## LATS1 rabbit monoclonal antibody

Catalog # H00009113-K

Specification

Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human LATS1 peptide using ARM Technology.
Immunogen	A synthetic peptide of human LATS1 is used for rabbit immunization. Customer or Abnova will decide on the preferred peptide sequence.
Host	Rabbit
Library Construction	Non-fusion antibody library from rabbit spleen (ARM Technology).
Expression	Overexpression vector and transfection into 293H cell line.
Reactivity	Human
Purification	Protein A
lsotype	lgG
Quality Control Testing	Antibody reactive against human LATS1 peptide by ELISA and mammalian transfected lysate by We stern Blot.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.
Deliverable	Up to three rabbit IgG clones of 100 ug each will be delivered to customer.
Note	<ol> <li>Customer may provide cell or tissue lysate for antibody screening.</li> <li>Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in cluding F(ab)<sub>2</sub>, lgG, scFv and different Fc and non-Fc conjugates per customer request.</li> </ol>

## Applications

• Western Blot (Transfected lysate)

Protocol Download

• ELISA

Gene Info — LATS1	
Entrez GenelD	<u>9113</u>
GeneBank Accession#	LATS1
Gene Name	LATS1
Gene Alias	WARTS, wts
Gene Description	LATS, large tumor suppressor, homolog 1 (Drosophila)
Omim ID	<u>603473</u>
Gene Ontology	Hyperlink
Gene Summary	The protein encoded by this gene is a putative serine/threonine kinase that localizes to the mitotic apparatus and complexes with cell cycle controller CDC2 kinase in early mitosis. The protein is p hosphorylated in a cell-cycle dependent manner, with late prophase phosphorylation remaining thr ough metaphase. The N-terminal region of the protein binds CDC2 to form a complex showing re duced H1 histone kinase activity, indicating a role as a negative regulator of CDC2/cyclin A. In ad dition, the C-terminal kinase domain binds to its own N-terminal region, suggesting potential nega tive regulation through interference with complex formation via intramolecular binding. Biochemica I and genetic data suggest a role as a tumor suppressor. This is supported by studies in knockout mice showing development of soft-tissue sarcomas, ovarian stromal cell tumors and a high sensiti vity to carcinogenic treatments. [provided by RefSeq

## Disease

- Adenocarcinoma
- Esophageal Neoplasms