anti-PRDM5-RAB-C505



Applications

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

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

Antibody information

rAb ID: anti-PRDM5-RAB-C505

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *PRDM5* 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* PRDM5 **Epitope:** binds to folded domain amino acids 2-226 **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-PRDM5-RAB-C505

Antigen information

Protein Name: PR domain-containing protein 5 HGNC Symbol: PRDM5 HGNC ID: 9349 Species: Homo sapiens

UniProt AC: Q9NQX1 UniProt KB: PRDM5_HUMAN

Protein Sequence:

MHHHHHHHHHHDLGTENLYFQSMLGMYVPDRFSLKSSRVQDGMGLYTARRVRKGEKFGPFAGEKRMPEDLDENMDYRL MWEVRGSKGEVLYILDATNPRHSNWLRFVHEAPSQEQKNLAAIQEGENIFYLAVEDIETDTELLIGYLDSDMEAEEEEQQIM TVIKEGEVENSRRQSTAGRKDRLGCKEDYACPQCESSFTSEDILAEHLQTLHQKPTEEKEFKCKNCGKKFPVKQALQRHV LQCTAKSSSSKGGYGLNDIFEAQKIEWHE

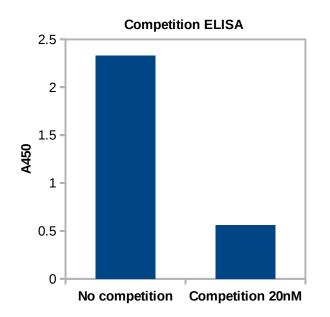
 Tag N-terminus:
 MHHHHHHHHHHHHHHDLGTENLYFQSM
 Tag C-terminus:
 SSKGGYGLNDIFEAQKIEWHE

 Vector Type:
 pNIC-Bio2 Vector Link:
 http://www.thesgc.org/sites/default/files/oxford_vectors/pNIC-Bio2.pdf

 Protein Sequence Position:
 2-226 Antigen source:
 E. coli Source Lab:
 SGC Source Lab ID:
 PRDM5-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), guenched with 1M H₃PO₄, 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: 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>

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

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