abcam

Product datasheet

Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free ab247912

יובעבלאר RabMAb

画像数9

製品の概要

特記事項

製品名 Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR4576] to Hsp105/HSP110 - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: IHC-P, IP, WB, ICC/IF

種交差性 交差種: Mouse, Rat, Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ab247912 is the carrier-free version of ab109624.

Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased 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.

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.

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 production

For more information see here.

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

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

バッファー pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

ポリ/モノ モノクローナル **クローン名** EPR4576

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab247912の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols.
IP		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 97 kDa.
ICC/IF		Use at an assay dependent concentration.

ターゲット情報

機能 Prevents the aggregation of denatured proteins in cells under severe stress, on which the ATP levels decrease markedly. Inhibits HSPA8/HSC70 ATPase and chaperone activities.

組織特異性 Highly expressed in testis. Present at lower levels in most brain regions, except cerebellum. Overexpressed in cancer cells.

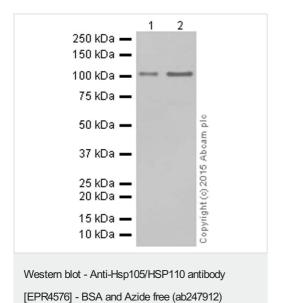
配列類似性 Belongs to the heat shock protein 70 family.

翻訳後修飾 Phosphorylation on Ser-509 may be important for regulation of the HSPA8/HSC70 chaperone

activity.

細胞内局在 Cytoplasm.

画像



All lanes: Anti-Hsp105/HSP110 antibody [EPR4576] (ab109624) at 1/5000 dilution (purified)

Lane 1: HeLa whole cell lysate Lane 2: HepG2 whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

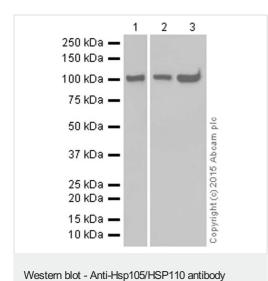
All lanes: HRP-conjugated anti-rabbit lgG, specific to the nonreduced form of IgG at 1/1000 dilution

Predicted band size: 97 kDa

Observed band size: 105 kDa

This data was developed using ab109624, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.



All lanes: Anti-Hsp105/HSP110 antibody [EPR4576] (ab109624) at 1/5000 dilution (purified)

Lane 1: C6 whole cell lysate Lane 2: PC-12 whole cell lysate

Lane 3: NIH/3T3 whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: HRP-conjugated anti-rabbit lgG, specific to the nonreduced form of IgG at 1/1000 dilution

Predicted band size: 97 kDa Observed band size: 105 kDa

[EPR4576] - BSA and Azide free (ab247912)

This data was developed using ab109624, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.

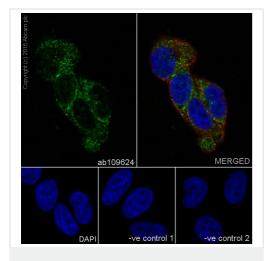
Secondary antibody only control.

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Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912)

This data was developed using <u>ab109624</u>, the same antibody clone in a different buffer formulation.

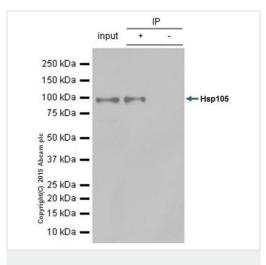
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human liver carcinoma tissue labelling Hsp105/HSP110 with purified ab109624 at a dilution of 1/500. Heat mediated antigen retrieval was performed using EDTA buffer pH 9. ab97051, a HRP-conjugated goat anti-rabbit lgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.



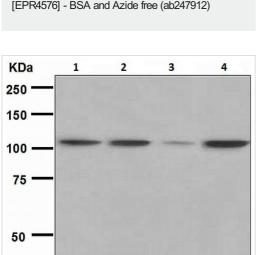
Immunocytochemistry/ Immunofluorescence - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912)

This data was developed using <u>ab109624</u>, the same antibody clone in a different buffer

formulation.Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling Hsp105/HSP110 with purified <u>ab109624</u> at a dilution of 1/250. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. <u>ab150077</u>, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/1000) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain. <u>ab7291</u>, a mouse anti-tubulin (1/1000) and <u>ab150120</u>, an Alexa Fluor[®] 594-conjugated goat anti-mouse lgG (1/1000) were also used. Control 1: primary antibody (1/250) and secondary antibody, <u>ab150120</u>, an Alexa Fluor[®] 594-conjugated goat anti-mouse lgG (1/1000). Control 2: <u>ab7291</u> (1/1000) and secondary antibody, <u>ab150077</u>, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/1000).



Immunoprecipitation - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912)



Western blot - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912)

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This data was developed using ab109624&

44; the same antibody clone in a different buffer formulation.ab109624 (purified) at 1/40 immunoprecipitating Hsp105/HSP110 in NIH/3T3 whole cell lysate.

Lane 1 (input): NIH/3T3 whole cell lysate (10µg)

Lane 2 (+): <u>ab109624</u> + NIH/3T3 whole cell lysate.

Lane 3 (-): Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab109624</u> in NIH/3T3 whole cell lysate.

For western blotting, VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/10,000 dilution.

Blocking and dilution buffer: 5% NFDM/TBST.

All lanes : Anti-Hsp105/HSP110 antibody [EPR4576] (**ab109624**) at 1/1000 dilution (unpurified)

Lane 1: HeLa cell lysate

Lane 2: HepG2 cell lysate

Lane 3: Human cerebellum lysate

Lane 4: MCF-7 cell lysate

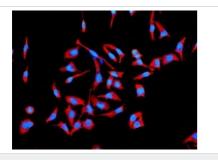
Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP-conjugated goat anti-rabbit lgG at 1/2000 dilution

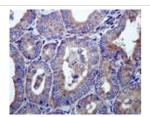
Predicted band size: 97 kDa

This data was developed using <u>ab109624</u>, the same antibody clone in a different buffer formulation.



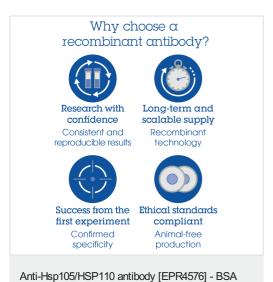
Immunocytochemistry/ Immunofluorescence - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912) This data was developed using <u>ab109624</u>, the same antibody clone in a different buffer

formulation.lmmunocytochemistry/lmmunofluorescence analysis of HeLa cells labelling Hsp105/HSP110 with unpurified <u>ab109624</u> at a dilution of 1/100.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Hsp105/HSP110 antibody [EPR4576] - BSA and Azide free (ab247912)

This data was developed using <u>ab109624</u>, the same antibody clone in a different buffer formulation.lmmunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human thyroid gland carcinoma tissue labelling Hsp105/HSP110 with unpurified <u>ab109624</u> at a dilution of 1/50. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



and Azide free (ab247912)

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