### abcam

#### Product datasheet

# Alexa Fluor® 647 Anti-BST2/Tetherin antibody [EPR23597-266] ab275193

Recombinant

#### 1 Image

#### Overview

Product name Alexa Fluor® 647 Anti-BST2/Tetherin antibody [EPR23597-266]

**Description** Alexa Fluor® 647 Rabbit monoclonal [EPR23597-266] to BST2/Tetherin

Host species Rabbit

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

Tested applications Suitable for: Flow Cyt
Species reactivity Reacts with: Mouse

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

**Positive control** Flow cyt: C57 BL/6 mouse splenocytes.

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<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**.

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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 long

term. Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, 68.98% PBS

Purity Protein A purified

**Clonality** Monoclonal

Clone number EPR23597-266

**Isotype** IgG

#### **Applications**

**Target** 

The Abpromise guarantee Our Abpromise guarantee covers the use of ab275193 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		1/500.

## **Function** May be involved in the sorting of secreted proteins (By similarity). May be involved in pre-B-cell growth. Antiretroviral defense protein, that blocks release of retrovirus from the cell surface.

Depleted unpon HIV-1 infection by viral VPU protein through 20S proteasome degradation.

Depleted upon infection by human Kaposi's sarcoma-associated herpesvirus (KSHV) through ubiquitination and subsequent degradation. May play a role in B-cell activation in rheumatoid

arthritis.

Tissue specificity Predominantly expressed in liver, lung, heart and placenta. Lower levels in pancreas, kidney,

skeletal muscle and brain. Overexpressed in multiple myeloma cells. Highly expressed during B-cell development, from pro-B precursors to plasma cells. Highly expressed on T-cells, monocytes,

NK cells and dendritic cells (at protein level).

**Sequence similarities** Belongs to the tetherin family.

**Domain** The extracellular coiled coil domain is important for virus retention at the cell surface and

prevention of virus spreading.

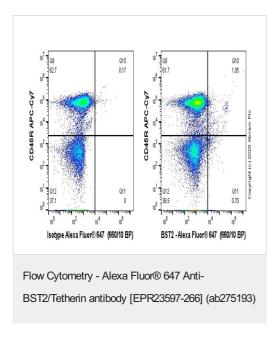
**Post-translational** Monoubiquitinated by KSHV E3 ubiquitin-protein ligase K5, leading to its targeting to late

**modifications** endosomes and degradation.

**Cellular localization** Golgi apparatus > trans-Golgi network. Cell membrane. Cell membrane. Late endosome.

Targeted to late endosomes upon KSHV infection and subsequent ubiquitination. Targeted to the

trans-Golgi network by viral VPU protein.



Flow cytometry staining of C57 BL/6 mouse splenocytes with ab275193 (right) or Rabbit IgG (monoclonal) Alexa Fluor 647 isotype (left). Cells were incubated for 30 min on ice in 1x PBS containing 10  $\mu$ g/mL anti CD16/CD32 and 10 % normal goat serum to block FC receptors and non-specific protein-protein interaction followed by the antibody (ab275193) or Rabbit IgG (monoclonal) Alexa Fluor 647 isotype (1x 106 in 100 $\mu$ L at 1  $\mu$ g/mL (1/500)) for 30 min on ice. The cells were simultaneously stained with CD45R. Acquisition of >30000 events were collected using a 40 mW Red laser (638nm) and 660/10 bandpass filter. Events were gated on live single cells.

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

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