anti-L3MBTL2-RAB-C225



Applications

Competition 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-L3MBTL2-RAB-C225

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *L3MBTL2* 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 L3MBTL2 **Epitope:** binds to folded domain amino acids 170-625

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-L3MBTL2-RAB-C225

Antigen information

Protein Name: Lethal 3 malignant brain tumor-like protein 2

HGNC Symbol: L3MBTL2 HGNC ID: 18594 Species: Homo sapiens

UniProt AC: Q969R5 UniProt KB: LMBL2 HUMAN

Protein Sequence:

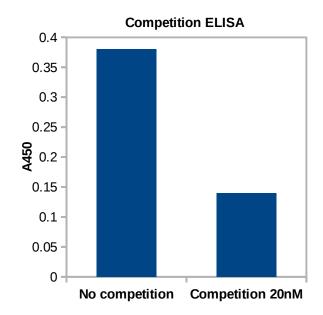
MHHHHHHHHHDLGTENLYFQSMTGQDALVLGFDWGKFLKDHSYKAAPVSCFKHVPLYDQWEDVMKGMKVEVLNSDAV LPSRVYWIASVIQTAGYRVLLRYEGFENDASHDFWCNLGTVDVHPIGWCAINSKILVPPRTIHAKFTDWKGYLMKRLVGSRT LPVDFHIKMVESMKYPFRQGMRLEVVDKSQVSRTRMAVVDTVIGGRLRLLYEDGDSDDDFWCHMWSPLIHPVGWSRRV GHGIKMSERRSDMAHHPTFRKIYCDAVPYLFKKVRAVYTEGGWFEEGMKLEAIDPLNLGNICVATVCKVLLDGYLMICVDG GPSTDGLDWFCYHASSHAIFPATFCQKNDIELTPPKGYEAQTFNWENYLEKTKSKAAPSRLFNMDCPNHGFKVGMKLEAV DLMEPRLICVATVKRVVHRLLSIHFDGWDSEYDQWVDCESPDIYPVGWCELTGYQLQPPVAAEPATPLKAKEATKKKKKQ

Vector Type: pNIC-H102 Vector Link: http://www.thesgc.org/sites/default/files/oxford_vectors/pNIC-H102.pdf

Protein Sequence Position: 170-625 Antigen source: E. coli Source Lab: SGC Source Lab ID: L3MBTL2A-A002

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/H₂O₂ substrate (Thermo Scientific), quenched with 1M H₃PO₄, and the absorbance at 450 nm (A450) was determined.

Spiked IP: Status:

Experimental Conditions: http://recombinant-antibodies.org/protocols/spiked-IP

Immunofluorescence:

Status:

Experimental Conditions: http://recombinant-antibodies.org/protocols/immunofluorescence

IP-MS – immunoprecipitation for mass spectrometric analysis:

Status: Pass

Experimental Conditions: http://recombinant-antibodies.org/protocols/IP-MS

ChIP – chromatin immunoprecipitation:

Status:

Experimental Conditions: Pending

IP - immunoprecipitation:

Status: pass

Experimental Conditions: Pending

SP Elisa:

Status:

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

Visit us at http://recombinant-antibodies.org/

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