abcam

ab316106

Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free

Recombinant | RabMAb | Advanced Validation

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	EPR26423-34
Purification technique	Affinity purification Protein A
Specificity	Unsuitable for Human WB.
Concentration	1.019 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

mIHC

Tested

Species	Human, Rat, Mouse

Dilution info	-
Notes	-

Not recommended

Species	Transfected cell line
Dilution info	-
Notes	-

IHC-P

Tested

Species	Transfected cell line, Human, Rat
Dilution info	-
Notes	-

Expected

Species	Mouse
Dilution info	Use at an assay dependent concentration.
Notes	-

Flow Cyt (Intra)

Tested

Species	Transfected cell line, Mouse
Dilution info	-
Notes	-

Expected

Species	Human, Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

WB

Tested

Species	Mouse, Rat
Dilution info	-
Notes	-

Not recommended

Species	Human, Transfected cell line
Dilution info	-
Notes	-

IP

Tested

Species	Mouse
Dilution info	-
Notes	-

Expected

Species	Human, Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Transfected cell line
Dilution info	-
Notes	-

ICC/IF

Tested

Species	Mouse, Rat
Dilution info	-
Notes	-

Expected

Species	Human
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Transfected cell line
Dilution info	-
Notes	-

IHC-Fr

Tested

Species	Mouse, Rat
Dilution info	-
Notes	-

Expected

Species	Human
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Transfected cell line
Dilution info	-
Notes	-

Target data

See full target information Gpr17

Function	Dual specificity receptor for uracil nucleotides and cysteinyl leukotrienes (CysLTs). Signals through G(i) and inhibition of adenylyl cyclase. May mediate brain damage by nucleotides and CysLTs following ischemia (By similarity).

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C

Notes

ab316106 is the carrirer-free version of ab316105.

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	GPR17 also known as G protein-coupled receptor 17 is a member of the G protein-coupled receptor (GPCR) family. This receptor has a molecular weight of approximately 40 kDa. It is expressed in various tissues with significant levels found in the central nervous system including the brain and spinal cord. GPR17 also shows expression in kidneys liver and heart. It is classified under the class A Rhodopsin-like family and is an orphan receptor meaning its natural ligand is not completely established but is thought to be associated with uridine nucleotides and cysteinyl leukotrienes.
Biological function summary	GPR17 is involved in several physiological processes particularly in the regulation of inflammation and myelination within the central nervous system. It plays a critical role in the differentiation of oligodendrocyte precursor cells into mature oligodendrocytes which are essential for myelin sheath formation and repair of neural damage. GPR17 is not known to function as part of a larger receptor complex but it interacts with intracellular signaling pathways to modulate these cellular processes.
Pathways	GPR17 integrates into inflammatory response pathways and neural cell development pathways. It is actively involved in the nucleotide signaling pathway which is important for responses to tissue damage and initiating repair processes. GPR17 shares functional connections with proteins such as P2Y receptors which also respond to nucleotides and are involved in diverse cellular responses including muscle contraction and platelet aggregation.
Associated diseases and disorders	GPR17's expression and role connect it to multiple sclerosis and brain ischemia. In multiple sclerosis dysregulation of GPR17 can affect the formation and maintenance of the myelin sheath contributing to neural degeneration. During brain ischemia GPR17 activation influences the extent of injury and subsequent recovery through its effects on inflammation and tissue repair. It is linked to proteins like CXCR7 which similarly influence neural damage and repair processes in ischemic 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

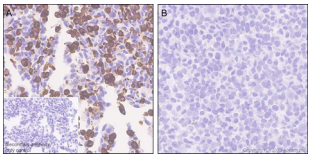
We do not recommend this combination. It is not covered by our product promise.

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.

35 product images



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

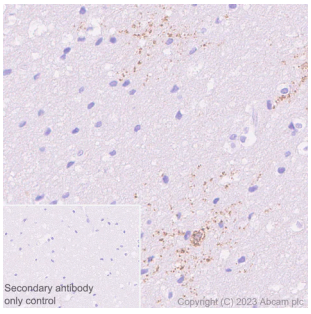
Immunohistochemical analysis of paraffin-embedded (A) HEK-293T (human epithelial cell line from embryonic kidney) transfected with a GPR17 expression vector containing a Myc-His tag. (B) HEK-293T transfected with empty vector containing a Myc-His tag. tissue labeling GPCR GPR17 with ab316105 at 1/5000 (0.102 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on (A) HEK-293T (human epithelial cell line from embryonic kidney) transfected with a GPR17 expression vector containing a Myc-His tag, negative staining on (B) HEK-293T transfected with empty vector containing a Myc-His tag. The section was incubated with ab316105 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 LeicaDS9800 (Bond™ Polymer Refine Detection).

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-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

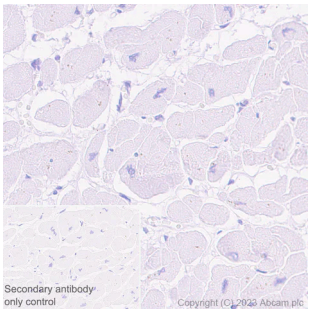
Immunohistochemical analysis of paraffin-embedded Human cerebrum tissue labeling GPCR GPR17 with ab316105 at 1/500 (1.018 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Positive staining on oligodendrocytes of human cerebrum (PMID: 28357182; PMID: 23801362). The section was incubated with ab316105 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-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Human cardiac muscle using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

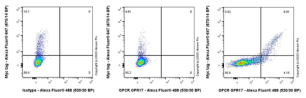
Immunohistochemical analysis of paraffin-embedded Human cardiac muscle tissue labeling GPCR GPR17 with ab316105 at 1/500 (1.018 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Negative control: no staining on human cardiac muscle. The section was incubated with ab316105 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-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

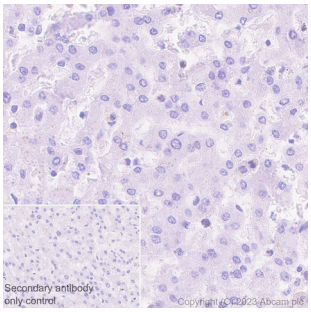
GPCR GPR17 Flow Cytometry (Intracellular) staining using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Isotype (Left) / 293T (human embryonic kidney epithelial cell) transfected with an empty expression vector containing a myc tag (Middle) / 293T transfected with a GPCR expression vector containing a myc tag (Right) cells labelling GPCR GPR17 with ab316105 at 1/500 dilution (0.1 ug)/Middle and Right compared with a Rabbit monoclonal IgG (ab172730) / Left isotype control.

Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.

Cells are co-stained with Myc tag conjugated to Alexa Fluor®647.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Human liver using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of paraffin-embedded Human liver tissue labeling GPCR GPR17 with ab316105 at 1/500 (1.018 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

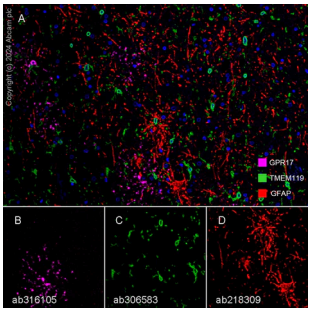
Negative control: no staining on human liver (PMID: 16990797). The section was incubated with ab316105 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

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

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Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Human cerebrum tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded human cerebrum tissue sections.

Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-TMEM119 (green; Opal™520) and anti-GFAP (red; Opal™570) on human cerebrum.

Panel B: anti-GPR17 staining oligodendrocytes in human cerebrum.

Panel C: anti-TMEM119 staining microglia in human cerebrum.

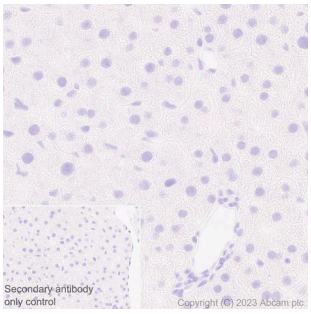
Panel D: anti-GFAP staining astrocytes in human cerebrum.

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab306583 at a 1/2000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Rat liver using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of paraffin-embedded Rat liver tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

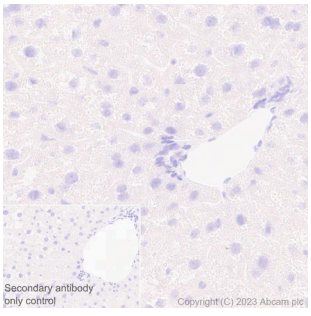
Negative control: no staining on rat liver (PMID: 16990797). The section was incubated with ab316105 for 30 mins at room temperature.
The immunostaining was performed on a Leica Biosystems BOND® RX instrument

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

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Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Mouse liver using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

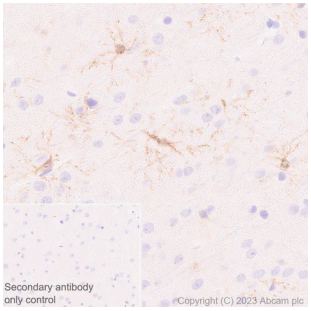
Immunohistochemical analysis of paraffin-embedded Mouse liver tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Negative control: no staining on mouse liver. The section was incubated with ab316105 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-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

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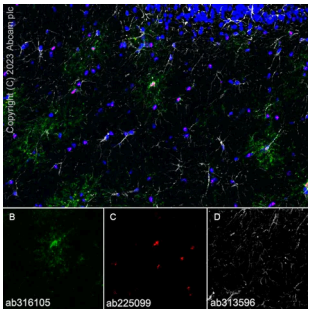
Immunohistochemical analysis of paraffin-embedded Rat cerebrum tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on oligodendrocytes of rat cerebrum (PMID: 21209081). The section was incubated with ab316105 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 LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Frozen sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

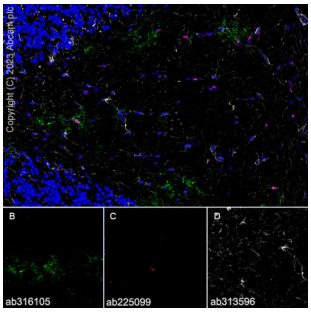
This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Rat cerebellum (fresh frozen) tissue labeling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution followed by ab150088 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) preadsorbed (ab150088) at 1/1000 (2 ug/mL) dilution (Green).

Panel A: merged staining of anti-GPCR GPR17 (ab316105, green), anti-Olig2 (ab225099, red) and anti-GFAP (ab313596, grey) on rat cerebellum.
Panel B: anti-GPCR GPR17 stained on rat cerebellum.
Panel C: anti-Olig2 stained in oligodendrocyte of rat cerebellum.

Panel D: anti-GFAP stained in astrocytes of rat cerebellum.
The nuclear counterstain was DAPI (Blue). The section was incubated with ab316105 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150088 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) preadsorbed (ab150088)at 1/1000 (2 ug/mL) dilution.



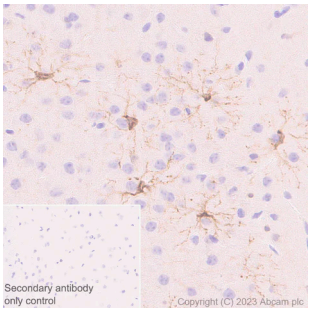
Immunohistochemistry (Frozen sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

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Secondary antibody control: Secondary antibody is ab150088 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) preadsorbed (ab150088)at 1/1000 (2 ug/mL) dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

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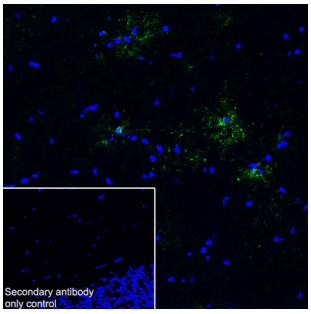
Immunohistochemical analysis of paraffin-embedded Mouse cerebrum tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Positive staining on oligodendrocytes of mouse cerebrum (PMID: 26453964). The section was incubated with ab316105 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 LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



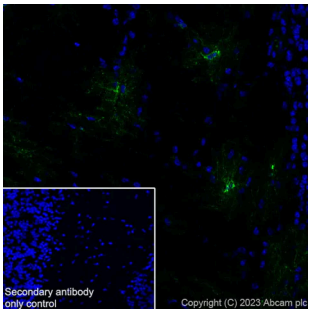
Immunohistochemistry (Frozen sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

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Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Rat cerebellum (fresh frozen) tissue labeling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution (Green).

Confocal image showing positive staining on rat cerebellum. The nuclear counterstain was DAPI (Blue). The section was incubated with ab316105 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbedat 1/1000 (2 ug/mL) dilution.



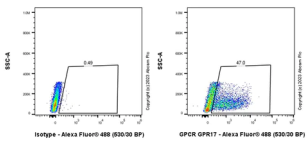
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Confocal image showing positive staining on mouse cerebellum. The nuclear counterstain was DAPI (Blue). The section was incubated with ab316105 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

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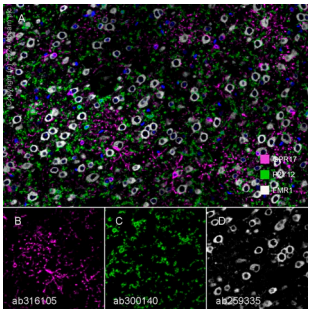


Flow Cytometry (Intracellular) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized Mouse primary neuron cells labelling GPCR GPR17 with ab316105 at 1/500 dilution (0.1 ug)/Right compared with a Rabbit monoclonal IgG (ab172730) / Left isotype control.

Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse cerebrum tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-FMR1 (gray; Opal™570) on mouse cerebrum.

Panel B: anti-GPR17 staining oligodendrocytes in mouse cerebrum.

Panel C: anti-P2RY12 staining microglia in mouse cerebrum.

Panel D: anti-FMR1 staining neurons in mouse cerebrum.

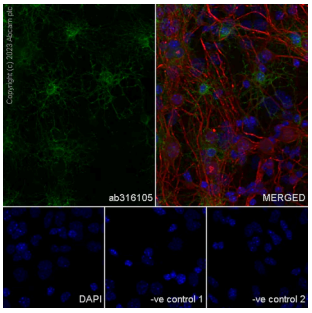
Nuclear DNA was labeled with DAPI (shown in blue).

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Immunocytochemistry/ Immunofluorescence - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunocytochemistry/ Immunofluorescence staining of mouse primary neural/glia cell using rabbit Anti-GPCR GPR17 antibody

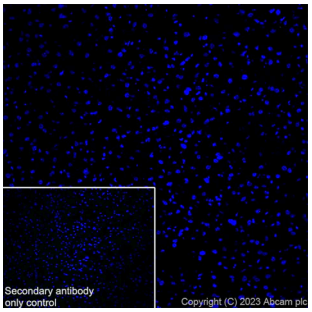
This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized mouse primary neural/glia cell cells labelling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 (2 ug/mL) dilution (Green).

Confocal image showing cytoplasmic staining in mouse oligodendrocytes.
Confocal scanning Z step was set as 0.3 µm followed by image processing with maximum Z projection.
Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).

ab11267 Anti-MAP2 mouse monoclonal antibody was used to counterstain at 1/500 (4 ug/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution.



Immunohistochemistry (Frozen sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

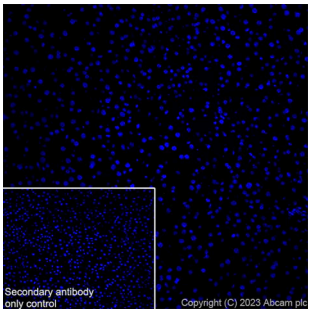
GPCR GPR17 Immunohistochemistry (Frozen sections) staining of Mouse liver (fresh frozen) using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Mouse liver (fresh frozen) tissue labeling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution (Green).

Negative control: confocal image showing no staining on mouse liver. The nuclear counterstain was DAPI (Blue). The section was incubated with ab316105 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbedat 1/1000 (2 ug/mL) dilution.



Immunohistochemistry (Frozen sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

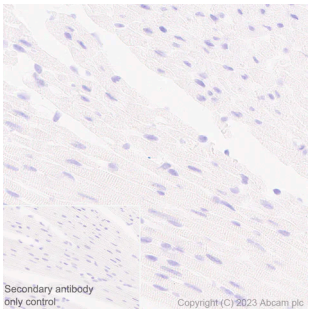
GPCR GPR17 Immunohistochemistry (Frozen sections) staining of Rat liver (fresh frozen) using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen Rat liver (fresh frozen) tissue labeling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution (Green).

Negative control: confocal image showing no staining on rat liver (PMID: 16990797). The nuclear counterstain was DAPI (Blue). The section was incubated with ab316105 for 60 mins at room temperature. The section was then mounted using Fluoromount®.The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Secondary antibody control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbedat 1/1000 (2 ug/mL) dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Mouse cardiac muscle using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

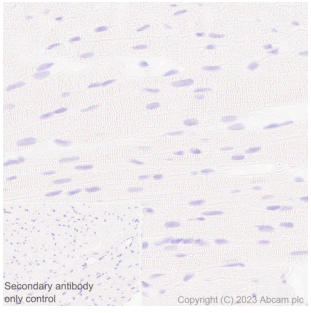
Immunohistochemical analysis of paraffin-embedded Mouse cardiac muscle tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Negative control: no staining on mouse cardiac muscle. The section was incubated with ab316105 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-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining of Rat cardiac muscle using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

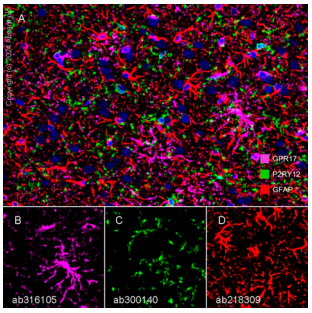
Immunohistochemical analysis of paraffin-embedded Rat cardiac muscle tissue labeling GPCR GPR17 with ab316105 at 1/2000 (0.255 ug/ml) dilution, followed by a ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Negative control: no staining on rat cardiac muscle. The section was incubated with ab316105 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



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Rat cerebrum tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded rat cerebrum tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on rat cerebrum.

Panel B: anti-GPR17 staining oligodendrocytes in rat cerebrum.

Panel C: anti-P2RY12 staining microglia in rat cerebrum.

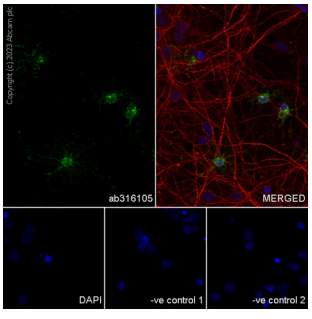
Panel D: anti-GFAP staining astrocytes in rat cerebrum.

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/140000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Immunocytochemistry/ Immunofluorescence - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Immunocytochemistry/ Immunofluorescence staining of rat primary neural/glia cell using rabbit Anti-GPCR GPR17 antibody

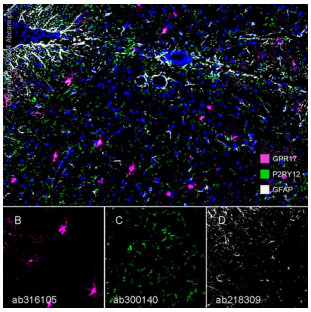
This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized rat primary neural/glia cell cells labelling GPCR GPR17 with ab316105 at 1/50 (10.18 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 (2 ug/mL) dilution (Green).

Confocal image showing cytoplasmic staining in rat oligodendrocytes.
Confocal scanning Z step was set as 0.3 µm followed by image processing with maximum Z projection.
Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).

ab11267 Anti-MAP2 mouse monoclonal antibody was used to counterstain at 1/500 (4 ug/ml) dilution (Red). The Nuclear counterstain was DAPI (Blue).

Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 (2 ug/mL) dilution.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Mouse spinal cord tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse spinal cord tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on mouse spinal cord.

Panel B: anti-GPR17 staining oligodendrocytes in mouse spinal cord.

Panel C: anti-P2RY12 staining microglia in mouse spinal cord.

Panel D: anti-GFAP staining astrocytes in mouse spinal cord.

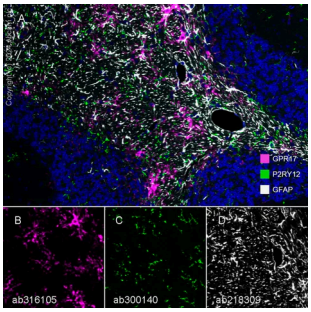
The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Mouse cerebellum tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse cerebellum tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on mouse cerebellum.

Panel B: anti-GPR17 staining oligodendrocytes in mouse cerebellum.

Panel C: anti-P2RY12 staining microglia in mouse cerebellum.

Panel D: anti-GFAP staining astrocytes in mouse cerebellum.

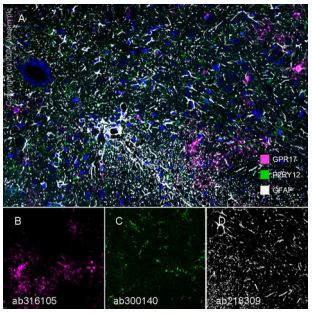
The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Rat spinal cord tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded rat spinal cord tissue sections.

Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on rat spinal cord.

Panel B: anti-GPR17 staining oligodendrocytes in rat spinal cord.

Panel C: anti-P2RY12 staining microglia in rat spinal cord.

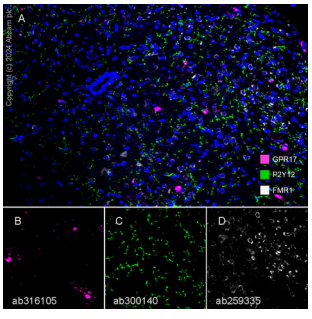
Panel D: anti-GFAP staining astrocytes in rat spinal cord.

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse spinal cord tissue.

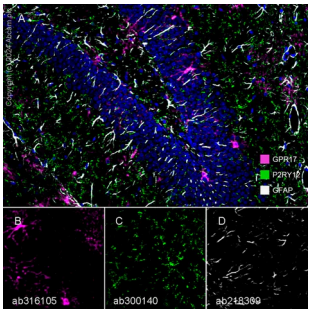
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and

anti-FMR1 (gray; Opal™570) on mouse spinal cord.
Panel B: anti-GPR17 staining oligodendrocytes in mouse spinal cord.
Panel C: anti-P2RY12 staining microglia in mouse spinal cord.
Panel D: anti-FMR1 staining neurons in mouse spinal cord.
Nuclear DNA was labeled with DAPI (shown in blue).
The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Mouse hippocampus tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse hippocampus tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on mouse hippocampus.

Panel B: anti-GPR17 staining oligodendrocytes in mouse hippocampus.

Panel C: anti-P2RY12 staining microglia in mouse hippocampus.

Panel D: anti-GFAP staining astrocytes in mouse hippocampus.

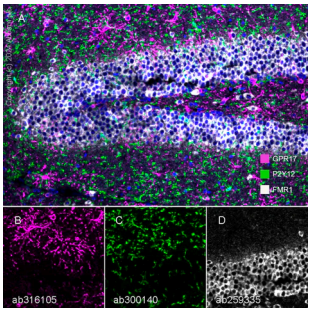
The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse hippocampus tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-FMR1 (gray; Opal™570) on mouse hippocampus.

Panel B: anti-GPR17 staining oligodendrocytes in mouse hippocampus.

Panel C: anti-P2RY12 staining microglia in mouse hippocampus.

Panel D: anti-FMR1 staining neurons in mouse hippocampus.

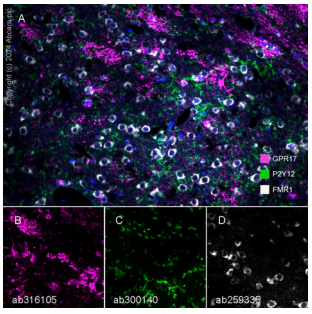
Nuclear DNA was labeled with DAPI (shown in blue).

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins. Nuclear counterstaining with DAPI.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded rat cerebrum tissue sections.

Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-FMR1 (gray; Opal™570) on rat cerebrum.

Panel B: anti-GPR17 staining oligodendrocytes in rat cerebrum.

Panel C: anti-P2RY12 staining microglia in rat cerebrum.

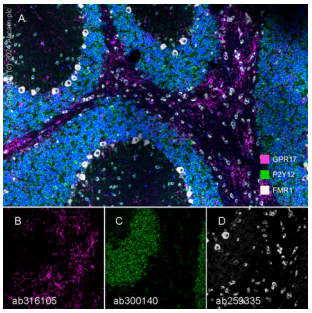
Panel D: anti-FMR1 staining neurons in rat cerebrum.

Nuclear DNA was labeled with DAPI (shown in blue).

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins. Nuclear counterstaining with DAPI.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse cerebellum tissue sections.

Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-FMR1 (gray; Opal™570) on mouse cerebellum.

Panel B: anti-GPR17 staining oligodendrocytes in mouse cerebellum.

Panel C: anti-P2RY12 staining microglia in mouse cerebellum.

Panel D: anti-FMR1 staining neurons and Purkinje cells in mouse cerebellum.

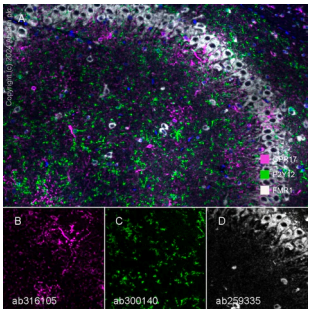
Nuclear DNA was labeled with DAPI (shown in blue).

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded rat hippocampus tissue sections.

Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-FMR1 (gray; Opal™570) on rat hippocampus.

Panel B: anti-GPR17 staining oligodendrocytes in rat hippocampus.

Panel C: anti-P2RY12 staining microglia in rat hippocampus.

Panel D: anti-FMR1 staining neurons in rat hippocampus.

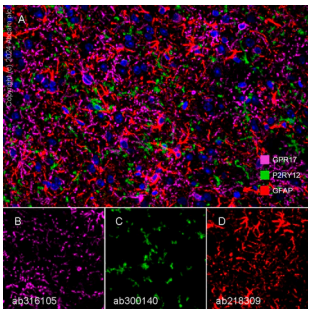
Nuclear DNA was labeled with DAPI (shown in blue).

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/40000 dilution, and ab259335 at a 1/10000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Multiplex immunohistochemistry - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Multiplex immunohistochemistry staining of Mouse cerebrum tissue using rabbit Anti-GPCR GPR17 antibody

Fluorescence multiplex immunohistochemical analysis of formalin/PFA-fixed paraffin-embedded mouse cerebrum tissue sections.
Panel A: merged staining of anti-GPR17 (magenta; Opal™690), anti-P2RY12 (green; Opal™520) and anti-GFAP (red; Opal™570) on mouse cerebrum.

Panel B: anti-GPR17 staining oligodendrocytes in mouse cerebrum.

Panel C: anti-P2RY12 staining microglia in mouse cerebrum.

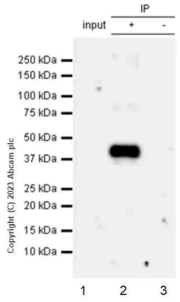
Panel D: anti-GFAP staining astrocytes in mouse cerebrum.

The section was incubated in three rounds of staining: in the order of ab316105 at a 1/500 dilution, ab300140 at a 1/140000 dilution, and ab218309 at a 1/1000 dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. Nuclear counterstaining with DAPI.

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

This data was developed using ab316105, the same antibody clone in a different buffer formulation.



Immunoprecipitation - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.

GPCR GPR17 was immunoprecipitated from 0.35 mg Mouse brain tissue lysate with ab316105 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab316105 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP)(ab131366) was used at 1/5000 dilution.

Lane 1: Mouse brain tissue lysate
Lane 2: ab316105 IP in Mouse brain tissue lysate
Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab316105 in mouse brain tissue lysate.

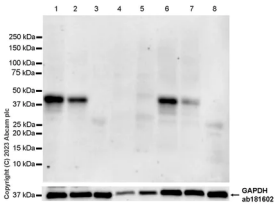
All lanes:
Immunoprecipitation - Anti-GPCR GPR17 antibody [EPR26423-34] (ab316105) at 1/30 dilution

All lanes:
Mouse brain tissue lysate

Secondary

All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 180s



Western blot - Anti-GPCR GPR17 antibody [EPR26423-34] - BSA and Azide free (ab316106)

GPCR GPR17 Western blot staining using rabbit Anti-GPCR GPR17 antibody

This data was developed using ab316105, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.

Negative control: heart, lung and spleen(PMID: 19625605), (PMID: 19838178).

In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.

All lanes:
Western blot - Anti-GPCR GPR17 antibody [EPR26423-34] (ab316105) at 1/1000 dilution

Lane 1:
Mouse P14 spinal cord tissue lysate at 40 µg

Lane 2:
Mouse P14 brain tissue lysate at 40 µg

Lane 3:
Mouse heart tissue lysate at 40 µg

Lane 4:
Mouse lung tissue lysate at 40 µg

Lane 5:
Mouse spleen tissue lysate at 40 µg

Lane 6:
Rat P14 spinal cord tissue lysate at 40 µg

Lane 7:
Rat P14 brain tissue lysate at 40 µg

Lane 8:
Rat heart tissue lysate at 40 µg

Secondary

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

Observed band size: 40 kDa, 36 kDa

Exposure time: 103s

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