# anti-L3MBTL3-RAB-C338



### Applications

Competitior ELISA	Western Blot	SPR	Spiked IP	Immunofluorescence	IP-MS	ChIP	
Pass					Pass		

\*rAb has been tested for the following applications. See below for the experimental details.

## Antibody information

rAb ID: anti-L3MBTL3-RAB-C338

**Description:** recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *L3MBTL3* protein under non-denaturing conditions; specificity and affinity tested.

Binder type: rAb Isotype: IgG1 Species: *Homo sapiens* Produced in: *E. coli* rAb tags: Avi-tag; no tag Specificity: reacts with *Homo sapiens* L3MBTL3 Epitope: binds to folded domain amino acids 229-551 Storage conditions: short term – store at n 4°C (over 6 months), long term - PBS -20°C or -80°C Link: http://recombinant-antibodies.org/binders/anti-L3MBTL3-RAB-C338

### Antigen information

Protein Name: Lethal 3 malignant brain tumor-like protein 3 HGNC Symbol: L3MBTL3 HGNC ID: 23035 Species: Homo sapiens UniProt AC: Q96JM7 UniProt KB: LMBL3\_HUMAN

### Protein Sequence:

MSGLNDIFEAQKIEWHEGSAGGSGKKAWCWASYLEEEKAVAVPAKLFKEHQSFPYNKNGFKVGMKLEGVDPEHQSVYCV LTVAEVCGYRIKLHFDGYSDCYDFWVNADALDIHPVGWCEKTGHKLHPPKGYKEEEFNWQTYLKTCKAQAAPKSLFENQ NITVIPSGFRVGMKLEAVDKKNPSFICVATVTDMVDNRFLVHFDNWDESYDYWCEASSPHIHPVGWCKEHRRTLITPPGYP NVKHFSWDKYLEETNSLPAPARAFKVKPPHGFQKKMKLEVVDKRNPMFIRVATVADTDDHRVKVHFDGWNNCYDYWIDA DSPDIHPVGWCSKTGHPLQPPLSPLELMEGGSGHHHHHH

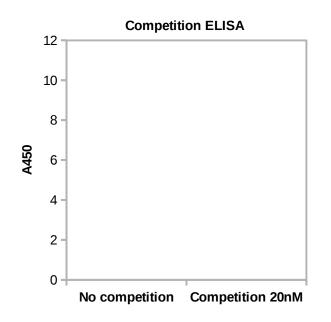
 Tag N-terminus:
 MSGLNDIFEAQKIEWHEGSAGGSG
 Tag C-terminus:
 GGSGHHHHHH

 Vector Type:
 p28BIOH-LIC Vector Link:
 http://www.thesgc.org/sites/default/files/toronto\_vectors/p28BIOH-LIC.pdf

 Protein Sequence Position:
 229-551
 Antigen source:
 E. coli
 Source Lab:
 SGC
 Source Lab ID:
 L3MBTL3-A004

 Description:
 affinity purified recombinant protein

### Validation data



**Single point competition phage ELISA** Plot represents specific binding of a target to the rAb-phage in solution (right bar) in comparison to binding to the target immobilized on the plate surface (left bar). Experimental conditions were calibrated to capture binders with dissociation Constant ( $K_D$ ): 20nM or lower.

Experimental Conditions: Culture supernatants containing rAbphage were diluted five-fold in phosphate-buffered saline, 0.5% (w/v) BSA, 0.1% (v/v) Tween 20 either with or without soluble antigen competitor at 20 nM. After 1 h incubation at room temperature, the mixtures were transferred to neutravidin coated plates preloaded with 50 µL of 20 nM biotinylated antigen and incubated for 15 min. The plates were washed with phosphatebuffered saline, 0.05% (v/v) Tween 20 and incubated for 30 min with horse radish peroxidase/anti-M13 antibody conjugate (1:5000 dilution). The plates were washed, developed with peroxidase 3,3',5,5'-Tetramethyl-benzidine/H<sub>2</sub>O<sub>2</sub> substrate (Thermo Scientific), guenched with 1M H<sub>3</sub>PO<sub>4</sub>, and the absorbance at 450 nm (A450) was determined.

Spiked IP: Status: Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/spiked-IP</u>

#### Immunofluorescence:

Status: Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/immunofluorescence</u>

#### IP-MS – immunoprecipitation for mass spectrometric analysis:

Status: Pass Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/IP-MS</u>

ChIP – chromatin immunoprecipitation: Status: Experimental Conditions: Pending

IP – immunoprecipitation: Status: pass Experimental Conditions: Pending

SP Elisa:

Status: Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/ELISA-IC50-EC50-direct-coating</u>

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