# abcam

# Product datasheet

# Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free ab238957



# 7 Images

#### Overview

**Product name** Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide

free

**Description** Rabbit monoclonal [EP2122Y] to Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) - BSA

and Azide free

**Host species** Rabbit

Specificity ab238957 will detect Ezrin, Radixin and Moesin. This antibody detects the phosphorylated target

and not the non-phosphorylated epitope.

Suitable for: WB, IHC-P, Dot blot **Tested applications** 

Unsuitable for: Flow Cyt

Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control IHC: human breast adenocarcinoma.

**General notes** ab238957 is the carrier-free version of ab76247.

> 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 cellbased 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<sup>®</sup> is a trademark of Fluidigm Canada Inc.

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<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**.

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. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EP2122Y

**Isotype** IgG

## **Applications**

# The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab238957 in the following tested applications.

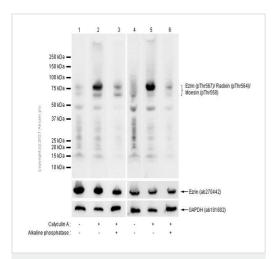
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Detects a band of approximately 75-81 kDa (predicted molecular weight: 69 kDa).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
Dot blot		Use at an assay dependent concentration.

**Application notes** 

Is unsuitable for Flow Cyt.

## **Images**



Western blot - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free (ab238957)

**All lanes:** Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] (ab76247) at 1/1000 dilution

**Lane 1 :** Untreated HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

Lane 2: HeLa (Human cervix adenocarcinoma epithelial cell) starved overnight, then treated with 100nM Calyculin A for 30 minutes whole cell lysate

Lane 3: HeLa (Human cervix adenocarcinoma epithelial cell) starved overnight, then treated with 100nM Calyculin A for 30 minutes whole cell lysate, then the membrane treated with Alkaline Phosphatase for 1 hour

**Lane 4 :** Untreated HEK-293 (Human embryonic kidney epithelial cell) whole cell lysate

**Lane 5**: EK-293 (Human embryonic kidney epithelial cell) starved overnight, then treated with 100nM Calyculin A for 60 minutes whole cell lysate

**Lane 6**: HEK-293 (Human embryonic kidney epithelial cell) starved overnight, then treated with 100nM Calyculin A for g0 minutes whole cell lysate, then the membrane treated with Alkaline Phosphatase for 1 hour

Lysates/proteins at 15 µg per lane.

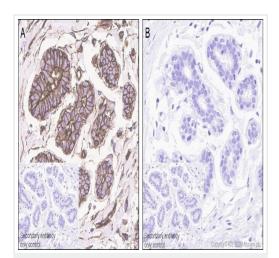
#### Secondary

**All lanes :** Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

**Predicted band size:** 69 kDa **Observed band size:** 75-81 kDa

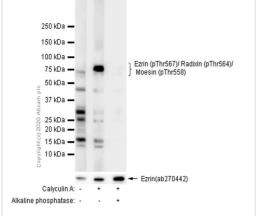
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Blocking and dilution buffer: 5% NFDM/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffin-

embedded sections) - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] -BSA and Azide free (ab238957) 250 kDa -Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558)



Western blot - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] -BSA and Azide free (ab238957)

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Immunohistochemical analysis of paraffin-embedded human breast tissue labeling Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) with ab76247 at 1/500 (1.64 µg/ml) dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Membranous staining on human breast.

The section was incubated with **ab76247** for 30 mins at room temperature. The immunostaining staining was performed on a Leica Biosystems BOND® RX instrument.

Secondary antibody only control: Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.

Counterstained with Hematoxylin.

All lanes: Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] (ab76247) at 1/1000 dilution

Lane 1: Untreated NIH/3T3 (Mouse embyonic fibroblast) whole cell lysate

Lane 2: Untreated NIH/3T3 (Mouse embyonic fibroblast) treated with 100nM Calyculin A for 30 minutes whole cell lysate

Lane 3: Untreated NIH/3T3 (Mouse embyonic fibroblast) treated with 100nM Calyculin A for 30 minutes whole cell lysate, then the membrane treated with Alkaline Phosphatase for 1 hour,

Lysates/proteins at 15 µg per lane.

# Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 69 kDa Observed band size: 75-81 kDa This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Blocking and dilution buffer: 5% NFDM/TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Immunohistochemical analysis of paraffin-embedded mouse testis tissue labeling Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) with <u>ab76247</u> at 1/500 (1.64 µg/ml) dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Membranous staining on human breast.

The section was incubated with <u>ab76247</u> for 30 mins at room temperature. The immunostaining staining was performed on a Leica Biosystems BOND<sup>®</sup> RX instrument.

Secondary antibody only control: Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.

Counterstained with Hematoxylin.

**All lanes :** Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] (ab76247) at 1/1000 dilution

Lane 1 : Untreated C6 (Rat glial tumor glial cell) whole cell lysate

Lane 2: Untreated C6 (Rat glial tumor glial cell) treated with

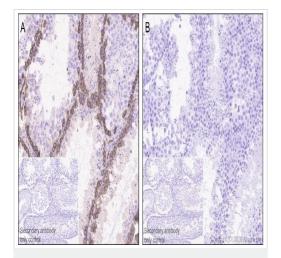
100ng/ml Calyculin A for 60 minutes whole cell lysate

Lane 3: Untreated C6 (Rat glial tumor glial cell) treated with 100ng/ml Calyculin A for 60 minutes whole cell lysate, then the membrane treated with Alkaline Phosphatase for 1 hour,

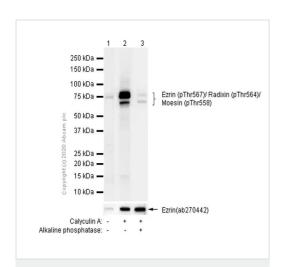
Lysates/proteins at 15  $\mu g$  per lane.

# antibody [EP2122V] - Secondary

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free (ab238957)



Western blot - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free (ab238957)

**Predicted band size:** 69 kDa **Observed band size:** 75-81 kDa

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Blocking and dilution buffer: 5% NFDM/TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

Immunohistochemical analysis of paraffin-embedded rat liver tissue labeling Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) with <u>ab76247</u> at 1/500 (1.64  $\mu$ g/ml) dilution followed by ready to use Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

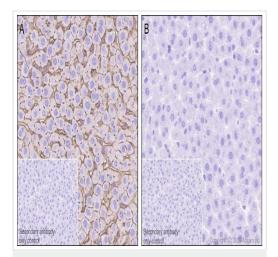
Membranous staining on human breast.

The section was incubated with <u>ab76247</u> for 30 mins at room temperature. The immunostaining staining was performed on a Leica Biosystems BOND<sup>®</sup> RX instrument.

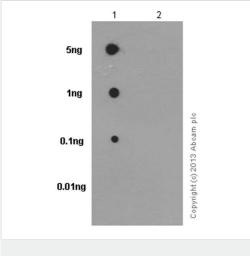
Secondary antibody only control: Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 20 mins.

Counterstained with Hematoxylin.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free (ab238957)



Dot Blot - Anti-Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) antibody [EP2122Y] - BSA and Azide free (ab238957) Dot blot analysis of Ezrin (pThr567)/ Radixin (pThr564)/ Moesin (pThr558) phospho peptide (Lane 1), Non-phospho peptide (Lane 2), labelling Ezrin (phospho Thr567)/ Radixin (phospho Thr564)/ Moesin (phospho Thr558) with <a href="mailto:ab76247">ab76247</a> at a dilution of 1/1000. Peroxidase conjugated goat anti-rabbit lgG (H+L) was used as the secondary antibody at a dilution of 1/2500.

Blocking and diluting buffer: 5% NFDM/TBST.

Exposure time: 3 minutes.

The sequence of non-phospho peptide: CGRDKYKTLRQIR. The sequence of phospho peptide: CGRDKYK-pT-LRQIR.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab76247).

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

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