

# **Product datasheet**

# Alexa Fluor® 647 Anti-EEA1 antibody [EPR4245] - Early Endosome Marker ab196186

Recombinant RabMAb

## **<u>1 References</u>** 3 Images

Overview	
Product name	Alexa Fluor® 647 Anti-EEA1 antibody [EPR4245] - Early Endosome Marker
Description	Alexa Fluor® 647 Rabbit monoclonal [EPR4245] to EEA1 - Early Endosome Marker
Host species	Rabbit
Conjugation	Alexa Fluor® 647. Ex: 652nm, Em: 668nm
Tested applications	Suitable for: Flow Cyt, ICC/IF
Species reactivity	Reacts with: Human
Immunogen	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
Positive control	ICC/IF: Jurkat cells. Flow Cyt: Jurkat cells.
General notes	This product is a recombinant monoclonal antibody, which offers several advantages including:
	<ul> <li>High batch-to-batch consistency and reproducibility</li> <li>Improved sensitivity and specificity</li> <li>Long-term security of supply</li> <li>Animal-free production</li> <li>For more information <u>see here</u>.</li> <li>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb<sup>®</sup> patents</u>.</li> <li>Alexa Fluor<sup>®</sup> is a registered trademark of Molecular Probes, Inc, a Thermo Fisher Scientific Company. The Alexa Fluor<sup>®</sup> dye included in this product is provided under an intellectual property license from Life Technologies Corporation. As this product contains the Alexa Fluor<sup>®</sup> dye, the purchase of this product conveys to the buyer the non-transferable right to use the purchased product and components of the product only in research conducted by the buyer (whether the buyer is an academic or for-profit entity). As this product contains the Alexa Fluor<sup>®</sup> dye the sale of this product is expressly conditioned on the buyer not using the product or its components, or any materials made using the product or its components, in any activity to generate revenue, which may include, but is not limited to use of the product or its components: in manufacturing; (ii) to provide a service, information, or data in return for payment (iii) for therapeutic, diagnostic or prophylactic purposes; or (iv) for resale, regardless of whether they are sold for use in research. For information on purchasing a license to this product for purposes other than research, contact Life Technologies Corporation, 5781 Van Allen Way, Carlsbad, CA 92008 USA or</li> </ul>

### 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. Store In the Dark.
Storage buffer	pH: 7.40 Preservative: 0.02% Sodium azide Constituents: 30% Glycerol (glycerin, glycerine), PBS, 1% BSA
Purity	Protein A purified
Clonality	Monoclonal
Clone number	EPR4245
lsotype	lgG

#### Applications

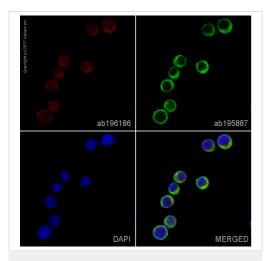
The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab196186 in the following tested applications.

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

Application	Abreviews	Notes
Flow Cyt		1/50. <u>ab199093</u> - Rabbit monoclonal IgG (Alexa Fluor® 647), is suitable for use as an isotype control with this antibody.
ICC/IF		1/50 - 1/167. This product gave a positive signal in Jurkat cells fixed with 80% methanol (5 min)

FunctionBinds phospholipid vesicles containing phosphatidylinositol 3-phosphate and participates in endosomal trafficking.Sequence similaritiesContains 1 C2H2-type zinc finger. Contains 1 FYVE-type zinc finger.DomainThe FYVE-type zinc finger domain mediates interactions with phosphatidylinositol 3-phosphate in membranes of early endosomes and penetrates bilayers. The FYVE domain insertion into Ptdlns(3)P-enriched membranes is substantially increased in acidic conditions.Cellular localizationCytoplasm. Early endosome membrane.	Target	
Contains 1 FYVE-type zinc finger.         Domain         The FYVE-type zinc finger domain mediates interactions with phosphatidylinositol 3-phosphate in membranes of early endosomes and penetrates bilayers. The FYVE domain insertion into PtdIns(3)P-enriched membranes is substantially increased in acidic conditions.	Function	
membranes of early endosomes and penetrates bilayers. The FYVE domain insertion into Ptdlns(3)P-enriched membranes is substantially increased in acidic conditions.	Sequence similarities	
Cellular localization         Cytoplasm. Early endosome membrane.	Domain	membranes of early endosomes and penetrates bilayers. The FYVE domain insertion into
	Cellular localization	Cytoplasm. Early endosome membrane.

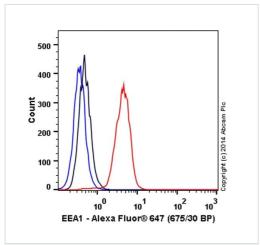
#### Images



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-EEA1 antibody [EPR4245] - Early Endosome Marker (ab196186)

ab196186 staining EEA1 in Jurkat cells. The cells were fixed with 80% methanol (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab196186 at 1/167 dilution (shown in red) and **ab195887**, Mouse monoclonal to alpha Tubulin (Alexa Fluor<sup>®</sup> 488), at 1/250 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



Flow Cytometry - Alexa Fluor® 647 Anti-EEA1 antibody [EPR4245] - Early Endosome Marker (ab196186) Overlay histogram showing Jurkat cells stained with ab196186 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab196186, 1/50 dilution) for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) Alexa Fluor® 647 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 solid-state 25mW red diode laser (635 nm) and 675/30 bandpass filter.

This antibody gave a positive signal in Jurkat cells fixed with 4% formaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



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