

## Product datasheet

# Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] $\alpha$ b51134

**KO VALIDATED** Recombinant RabMAB

[24 References](#) [7 Images](#)

### Overview

<b>Product name</b>	Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y]
<b>Description</b>	Rabbit monoclonal [EP1465Y] to Serine/threonine-protein kinase 4/MST-1
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> Flow Cyt (Intra), WB, IHC-P, ICC/IF, IP
<b>Species reactivity</b>	<b>Reacts with:</b> Mouse, Rat, Human
<b>Immunogen</b>	Synthetic peptide within Human Serine/threonine-protein kinase 4/MST-1 aa 1-100 (N terminal). The exact sequence is proprietary.
<b>Positive control</b>	WB: Jurkat, Ramos and HeLa cell lysate and mouse spleen, rat spleen and human urinary bladder tissues. ICC/IF: Raw264.7 cells. IHC-P: Human gastric carcinoma. IP: Jurkat cell lysate. Flow Cyt (intra): HeLa cells.
<b>General notes</b>	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"><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></ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAB<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAB<sup>®</sup> patents</a>.</p>

### Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C. Avoid freeze / thaw cycle.
<b>Storage buffer</b>	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.21% BSA
<b>Purity</b>	Protein A purified

<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EP1465Y
<b>Isotype</b>	IgG

## Applications

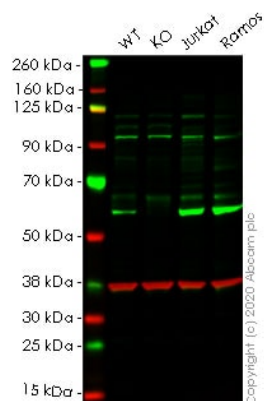
**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab51134 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt (Intra)		1/50.
WB		1/10000. Detects a band of approximately 59 kDa (predicted molecular weight: 56 kDa).
IHC-P		1/50 - 1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. We strongly recommend that customers perform an antigen retrieval step.
ICC/IF		1/100 - 1/250.
IP		1/30 - 1/100.

## Target

<b>Function</b>	Stress-activated, pro-apoptotic kinase which, following caspase-cleavage, enters the nucleus and induces chromatin condensation followed by internucleosomal DNA fragmentation. Key component of the Hippo signaling pathway which plays a pivotal role in organ size control and tumor suppression by restricting proliferation and promoting apoptosis. The core of this pathway is composed of a kinase cascade wherein MST1/MST2, in complex with its regulatory protein SAV1, phosphorylates and activates LATS1/2 in complex with its regulatory protein MOB1, which in turn phosphorylates and inactivates YAP1 oncoprotein and WWTR1/TAZ. Phosphorylation of YAP1 by LATS2 inhibits its translocation into the nucleus to regulate cellular genes important for cell proliferation, cell death, and cell migration. MST1/MST2 are required to repress proliferation of mature hepatocytes, to prevent activation of facultative adult liver stem cells (oval cells), and to inhibit tumor formation (By similarity). Phosphorylates 'Ser-14' of histone H2B (H2BS14ph) during apoptosis. Phosphorylates FOXO3 upon oxidative stress, which results in its nuclear translocation and cell death initiation.
<b>Tissue specificity</b>	Ubiquitously expressed.
<b>Sequence similarities</b>	Belongs to the protein kinase superfamily. STE Ser/Thr protein kinase family. STE20 subfamily. Contains 1 protein kinase domain. Contains 1 SARA domain.
<b>Post-translational modifications</b>	Autophosphorylated on serine and threonine residues.
<b>Cellular localization</b>	Cytoplasm. Nucleus. The caspase-cleaved form cycles between the nucleus and cytoplasm.

## Images



Western blot - Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134)

**All lanes** : Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134) at 1/1000 dilution

**Lane 1** : Wild-type HeLa cell lysate

**Lane 2** : STK4 knockout HeLa cell lysate

**Lane 3** : Jurkat cell lysate

**Lane 4** : Ramos cell lysate

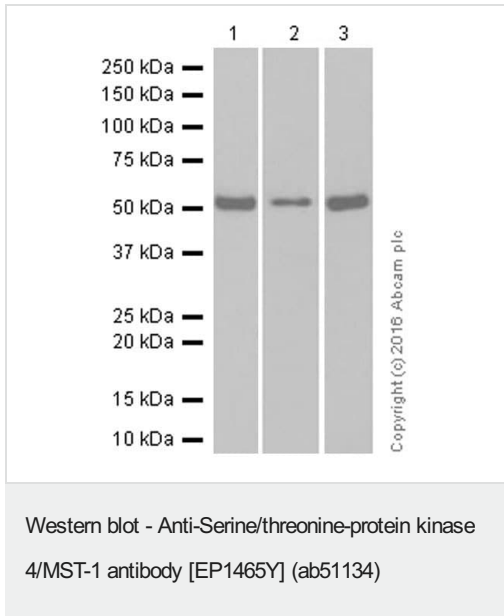
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 56 kDa

**Lanes 1-4:** Merged signal (red and green). Green - ab51134 observed at 52 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab51134 Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] was shown to specifically react with MST-1 in wild-type HeLa cells. Loss of signal was observed when knockout cell line **ab265442** (knockout cell lysate **ab258215**) was used. Wild-type and MST-1 knockout samples were subjected to SDS-PAGE. ab51134 and Anti-GAPDH antibody [6C5] - Loading Control (**ab8245**) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (**ab216776**) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



**All lanes :** Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134) at 1/50000 dilution

**Lane 1 :** Jurkat (human acute T cell leukemia) whole cell lysate

**Lane 2 :** Mouse spleen

**Lane 3 :** Rat spleen

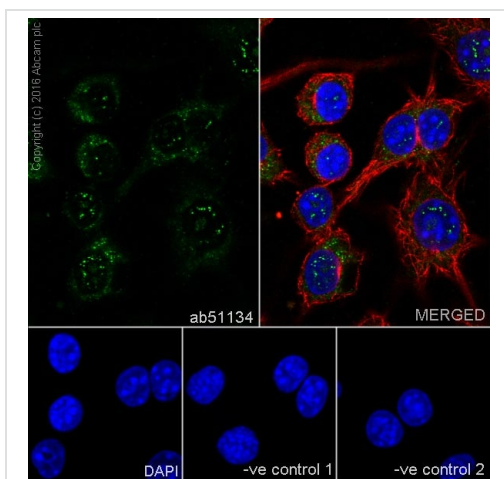
Lysates/proteins at 1/20 dilution per lane.

### Secondary

**All lanes :** Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/2000 dilution

**Predicted band size:** 56 kDa

**Diluting and blocking buffer:** 5% NFD/MTBST

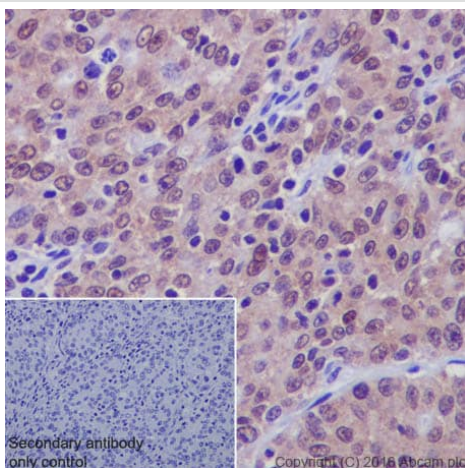


ab51134 staining Serine/threonine-protein kinase 4/MST-1 in Raw264.7 (mouse abelson murine leukemia virus-induced tumor) cells by ICC/IF (Immunocytochemistry/immunofluorescence). Cells were fixed with 100% methanol and permeabilized with 0.1% Triton X-100. Samples were incubated with primary antibody at a dilution of 1/100. A goat anti rabbit IgG (Alexa Fluor® 488) (**ab150077**) was used as the secondary antibody at a concentration of 1/1000.

**ab7291** anti-Tubulin (mouse mAb) (1/1000) and **ab150120** AlexaFluor®594 Goat anti-Mouse secondary (1/1000) were used as counterstains for primary antibody ab51134 and secondary antibody **ab150077** respectively and DAPI was used as a nuclear counterstain.

**Negative control 1:** Rabbit primary antibody and anti-mouse secondary antibody (**ab150120**)

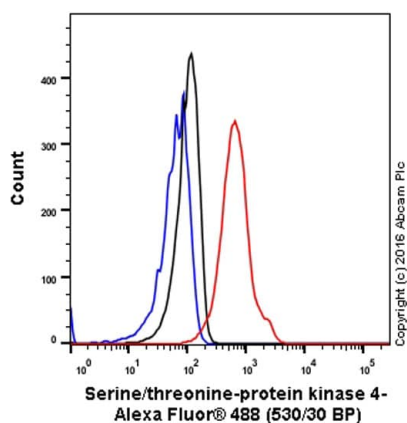
**Negative control 2:** Mouse primary antibody (**ab7291**) and anti-rabbit secondary antibody (**ab150077**)



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134)

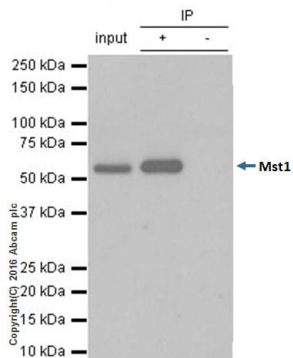
ab51134 staining Serine/threonine-protein kinase 4/MST-1 in human gastric carcinoma tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/50. A goat anti-rabbit IgG H&L (HRP) **ab97051** was used as the secondary antibody at a dilution of 1/500.

**Negative control 1:** PBS in place of primary antibody.



Flow Cytometry (Intracellular) - Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134)

Intracellular Flow Cytometry analysis of HeLa cells labelling Serine/threonine-protein kinase 4/MST-1 with purified ab51134 at a dilution of 1/50 (red). Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. An Alexa Fluor<sup>®</sup> 488-conjugated goat anti-rabbit IgG (1/2000) was used as the secondary antibody. Black - Isotype control, rabbit monoclonal IgG. Blue - Unlabelled control, cells without incubation with primary and secondary antibodies.



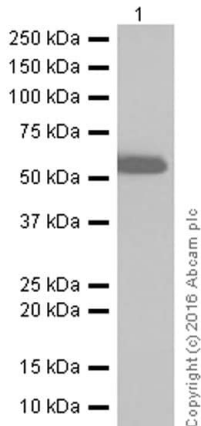
Immunoprecipitation - Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134)

ab51134 immunoprecipitating Serine/threonine-protein kinase 4/MST-1. 10µg of cell lysate was incubated with primary antibody at a dilution of 1/30 and VeriBlot for IP Detection Reagent (HRP) (**ab131366**) at a dilution of 1/1000.

**Lane 1:** Jurkat (human acute T cell leukemia) whole cell lysate (10ug)

**Lane 2:** Jurkat (human acute T cell leukemia) whole cell lysate

**Lane 3:** Rabbit monoclonal IgG (**ab172730**) instead of ab51134 in Jurkat (human acute T cell leukemia) whole cell lysate



Western blot - Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134)

Anti-Serine/threonine-protein kinase 4/MST-1 antibody [EP1465Y] (ab51134) at 1/10000 dilution + HeLa (human cervix adenocarcinoma) whole cell lysate at 20 µg

**Secondary**

Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/2000 dilution

**Predicted band size:** 56 kDa

**Diluting and blocking buffer:** 5% NFDM/TBST

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