abcam

Product datasheet

Anti-PAX7 antibody [PAX7/1187] ab218472

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Overview

Product name Anti-PAX7 antibody [PAX7/1187]

Description Mouse monoclonal [PAX7/1187] to PAX7

Host species Mouse

Tested applications Suitable for: IHC-P

Species reactivity Reacts with: Human

Immunogen Recombinant fragment within Human PAX7 aa 301-505. The exact sequence is proprietary.

Database link: P23759

Positive control IHC-P: Human rhabdomyosarcoma and testicular carcinoma tissues.

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or

contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

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.

Storage buffer pH: 7.2

Preservative: 0.05% Sodium azide Constituents: 99% PBS, 0.05% BSA

Purity Protein G purified

Clonality Monoclonal
Clone number PAX7/1187

Light chain type lgG1 kappa

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Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab218472 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
IHC-P	*****(1)	Use a concentration of 1 - 2 µg/ml. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. Primary incubation is for 30 min at RT.

Target

Function

Probable transcription factor. May have a role in myogenesis.

Involvement in disease

Defects in PAX7 are a cause of rhabdomyosarcoma type 2 (RMS2) [MIM:268220]. It is a form of rhabdomyosarcoma, a highly malignant tumor of striated muscle derived from primitive mesenchimal cells and exhibiting differentiation along rhabdomyoblastic lines.

Rhabdomyosarcoma is one of the most frequently occurring soft tissue sarcomas and the most common in children. It occurs in four forms: alveolar, pleomorphic, embryonal and botryoidal

common in children. It occurs in four forms: alveolar, pleomorphic, embryonal and botryoidal rhabdomyosarcomas. Note=A chromosomal aberration involving PAX7 is found in rhabdomyosarcoma. Translocation t(1;13)(p36;q14) with FOXO1. The resulting protein is a

transcriptional activator.

Sequence similarities

Belongs to the paired homeobox family.

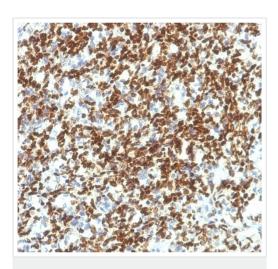
Contains 1 homeobox DNA-binding domain.

Contains 1 paired domain.

Cellular localization

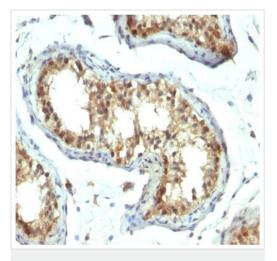
Nucleus.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PAX7 antibody
[PAX7/1187] (ab218472)

Immunohistochemical analysis of formalin-fixed, paraffin-embedded human rhabdomyosarcoma tissue section labeling PAX7 with ab218472 at 1 μ g/mL.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PAX7 antibody
[PAX7/1187] (ab218472)

Immunohistochemical analysis of formalin-fixed, paraffin-embedded human testicular carcinoma tissue section labeling PAX7 with ab218472 at 1 μ g/mL.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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