chaperone leave HSP90 in an ADP-bound partially open conformation and finally, ADP is released from HSP90 which acquires an open conformation for the next cycle (PubMed:26991466, PubMed:27295069). Plays a critical role in mitochondrial import, delivers preproteins to the mitochondrial import receptor TOMM70 (PubMed:12526792). Apart from its chaperone activity, it also plays a role in the regulation of the transcription machinery. HSP90 and its co-chaperones modulate transcription at least at three different levels (PubMed:25973397). In the first place, they alter the steady-state levels of certain transcription factors in response to various physiological cues (PubMed:25973397). Second, they modulate the activity of certain epigenetic modifiers, such as histone deacetylases or DNA methyl transferases, and thereby respond to the change in the environment (PubMed:25973397). Third, they participate in the eviction of histones from the promoter region of certain genes and thereby turn on gene expression (PubMed:25973397). Binds bacterial lipopolysaccharide (LPS) and mediates LPS-induced inflammatory response, including TNF secretion by monocytes (PubMed:11276205). Antagonizes STUB1-mediated inhibition of TGF-beta signaling via inhibition of STUB1-mediated SMAD3 ubiquitination and degradation (PubMed:24613385). Mediates the association of TOMM70 with IRF3 or TBK1 in mitochondrial outer membrane which promotes host antiviral response (PubMed:20628368, PubMed:25609812). (Microbial infection) Seems to interfere

with N.meningitidis NadA-mediated invasion of human cells. Decreasing HSP90 levels increases adhesion and entry of E.coli expressing NadA into human Chang cells; increasing its levels leads to decreased adhesion and invasion.

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C
Storage information	Do Not Freeze

Notes

Patented technology
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

What are the advantages of a recombinant monoclonal antibody?
This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free batch production

For more information, read more on recombinant antibodies.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with 1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

Compatibility
This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® is a trademark of Fluidigm Canada Inc.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	Hsp90 alpha also known as Hsp90AA1 or 5G5 is a molecular chaperone with an approximate molecular weight of 90 kDa. It plays an important role in the folding stability and function of many client proteins. This protein is expressed ubiquitously in eukaryotic cells with high levels in the cytoplasm and the endoplasmic reticulum. Hsp90 alpha operates through a unique ATPase-driven chaperone cycle that modulates protein conformations helping in the assembly and disassembly of protein complexes.
Biological function summary	Hsp90 alpha assists in maintaining protein homeostasis and is part of a larger chaperome complex including co-chaperones and other chaperones such as Hsp70. It plays an important role in stress response by stabilizing proteins and preventing aggregation. Hsp90 alpha is essential for the normal function of several kinases and transcription factors which are important for cell signaling and regulation processes. It functions as a mediator for cellular response to environmental cues.
Pathways	Hsp90 alpha is an integral component in the signal transduction and cell cycle control pathways. It interacts with various signaling proteins and receptors like AKT and steroid hormone receptors to facilitate their proper folding and activation. Furthermore Hsp90 alpha influences the MAPK/ERK pathway playing a role in cell proliferation and differentiation. Its interaction with Hsp90 protein clients creates a regulatory framework across multiple pathways allowing precise control over cellular function.
Associated diseases and disorders	Hsp90 alpha is linked to cancer and neurodegenerative diseases. Its enhanced expression in tumors correlates with the stabilization of oncogenic client proteins such as HER2 contributing to cancer progression. Additionally Hsp90 alpha relates to disorders like Alzheimer's disease due to its role in handling misfolded proteins. The protein interacts with co-chaperones like H1L1 which may influence its impact on disease outcomes highlighting the therapeutic potential of targeting Hsp90 alpha in these conditions.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

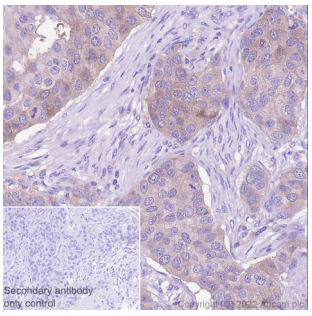
We do not recommend this combination. It is not covered by our product promise.

We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

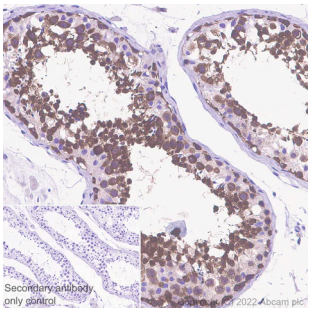
Full details and terms and conditions can be found here:
Terms & Conditions.

12 product images



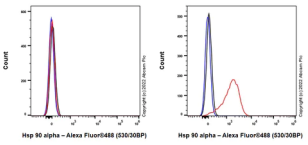
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue labeling Hsp90 alpha with ab303516 at 1/1000 dilution (0.462 µg/mL), followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Positive staining was observed on human breast carcinoma. The section was incubated with ab303516 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) at Ready to use dilution. Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



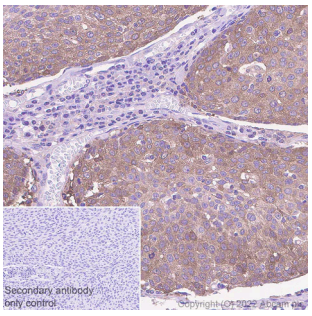
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin-embedded Human testis tissue labeling Hsp90 alpha with ab303516 at 1/1000 (0.462 ug/ml) followed by a ready to use Rabbit specific IHC polymer detection kit HRP/Dab (ab209101). Positive staining on human testis. The section was incubated with ab303516 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use Rabbit specific IHC polymer detection kit HRP/Dab (ab209101).Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



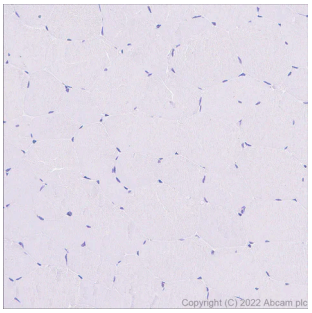
Flow Cytometry (Intracellular) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Wild-type Wild-type 293T (human embryonic kidney epithelial cell, Right) / Hsp90 alpha knockout 293T(Left) cells labelling Hsp90 alpha with ab303516 at 1/50 dilution (1ug) (Red) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Positive staining on 293T cells (ab255449), while no staining on Hsp90 alpha knockout 293T cells (ab266591).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

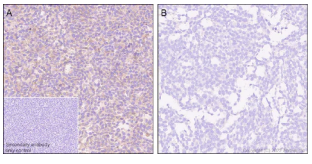
This data was developed using ab303516, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin-embedded Human lung carcinoma tissue labeling Hsp90 alpha with ab303516 at 1/1000 (0.462 ug/ml) followed by a ready to use Rabbit specific IHC polymer detection kit HRP/Dab (ab209101). Positive staining on human lung carcinoma. The section was incubated with ab303516 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use Rabbit specific IHC polymer detection kit HRP/Dab (ab209101).Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

Hsp 90 alpha Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-Hsp 90 alpha antibody

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded human skeletal muscle tissue labeling Hsp90 alpha with ab303516 at 1/1000 dilution (0.462 µg/mL), followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Negative control: no staining is observed on human skeletal muscle. The section was incubated with ab303516 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



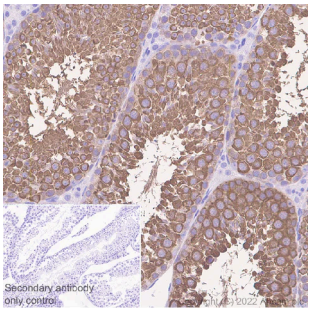
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded (A) parental HEK293T tissue labeling Hsp90 alpha with ab303516 at 1/2000 dilution (0.231 µg/mL), followed by a ready to use Rabbit specific

IHC polymer detection kit HRP/DAB (ab209101). Positive staining was observed on (A) parental HEK293T cell pellet, no staining is observed on (B) Hsp90 alpha KO HEK293T cell pellet. The section was incubated with ab303516 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) added without primary.

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

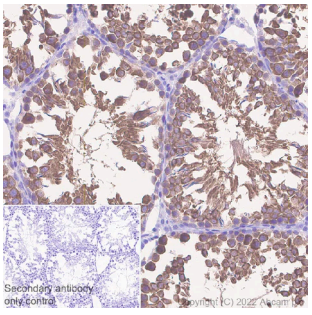


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded mouse testis tissue labeling Hsp90 alpha with ab303516 at 1/1000 dilution (0.462 µg/mL), followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Positive staining was observed on mouse testis. The section was incubated with ab303516 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) at Ready to use dilution.

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins

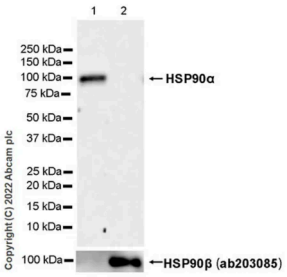


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded rat testis tissue labeling Hsp90 alpha with ab303516 at 1/1000 dilution (0.462 µg/mL), followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Positive staining was observed on rat testis. The section was incubated with ab303516 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) at Ready to use dilution.

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins



Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

Hsp 90 alpha Western blot staining using rabbit Anti-Hsp 90 alpha antibody

This data was developed using ab303516, the same antibody clone in a different buffer formulation.
Blocking / Diluting buffer and concentration: 5% NFDM/TBST.

Negative control: skeletal muscle (PMID:21209834).

This antibody does not cross-react with human Hsp90-beta.

All lanes:
Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] (ab303516) at 1/1000 dilution

Lane 1:
Human Hsp90-alpha active full length protein at 500 ng

Lane 2:
Human Hsp90-beta active full length protein at 500 ng

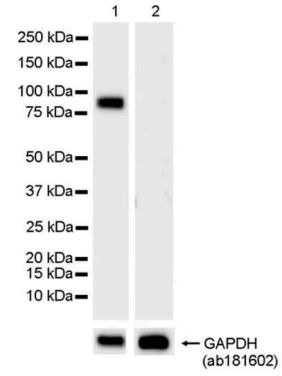
Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Predicted band size: 85 kDa

Observed band size: 90 kDa

Exposure time: 114s



Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Blocking / Diluting buffer and concentration: 5% NFDM/TBST Negative control: skeletal muscle (PMID:21209834)
All lanes:
Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] (ab303516) at 1/1000 dilution

Lane 1:
Human testis tissue lysate at 20 µg

Lane 2:
Human skeletal muscle tissue lysate at 20 µg

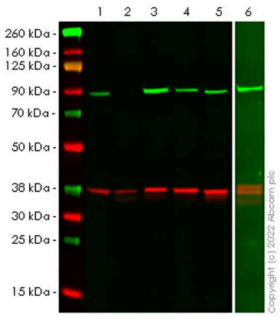
Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Predicted band size: 85 kDa

Observed band size: 90 kDa

Exposure time: 48s



Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Blocking / Diluting buffer and concentration: 5% NFDM/TBST The samples were run on a Bis-Tris gel. Performed under reducing conditions. False colour image of Western blot: Anti-Hsp90 alpha antibody (ab303516) staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab303516 was shown to bind specifically to Hsp90 alpha. A band was observed at 90 kDa in wild-type 293T cell lysates with no signal observed at this size in Hsp90 alpha knockout cell line. To generate this image, wild-type and Hsp90 alpha knockout 293T cell lysates were analyzed. First, samples were run on an SDS-PAGE gel then transferred onto an immobilon-FL PVDF membrane. Membranes were blocked in in Intercept® (TBS) Blocking Buffer diluted with an equal volume of 0.1% TBS before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged.
All lanes:
Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] (ab303516) at 1/1000 dilution

Lane 1:
Wild-type 293T (human embryonic kidney epithelial cell) whole cell lysate at 20 µg

Lane 2:
Hsp90 alpha knockout 293T whole cell lysate at 20 µg

Lane 3:
Human testis tissue lysate, at 20 µg

Lane 4:
Human kidney tissue lysate at 20 µg

Lane 5:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate at 20 µg

Lane 6:
C6 (rat glial tumor glial cell), whole cell lysate at 20 µg

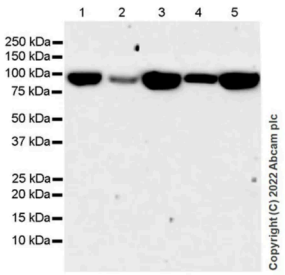
Secondary

Lanes 1 - 6:
Western blot - Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/20000 dilution

Lanes 1 - 6:
Western blot - Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) at 1/20000 dilution

Predicted band size: 85 kDa

Observed band size: 90 kDa



Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] - BSA and Azide free (ab303517)

This data was developed using ab303516, the same antibody clone in a different buffer formulation. Blocking / Diluting buffer and concentration: 5% NFDM/TBST
All lanes:
Western blot - Anti-Hsp 90 alpha antibody [EPR27270-11] (ab303516) at 1/1000 dilution

Lane 1:
Mouse brain tissue lysate at 20 µg

Lane 2:
Mouse kidney tissue lysate at 20 µg

Lane 3:
Mouse testis tissue lysate at 20 µg

Lane 4:
Rat brain tissue lysate at 20 µg

Lane 5:
Rat testis tissue lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Predicted band size: 85 kDa

Observed band size: 90 kDa

Exposure time: 114s

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.