abcam

Product datasheet

Alexa Fluor® 647 Anti-COX2 / Cyclooxygenase 2 antibody [EPR12012] ab225273

Recombinant RabMAb

3 Images

Overview

1 References

Product name Alexa Fluor® 647 Anti-COX2 / Cyclooxygenase 2 antibody [EPR12012]

Description Alexa Fluor® 647 Rabbit monoclonal [EPR12012] to COX2 / Cyclooxygenase 2

Host species Rabbit

Conjugation Alexa Fluor® 647. Ex: 652nm, Em: 668nm

Tested applications Suitable for: ICC/IF, IHC-P

Species reactivity Reacts with: Human

Predicted to work with: Mouse

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

Positive control IHC-P: Normal human colon tissue sections ICC/IF: U-87 MG cells

General notes 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**.

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Properties

Form Liquid

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

term. Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

Purity Protein A purified

ClonalityMonoclonalClone numberEPR12012

Isotype IgG

Applications

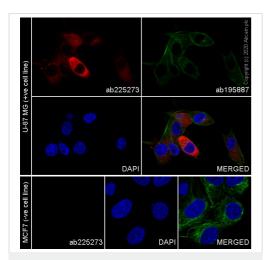
The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab225273 in the following tested applications.

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

| Application | Abreviews | Notes |
|-------------|-----------|---|
| ICC/IF | | Use at an assay dependent concentration. |
| IHC-P | | 1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. |

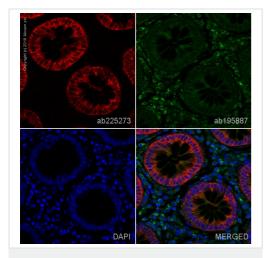
| larget | | |
|----------------------------------|--|--|
| Function | Mediates the formation of prostaglandins from arachidonate. May have a role as a major mediator of inflammation and/or a role for prostanoid signaling in activity-dependent plasticity. | |
| Pathway | Lipid metabolism; prostaglandin biosynthesis. | |
| Sequence similarities | Belongs to the prostaglandin G/H synthase family. Contains 1 EGF-like domain. | |
| Post-translational modifications | S-nitrosylation by NOS2 (iNOS) activates enzme activity. S-nitrosylation may take place on different Cys residues in addition to Cys-561. | |
| Cellular localization | Microsome membrane. Endoplasmic reticulum membrane. | |

Images



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-COX2 / Cyclooxygenase 2 antibody [EPR12012] (ab225273)

Immunocytochemistry/immunofluorescence analysis of U-87 MG (human glioblastoma-astrocytoma epithelial cell) labelling COX2 with ab225273 at 10 µg/mL. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. Cells were counterstained with Anti-alpha Tubulin antibody - Microtubule Marker (Alexa Fluor® 448) (ab195887) at 1/200 dilution (red). Nuclear DNA was labelled with DAPI (blue). MCF7 (human breast adenocarcinoma epithelial cell) was used as a negative control.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 647 Anti-COX2 / Cyclooxygenase 2 antibody [EPR12012] (ab225273)

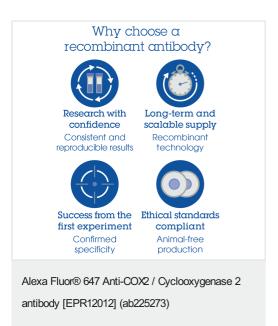
IHC image of COX2 / Cyclooxygenase 2 staining in a section of formalin-fixed paraffin-embedded normal human colon*.

The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins performed on a Leica BOND™. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab225273 at 1/100 dilution (shown in red) and counterstained using ab195887, Mouse monoclonal to alpha Tubulin (Alexa Fluor[®] 488), at 1/250 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue). The section was then mounted using Fluoromount[®].

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

For other IHC staining systems (automated and non-automated), customers should optimize variable parameters such as antigen retrieval conditions, antibody concentrations and incubation times.

*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre.



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