abcam

Product datasheet

Anti-PRKAR1A (phospho S77) antibody [EPMAYR1-127] ab139682



1 References 2 Images

Overview

Product name Anti-PRKAR1A (phospho S77) antibody [EPMAYR1-127]

Description Rabbit monoclonal [EPMAYR1-127] to PRKAR1A (phospho S77)

Host species Rabbit

Specificity ab139682 only detects Protein Kinase A regulatory subunit I alpha/PRKAR1A phosphorylated at

serine 77.

Tested applications Suitable for: WB

Unsuitable for: ICC/IF or IHC-P

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

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

Positive control Fetal heart tissue lysate.

General notes This antibody was developed as part of a collaboration between Abcam and the lab of Dr. Ozgur

Ogut at the Mayo Clinic.

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**.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

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Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

Purity Tissue culture supernatant

Clonality Monoclonal

Clone number EPMAYR1-127

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab139682 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
WB		1/1000 - 1/10000. Predicted molecular weight: 43 kDa.

Application notes Is unsuitable for ICC/IF or IHC-P.

Target

Tissue specificity Four types of regulatory chains are found: I-alpha, I-beta, II-alpha, and II-beta. Their expression

varies among tissues and is in some cases constitutive and in others inducible.

Involvement in disease Defects in PRKAR1A are the cause of Carney complex type 1 (CNC1) [MIM:160980]. CNC is a

multiple neoplasia syndrome characterized by spotty skin pigmentation, cardiac and other

myxomas, endocrine tumors, and psammomatous melanotic schwannomas.

Defects in PRKAR1A are the cause of intracardiac myxoma (INTMYX) [MIM:255960]. Inheritance

is autosomal recessive.

Defects in PRKAR1A are the cause of primary pigmented nodular adrenocortical disease type 1 (PPNAD1) [MIM:610489]. Primary pigmented nodular adrenocortical disease is a rare bilateral adrenal defect causing ACTH-independent Cushing syndrome. Macroscopic appearance of the adrenals is characteristic with small pigmented micronodules observed in the cortex. PPNAD1 is most often diagnosed in patients with Carney complex, but it can also be observed in patients

without other manifestations or familial history.

Sequence similarities Belongs to the cAMP-dependent kinase regulatory chain family.

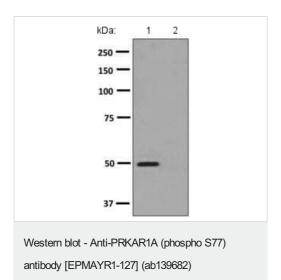
Contains 2 cyclic nucleotide-binding domains.

Post-translational modifications

The pseudophosphorylation site binds to the substrate-binding region of the catalytic chain,

resulting in the inhibition of its activity.

Images



All lanes : Anti-PRKAR1A (phospho S77) antibody [EPMAYR1-127] (ab139682) at 1/1000 dilution

Lane 1: Fetal heart lysate

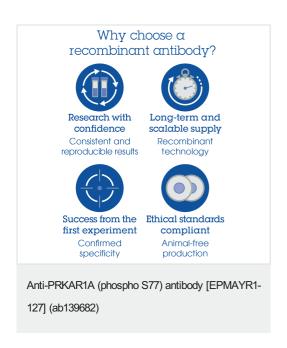
Lane 2: Fetal heart lysate treated with Alkaline Phosphatase

Lysates/proteins at 10 µg per lane.

Secondary

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

Predicted band size: 43 kDa



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