abcam

Product datasheet

Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free ab233702

יעלאעבע RabMAb

画像数 14

製品の概要

製品名 Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR20842] to ABAT/GABA-T - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), WB, IHC-P, IHC-Fr, IP, 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.

ポジティブ・コントロール IHC-P: Human kidney tissue.

特記事項 ab233702 is the carrier-free version of ab216465.

> 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.

製品の特性

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 EPR20842

アイソタイプ IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab233702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
Flow Cyt (Intra)		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 56 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
IHC-Fr		Use at an assay dependent concentration.
IP		Use at an assay dependent concentration.
ICC/IF		Use at an assay dependent concentration.

ターゲット情報

機能 Catalyzes the conversion of gamma-aminobutyrate and L-beta-aminoisobutyrate to succinate

semialdehyde and methylmalonate semialdehyde, respectively. Can also convert delta-

aminovalerate and beta-alanine.

組織特異性 Liver > pancreas > brain > kidney > heart > placenta.

関連疾患 Defects in ABAT are a cause of gamma-aminobutyrate transaminase deficiency (GABA-AT

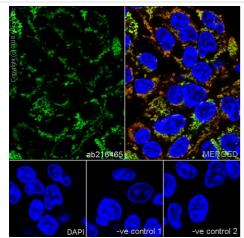
deficiency) [MIM:613163]. The phenotype of this deficiency includes psychomotor retardation,

hypotonia, hyperreflexia, lethargy, refractory seizures, and EEG abnormalities.

配列類似性 Belongs to the class-Ill pyridoxal-phosphate-dependent aminotransferase family.

細胞内局在 Mitochondrion matrix.

画像



Immunocytochemistry/ Immunofluorescence - Anti-

Immunofluorescent analysis of 100% methanol-fixed MCF7 (human breast adenocarcinoma epithelial cell) cells labeling ABAT/GABA-T with ab216465 at 1/100 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 (green). Confocal image showing mitochondrial staining in MCF7 cell line. The nuclear counter stain is DAPI (blue).

Immunofluorescent analysis of 100% methanol-fixed HepG2 (human hepatocellular carcinoma epithelial cell) cells labeling ABAT/GABA-T with ab216465 at 1/100 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor[®] 488) (ab150077) secondary antibody at 1/1000 (green). Confocal image showing mitochondrial staining in HepG2 cell line is observed. The nuclear counter stain is DAPI (blue).

Mitochondria are stained with <u>ab33985</u> Anti-COX IV (mouse mAb) - Mitochondrial Marker followed by <u>ab150120</u> AlexaFluor[®]594 Goat

-ve control 1: ab216465 at 1/100 dilution followed by ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution.

-ve control 2: ab33985 Anti-COX IV (mouse mAb) - Mitochondrial

Marker at 1/1000 dilution followed by ab150077 (AlexaFluor®488

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and

anti-Mouse secondary both at 1/1000 dilution (red).

Goat anti-Rabbit secondary) at 1/1000 dilution.

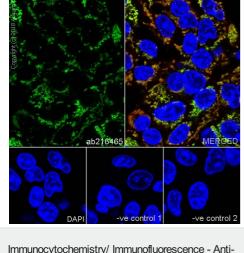
sodium azide (ab216465).

Mitochondria are stained with **ab33985** Anti-COX IV (mouse mAb) - Mitochondrial Marker followed by <u>ab150120</u> AlexaFluor[®]594 Goat anti-Mouse secondary both at 1/1000 dilution (red).

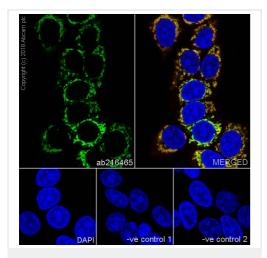
-ve control 1: **ab216465** at 1/100 dilution followed by **ab150120** (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution.

-ve control 2: ab33985 Anti-COX IV (mouse mAb) - Mitochondrial Marker at 1/1000 dilution followed by ab150077 (AlexaFluor®488 Goat anti-Rabbit secondary) at 1/1000 dilution.

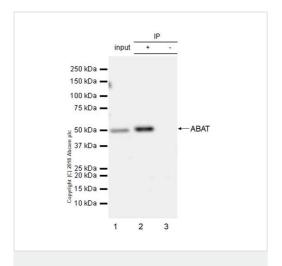
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



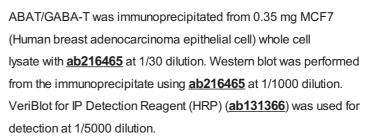
ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)



Immunocytochemistry/ Immunofluorescence - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)



Immunoprecipitation - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)



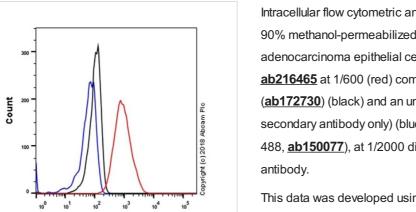
Lane 1: MCF7 (Human breast adenocarcinoma epithelial cell) whole cell lysate 10 µg (Input).

Lane 2: ab216465 IP in MCF7 whole cell lysate (+).

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab216465</u> in MCF7 whole cell lysate (-).

Blocking/Dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 3 minutes.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).

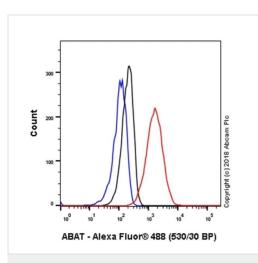


Flow Cytometry (Intracellular) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

ABAT - Alexa Fluor® 488 (530/30 BP)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol-permeabilized MCF7 (Human breast adenocarcinoma epithelial cell) cells labeling ABAT/GABA-T with **ab216465** at 1/600 (red) compared with a Rabbit monoclonal IgG (**ab172730**) (black) and an unlabelled control (cellsincubated with secondary antibody only) (blue). Goat anti rabbit IgG (Alexa Fluor[®] 488, **ab150077**), at 1/2000 dilution was used as the secondary antibody.

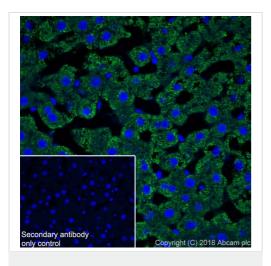
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



Flow Cytometry (Intracellular) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol-permeabilized HepG2 (Human hepatocellular carcinoma epithelial cell) cell line labeling ABAT/GABA-T with ab216465 at 1/60 (red) compared with a Rabbit monoclonal IgG (ab172730) (black) and an unlabelled control (cellsincubated with secondary antibody only) (blue). Goat anti rabbit IgG (Alexa Fluor® 488, ab150077), at 1/2000 dilution was used as the secondary antibody.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



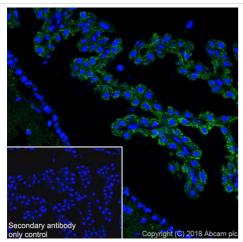
Immunohistochemistry (Frozen sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse liver tissue labeling ABAT/GABA-T with <u>ab216465</u> at 1/500 dilution (green), followed by <u>ab150077</u> AlexaFluor[®]488 Goat anti-Rabbit secondary at a 1/1000 dilution. Cytoplasmic staining in rat liver is observed. Counter stained with DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab150077</u> AlexaFluor[®]488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat-mediated antigen retrieval by using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



Immunohistochemistry (Frozen sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse choroid plexus tissue labeling ABAT/GABA-T with ab216465 at 1/500 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000 dilution. Cytoplasmic staining in mouse choroid plexus (PMID:25239459, PMID: 11459221) is observed. Counter stained with DAPI (blue).

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100

ABAT/GABA-T with ab216465 at 1/500 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000

(PMID:25239459, PMID: 11459221) is observed. Counter stained

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 AlexaFluor®488 Goat

Perform heat-mediated antigen retrieval by using sodium citrate

This data was developed using the same antibody clone in a

different buffer formulation containing PBS, BSA, glycerol, and

permeabilized frozen rat choroid plexus tissue labeling

dilution. Cytoplasmic staining in rat choroid plexus

buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

anti-Rabbit used at a 1/1000 dilution.

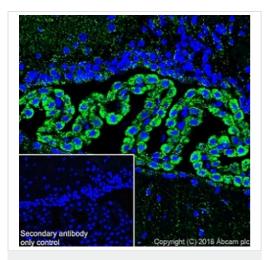
sodium azide (ab216465).

with DAPI (blue).

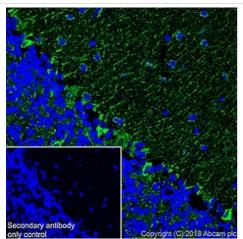
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 AlexaFluor®488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat-mediated antigen retrieval by using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



Immunohistochemistry (Frozen sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)



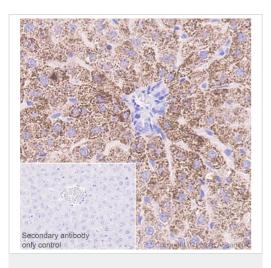
Immunohistochemistry (Frozen sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of paraffin-embedded rat liver tissue labeling ABAT/GABA-T with ab216465 at 1/500 dilution, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Granular cytoplasmic staining in rat liver (PMID: 25771305; PMID: 25738457) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit IgG H&L (HRP).

Perform heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab216465</u>).



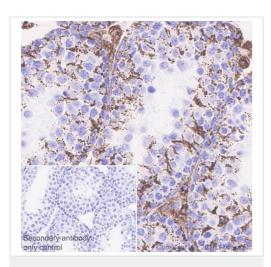
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100

antibody, secondary antibody is ab150077 AlexaFluor®488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat-mediated antigen retrieval by using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



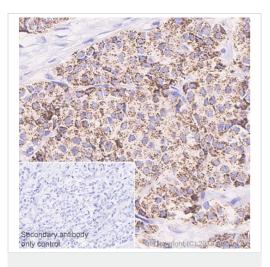
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of paraffin-embedded mouse testis tissue labeling ABAT/GABA-T with <u>ab216465</u> at 1/500 dilution, followed by a ready to use Goat Anti-Rabbit lgG H&L (HRP). Granular cytoplasmic staining in mouse testis (PMID: 25771305; PMID: 25738457) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



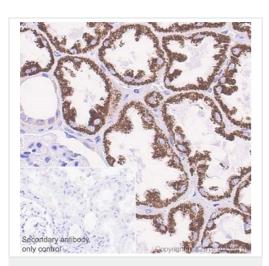
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ABAT/GABA-T antibody
[EPR20842] - BSA and Azide free (ab233702)

Immunohistochemical analysis of paraffin-embedded human breast cancer tissue labeling ABAT/GABA-T with <u>ab216465</u> at 1/500 dilution, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Granular cytoplasmic staining in human breast cancer (PMID: 25771305; PMID: 25738457) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ABAT/GABA-T antibody
[EPR20842] - BSA and Azide free (ab233702)

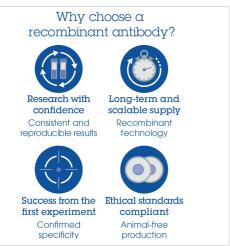
Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling ABAT/GABA-T with <u>ab216465</u> at 1/500 dilution, followed by a ready to use Goat Anti-Rabbit lgG H&L (HRP).

Granular cytoplasmic staining in human kidney (PMID: 25771305; PMID: 25738457) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit IgG H&L (HRP).

Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab216465).



Anti-ABAT/GABA-T antibody [EPR20842] - BSA and Azide free (ab233702)

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