# abcam

# Product datasheet

# Anti-MALT1/MLT antibody [EPR604] - BSA and Azide free ab247917

יובעדער RabMAb

画像数 2

#### 製品の概要

特記事項

製品名 Anti-MALT1/MLT antibody [EPR604] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR604] to MALT1/MLT - BSA and Azide free

由来種 Rabbit

アプリケーション **適用あり:** WB

適用なし: Flow Cyt,IHC-P or IP

種交差性 交差種: Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ab247917 is the carrier-free version of ab109724.

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

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

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#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

バッファー pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

**ポリ/モノ** モノクローナル

**クローン名** EPR604

アイソタイプ lgG

#### アプリケーション

**The Abpromise guarantee** <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab247917の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB		Use at an assay dependent concentration. Predicted molecular weight: 92 kDa.

追加情報 Is unsuitable for Flow Cyt,IHC-P or IP.

## ターゲット情報

機能 Enhances BCL10-induced activation of NF-kappa-B. Involved in nuclear export of BCL10. Binds to TRAF6, inducing TRAF6 oligomerization and activation of its ligase activity. Has ubiquitin

ligase activity. MALT1-dependent BCL10 cleavage plays an important role in T-cell antigen

receptor-induced integrin adhesion.

組織特異性 Highly expressed in peripheral blood mononuclear cells. Detected at lower levels in bone marrow.

thymus and lymph node, and at very low levels in colon and lung.

関連疾患 Note=A chromosomal aberration involving MALT1 is recurrent in low-grade mucosa-associated

lymphoid tissue (MALT lymphoma). Translocation t(11;18)(q21;q21) with BIRC2. This translocation is found in approximately 50% of cytogenetically abnormal low-grade MALT

lymphoma.

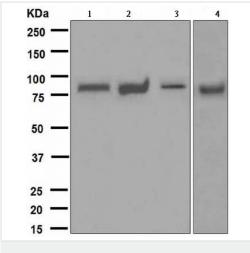
配列類似性 Belongs to the peptidase C14B family.

Contains 1 death domain.

Contains 2 lg-like C2-type (immunoglobulin-like) domains.

細胞内局在 Cytoplasm > perinuclear region. Nucleus. Shuttles between the nucleus and cytoplasm. Found in

perinuclear structures together with BCL10.



Western blot - Anti-MALT1/MLT antibody [EPR604] - BSA and Azide free (ab247917)

**All lanes :** Anti-MALT1/MLT antibody [EPR604] (<u>ab109724</u>) at 1/1000 dilution

Lane 1 : Jurkat lysate Lane 2 : K562 lysate

Lane 3: Bone marrow lysate

Lane 4: PBMC lysates

Lysates/proteins at 10 µg per lane.

# **Secondary**

**All lanes :** Standard HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 92 kDa

This data was developed using <u>ab109724</u>, the same antibody clone in a different buffer formulation.



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

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