# abcam

# Product datasheet

# HRP Anti-FABP4 antibody [EPR3579] ab216528

Recombinant

RabMAb

## 3 Images

#### Overview

Product name HRP Anti-FABP4 antibody [EPR3579]

**Description** HRP Rabbit monoclonal [EPR3579] to FABP4

Host species Rabbit
Conjugation HRP

Tested applications
Suitable for: IHC-P, WB
Species reactivity
Reacts with: Human

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

Positive control WB: Human adipose, human heart. IHC-P: normal human breast tissue sections

**General notes**This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

Improved sensitivity and specificityLong-term security of supplyAnimal-free production

For more information see here.

Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

#### **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: 0.1% Proclin 300 Solution

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

Purity Protein A purified

Clonality Monoclonal
Clone number EPR3579

**Isotype** IgG

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#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab216528 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		1/50. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		1/5000. Detects a band of approximately 15 kDa (predicted molecular weight: 15 kDa).

#### **Target**

Function Lipid transport protein in adipocytes. Binds both long chain fatty acids and retinoic acid. Delivers

long-chain fatty acids and retinoic acid to their cognate receptors in the nucleus.

**Sequence similarities**Belongs to the calycin superfamily. Fatty-acid binding protein (FABP) family.

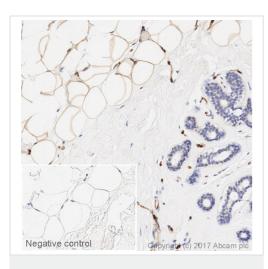
**Domain** Forms a beta-barrel structure that accommodates hydrophobic ligands in its interior.

**Cellular localization**Cytoplasm. Nucleus. Depending on the nature of the ligand, a conformation change exposes a

nuclear localization motif and the protein is transported into the nucleus. Subject to constitutive

nuclear export.

#### **Images**

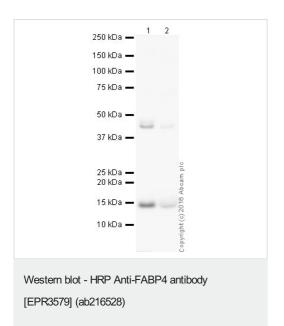


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - HRP Anti-FABP4 antibody
[EPR3579] (ab216528)

IHC image of FABP4 staining in a section of formalin-fixed paraffinembedded normal human breast\*, performed on a Leica BOND™. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20mins. The section was then incubated with ab216528, 1/50 dilution, for 15 mins at room temperature. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. The inset negative control image is taken from an identical assay without primary antibody.

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

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



**All lanes :** HRP Anti-FABP4 antibody [EPR3579] (ab216528) at 1/5000 dilution

Lane 1: Human Adipose

Lane 2: Heart (Human) Tissue Lysate - adult normal tissue

Lysates/proteins at 10 µg per lane.

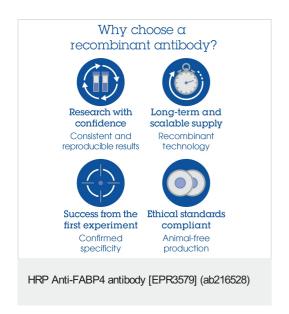
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 15 kDa
Observed band size: 15 kDa

Exposure time: 5 seconds

This blot was produced using a 4-12% Bis-tris gel under the MES buffer system. The gel was run at 200V for 35 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab216528 overnight at 4°C. Antibody binding was visualised using ECL development solution ab133406.



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

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