# abcam

#### Product datasheet

## Alexa Fluor® 647 Anti-ERK1 antibody [Y72] ab190579



Recombinant RabMAb

### 3 Images

#### Overview

**Product name** Alexa Fluor® 647 Anti-ERK1 antibody [Y72]

**Description** Alexa Fluor® 647 Rabbit monoclonal [Y72] to ERK1

**Host species** Rabbit

Alexa Fluor® 647. Ex: 652nm, Em: 668nm Conjugation

**Tested applications** Suitable for: Flow Cyt (Intra)

Species reactivity Reacts with: Human

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

ab32537 reacts with an epitope located in the N terminal region of ERK1. **Epitope** 

Positive control Flow Cyt (intra): HeLa cells and wildtype HAP1 cells.

**General notes** 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**.

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#### **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.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

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

**Purity** Protein A purified

**Clonality** Monoclonal

Clone number Y72 Isotype IgG

#### **Applications**

**Images** 

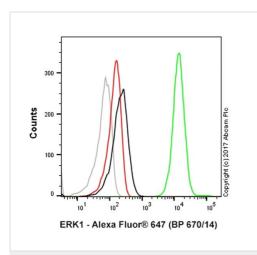
The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab190579 in the following tested applications.

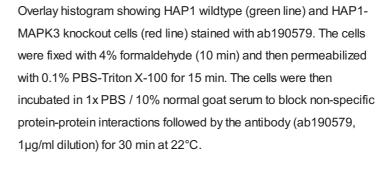
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt (Intra)		Use a concentration of 1 µg/ml.

Target		
Function	Involved in both the initiation and regulation of meiosis, mitosis, and postmitotic functions in differentiated cells by phosphorylating a number of transcription factors such as ELK-1. Phosphorylates EIF4EBP1; required for initiation of translation. Phosphorylates microtubule-associated protein 2 (MAP2). Phosphorylates SPZ1 (By similarity). Phosphorylates heat shock factor protein 4 (HSF4).	
Sequence similarities	Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. MAP kinase subfamily.  Contains 1 protein kinase domain.	
Domain	The TXY motif contains the threonine and tyrosine residues whose phosphorylation activates the MAP kinases.	
Post-translational modifications	Dually phosphorylated on Thr-202 and Tyr-204, which activates the enzyme. Dephosphorylated by PTPRJ at Tyr-204.	



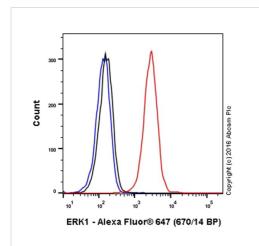
Flow Cytometry (Intracellular) - Alexa Fluor® 647 Anti-ERK1 antibody [Y72] (ab190579)



A rabbit monoclonal IgG isotype control antibody (<u>ab199093</u>) was used at the same concentration and conditions as the primary antibody (HAP1 wildtype - black line, HAP1-MAPK3 knockout - grey line). Unlabelled sample was also used as a control (this line is not shown for the purpose of simplicity).

Acquisition of >5,000 events were collected using a 40 mW red laser (640nm) and 670/14 bandpass filter.

This antibody can also be used in HAP1 cells fixed with 80% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 15 min under the same conditions.



Flow Cytometry (Intracellular) - Alexa Fluor® 647 Anti-ERK1 antibody [Y72] (ab190579)

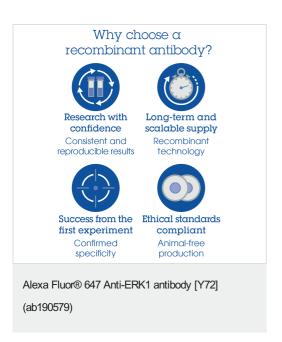
Overlay histogram showing HeLa cells stained with ab190579 (red line). The cells were fixed with 4% formaldehyde and then permeabilized with 0.1% PBS-Triton X-100 for 15 min.

The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab190579, 1/5000 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was Rabbit IgG (monoclonal) Alexa Fluor<sup>®</sup> 647 (<u>ab199093</u>) used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 40 mW Red laser (640nm) and 670/14 bandpass filter.

This antibody gave a positive signal in HeLa cells fixed with 80% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 15 min used under the same conditions.



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

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