abcam

Product datasheet

Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free ab231859

KO 評価済 リコンビナント RabMAb

1 References 画像数 9

製品の概要

製品名	Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free
製品の詳細	Rabbit monoclonal [EPR19089-202] to MEF2C - BSA and Azide free
由来種	Rabbit
アプリケーション	**適用あり:** Flow Cyt (Intra), WB, ICC/IF, ChIP, IHC-P
種交差性	**交差種:** Mouse, Rat, Human
免疫原	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
ポジティブ・コントロール	WB: Human MEF2C recombinant protein; Raji, Jurkat, Ramos, Daudi, RAW264.7, NIH/3T3,C6, Wild-type THP-1 and Daudi whole cell lysates; mouse and rat brain lysate. IHC-P: Human skeletal muscle and endometrial carcinoma tissue; Mouse and rat spleen tissue. ICC/IF: Raji cells. Flow: Jurkat cells. ChIP: Chromatin prepared from HUVEC cells.
特記事項	ab231859 is the carrier-free version of **ab211493**. 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 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. 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. 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	EPR19089-202
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ab231859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. Detects a band of approximately 50-60 kDa (predicted molecular weight: 51 kDa).
ICC/IF		Use at an assay dependent concentration.
ChIP		Use at an assay dependent concentration.
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.

ターゲット情報

機能	Transcription activator which binds specifically to the MEF2 element present in the regulatory regions of many muscle-specific genes. Controls cardiac morphogenesis and myogenesis, and is also involved in vascular development. Plays an essential role in hippocampal-dependent learning and memory by suppressing the number of excitatory synapses and thus regulating basal and evoked synaptic transmission. Crucial for normal neuronal development, distribution, and electrical activity in the neocortex. Necessary for proper development of megakaryocytes and platelets and for bone marrow B lymphopoiesis. Required for B-cell survival and proliferation in response to BCR stimulation, efficient IgG1 antibody responses to T-cell-dependent antigens and for normal induction of germinal center B cells. May also be involved in neurogenesis and in the development of cortical architecture (By similarity). Isoform 3 and isoform 4, which lack the repressor domain, are more active than isoform 1 and isoform 2.
組織特異性	Expressed in brain and skeletal muscle.
関連疾患	Defects in MEF2C are the cause of mental retardation-stereotypic movements-epilepsy and/or cerebral malformations (MRSME) [MIM:613443]. It is a disorder characterized by severe mental retardation, absent speech, hypotonia, poor eye contact and stereotypic movements. Dysmorphic

features include high broad forehead with variable small chin, short nose with anteverted nares, large open mouth, upslanted palpebral fissures and prominent eyebrows. Some patients have seizures.

配列類似性

Belongs to the MEF2 family.
Contains 1 MADS-box domain.
Contains 1 Mef2-type DNA-binding domain.

発生段階

Expression is highest during the early stages of postnatal development, at later stages levels greatly decrease.

ドメイン

The beta domain, missing in a number of isoforms, is required for enhancement of transcriptional activity.

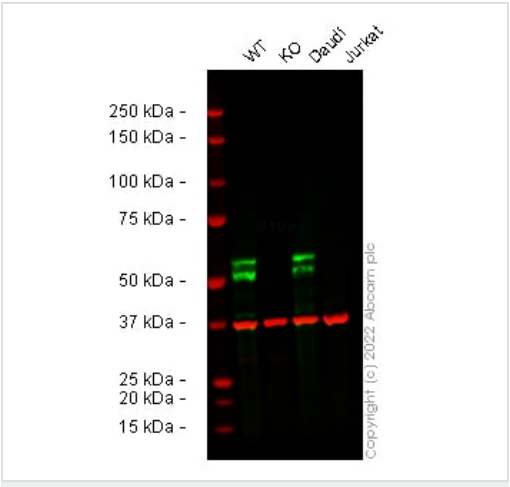
翻訳後修飾

Phosphorylation on Ser-59 enhances DNA binding activity (By similarity). Phosphorylation on Ser-396 is required for Lys-391 sumoylation and inhibits transcriptional activity.
Acetylated by p300 on several sites in diffentiating myocytes. Acetylation on Lys-4 increases DNA binding and transactivation.
Sumoylated on Lys-391 by SUMO2 but not by SUMO1 represses transcriptional activity.
Proteolytically cleaved in cerebellar granule neurons, probably by caspase 7, following neurotoxicity. Preferentially cleaves the CDK5-mediated hyperphosphorylated form which leads to neuron apoptosis and transcriptional inactivation.

細胞内局在

Nucleus.

画像



Western blot - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

All lanes : Anti-MEF2C antibody [EPR19089-202] - ChIP Grade (**ab211493**) at 1/1000 dilution

Lane 1 : Wild-type THP-1 cell lysate
Lane 2 : MEF2C knockout THP-1 cell lysate
Lane 3 : Daudi cell lysate
Lane 4 : Jurkat cell lysate

Lysates/proteins at 20 µg per lane.

Secondary
All lanes : Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution

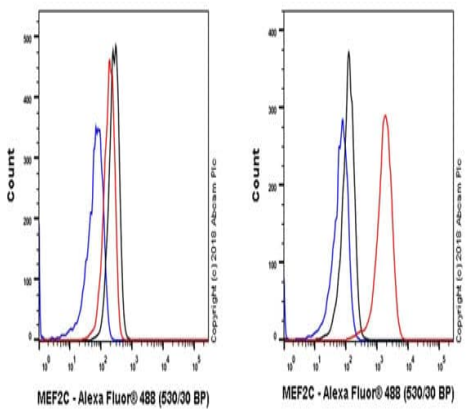
Performed under reducing conditions.

Predicted band size: 51 kDa
Observed band size: 55/60 kDa

False colour image of Western blot: Anti-MEF2C antibody [EPR19089-202] - ChIP Grade 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, **ab211493** was shown to bind specifically to MEF2C. A band was observed at 55/60 kDa in wild-type THP-1 cell lysates with no signal observed at this size in MEF2C knockout cell line. To generate this image, wild-type and MEF2C knockout THP-1 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween® 20 (TBS-T) 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. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

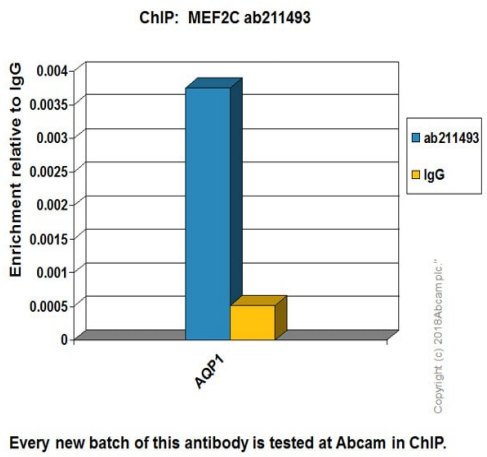


Flow Cytometry (Intracellular) - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol permeabilsed Jurkat (Human T cell leukemia T lymphocyte, Left) / Raji (Human Burkitt's lymphoma B lymphocyte, Right) cell lines labelling MEF2C with **ab211493** at 1/500 dilution (Red) compared with the isotype control Rabbit monoclonal IgG (**ab172730**) (Black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). Goat anti rabbit IgG Alexa Fluor® 488 (**ab150077**), at 1/2000 dilution was used as the secondary antibody.

Negative control: Jurkat (PMID: 27876533).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

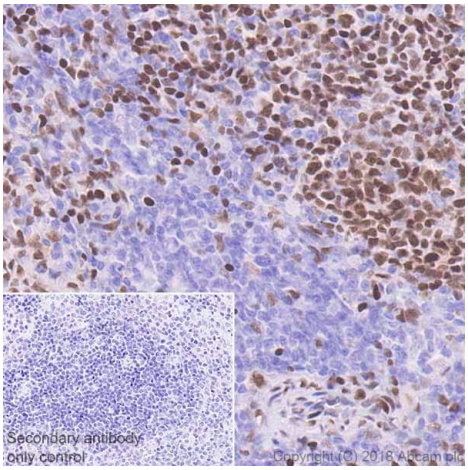


ChIP - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Chromatin was prepared from HUVEC cells according to the Abcam X-ChIP protocol. Cells were fixed with formaldehyde for 10 minutes. The ChIP was performed with: 25 µg of chromatin, 5 µg of **ab211493** (blue), and 20 µl of protein A/G sepharose beads slurry (10 µl of sepharose A beads + 10 µl of sepharose G beads). 5 µg of rabbit normal IgG was added to the beads control (yellow). The immunoprecipitated DNA was quantified by real time PCR (SYBR green chemistry).

ChIP was performed according to the literature (PMID: 26923194).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

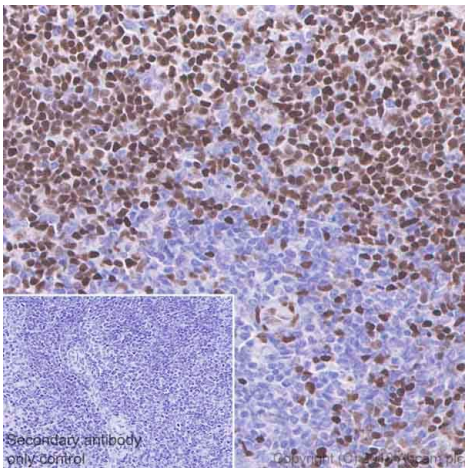


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Immunohistochemical analysis of paraffin-embedded rat spleen tissue labelling MEF2C with **ab211493** at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ready to use. Nuclear staining in B lymphocytes of rat spleen but not in T cells in the periarterial lymphatic sheath is observed (PMID: 8506376, PMID 15703219). Counterstained with hematoxylin.
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP)ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.

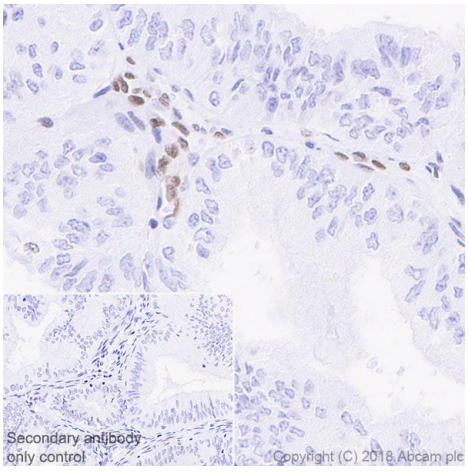


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Immunohistochemical analysis of paraffin-embedded mouse spleen tissue labelling MEF2C with **ab211493** at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ready to use. Nuclear staining in B lymphocytes of mouse spleen but not in T cells in the periarterial lymphatic sheath is observed (PMID: 8506376, PMID 15703219). Counterstained with hematoxylin.
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP)ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.

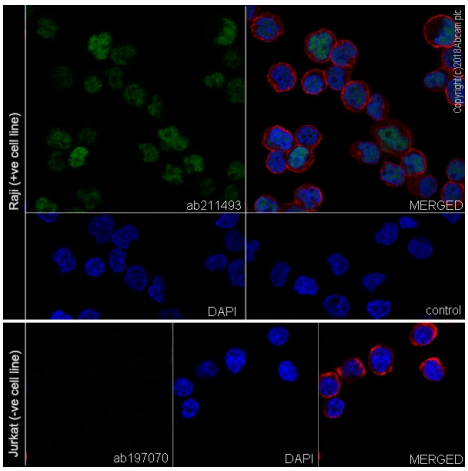


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Immunohistochemical analysis of paraffin-embedded human endometrial carcinoma tissue labelling MEF2C with **ab211493** at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ready to use. Nuclear staining in leukocytes but not in tumor cells of human endometrial carcinoma is observed (PMID: 8506376, PMID 15703219). Counterstained with hematoxylin.
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP)ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.

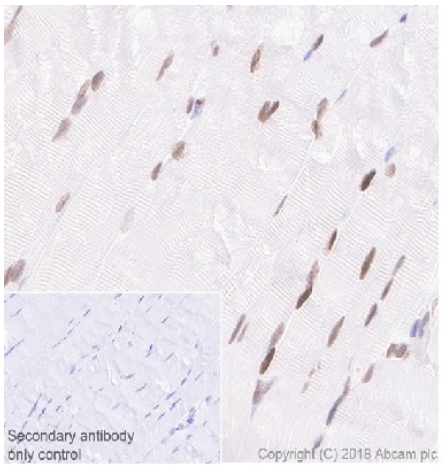


Immunocytochemistry/ Immunofluorescence - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilised Raji (human Burkitt's lymphoma B lymphocyte) cells labelling MEF2C with **ab211493** at 1/500 dilution, followed by AlexaFluor®488 Goat anti-Rabbit secondary (**ab150077**) at 1/1000 dilution (green). Confocal image showing nuclear staining in Raji cell line. **Negative control:** Jurkat (PMID: 27876533). DAPI was used as the nuclear counterstain, and the Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) (**ab195889** antibody was used as a counterstain at 1/200 dilution.

The negetive controls are as follows:
-ve control 1: **ab197070** on jurkat (human T cell leukemia cell line from peripheral blood) cells.
-ve control 2: Jurkat cells stained with DAPI.
-ve control 3: Merged negetive contol images.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

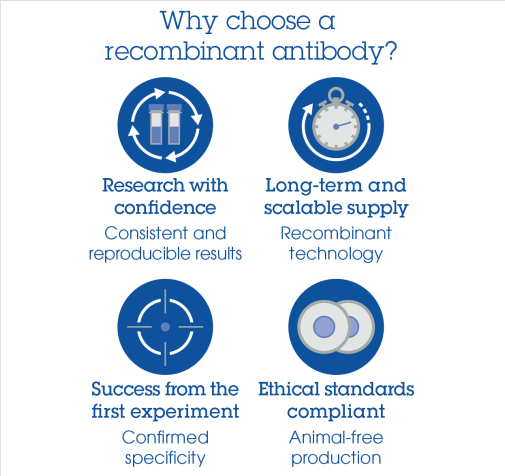


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

Immunohistochemical analysis of paraffin-embedded human skeletal muscle tissue labelling MEF2C with **ab211493** at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ready to use. Nuclear staining in human skeletal muscle cells is observed(PMID: 8506376, PMID 15703219). Counterstained with hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ready to use.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab211493**).

Perform heat mediated antigen retrieval with EDTA buffer pH 9 before commencing with IHC staining protocol.



Anti-MEF2C antibody [EPR19089-202] - BSA and Azide free (ab231859)

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