

## Product datasheet

# Anti-LATS1/WARTS antibody ab234820

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### Overview

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<b>Product name</b>	Anti-LATS1/WARTS antibody
<b>Description</b>	Rabbit polyclonal to LATS1/WARTS
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> IHC-P, ICC/IF, IP
<b>Species reactivity</b>	<b>Reacts with:</b> Human
<b>Immunogen</b>	Recombinant fragment corresponding to Human LATS1/WARTS aa 1-280. Database link: <a href="#">O95835</a>
<b>Positive control</b>	IHC-P: Human liver cancer, placenta and adrenal gland tissues. ICC/IF: HeLa cells. IP: K562 cell lysate.
<b>General notes</b>	<p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&amp;As</p>

### Properties

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<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 long term. Avoid freeze / thaw cycle.
<b>Storage buffer</b>	pH: 7.40 Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol (glycerin, glycerine), PBS
<b>Purity</b>	Protein G purified
<b>Purification notes</b>	Purity >95%.
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG

## Applications

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**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab234820 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
IHC-P		1/20 - 1/200.
ICC/IF		1/50 - 1/200.
IP		1/200 - 1/2000.

## Target

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**Function** Negative regulator of YAP1 in the Hippo signaling pathway that 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 STK3/MST2 and STK4/MST1, 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 LATS1 inhibits its translocation into the nucleus to regulate cellular genes important for cell proliferation, cell death, and cell migration. Acts as a tumor suppressor which plays a critical role in maintenance of ploidy through its actions in both mitotic progression and the G1 tetraploidy checkpoint. Negatively regulates G2/M transition by down-regulating CDK1 kinase activity. Involved in the control of p53 expression. Affects cytokinesis by regulating actin polymerization through negative modulation of LIMK1. May also play a role in endocrine function.

**Tissue specificity** Expressed in all adult tissues examined except for lung and kidney.

**Sequence similarities** Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. Contains 1 AGC-kinase C-terminal domain. Contains 1 protein kinase domain. Contains 1 UBA domain.

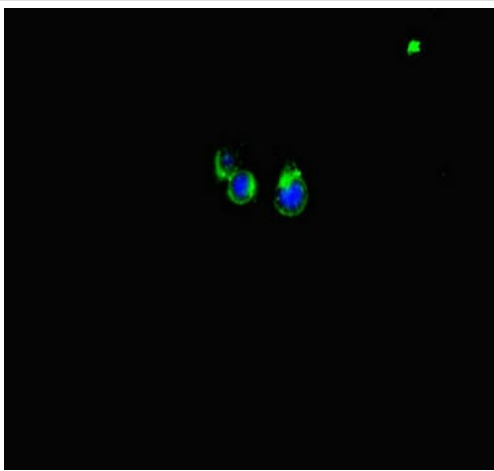
**Post-translational modifications** Autophosphorylated and phosphorylated during M-phase of the cell cycle. Phosphorylated by STK3/MST2 at Ser-909 and Thr-1079, which results in its activation. Phosphorylation at Ser-464 by NUA1 and NUA2 leads to decreased protein level and is required to regulate cellular senescence and cellular ploidy.

**Cellular localization** Cytoplasm, cytoskeleton, microtubule organizing center, centrosome. Localizes to the centrosomes throughout interphase but migrates to the mitotic apparatus, including spindle pole bodies, mitotic spindle, and midbody, during mitosis.

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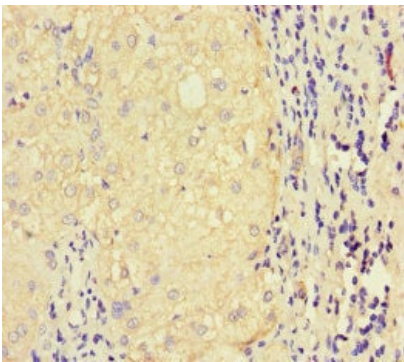
## Images

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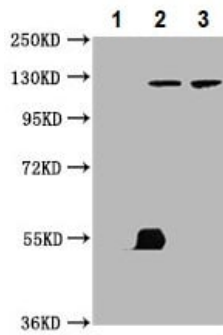
Immunocytochemistry/ Immunofluorescence - Anti-LATS1/WARTS antibody (ab234820)

HeLa (human epithelial cell line from cervix adenocarcinoma) cells stained for LATS1/WARTS (green) using ab234820 at 1/100 dilution in ICC/IF, followed by Alexa Fluor 488<sup>®</sup> conjugated Goat Anti-Rabbit IgG (H+L).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-LATS1/WARTS antibody (ab234820)

Paraffin-embedded human liver cancer tissue stained for LATS1/WARTS using ab234820 at 1/100 dilution in immunohistochemical analysis.



Immunoprecipitation - Anti-LATS1/WARTS antibody (ab234820)

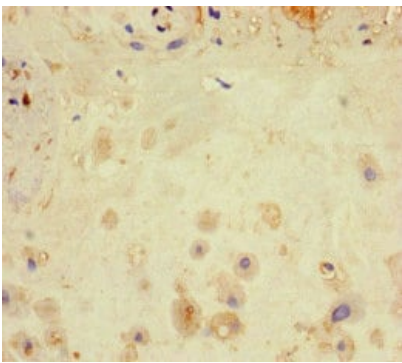
LATS1/WARTS was immunoprecipitated from 500 µg K562 (human chronic myelogenous leukemia cell line from bone marrow) whole cell lysate with 6 µg ab234820.

**Lane 1:** Rabbit control IgG IP (1 µg) in K562 whole cell lysate.

**Lane 2:** ab234820 IP in K562 whole cell lysate.

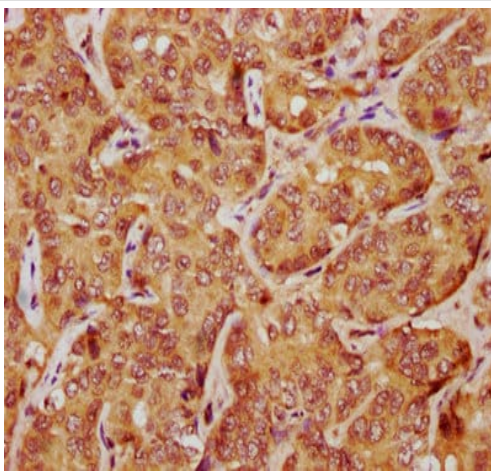
**Lane 3:** K562 whole cell lysate 10 µg (Input).

For western blotting, a HRP-conjugated Protein G antibody was used as the secondary antibody at 1/2000 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-LATS1/WARTS antibody (ab234820)

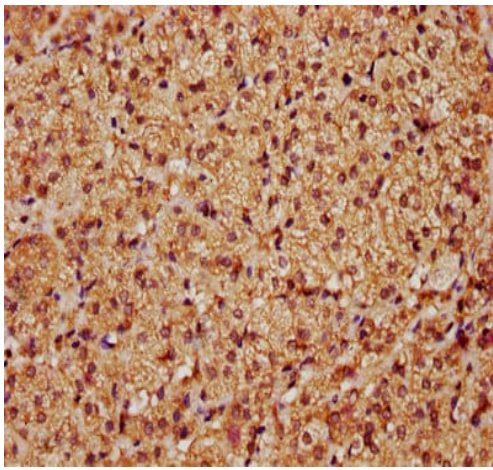
Paraffin-embedded human placenta tissue stained for LATS1/WARTS using ab234820 at 1/100 dilution in immunohistochemical analysis.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-LATS1/WARTS antibody (ab234820)

Paraffin-embedded human liver cancer tissue stained for LATS1/WARTS using ab234820 at 1/100 dilution in immunohistochemical analysis.

Performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-LATS1/WARTS antibody (ab234820)

Paraffin-embedded human adrenal gland tissue stained for LATS1/WARTS using ab234820 at 1/100 dilution in immunohistochemical analysis.

Performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody.

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

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