# anti-SMYD3-RAB-C233



### 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-SMYD3-RAB-C233

**Description:** recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *SMYD3* protein under non-denaturing conditions; specificity and affinity tested.

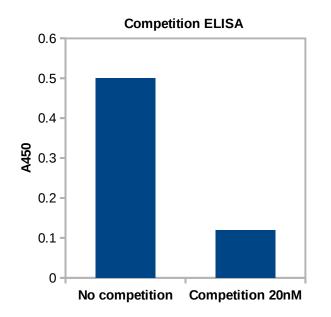
Binder type: rAb Isotype: IgG1 Species: *Homo sapiens* Produced in: *E. coli* rAb tags: Avi-tag; no tag Specificity: reacts with *Homo sapiens* SMYD3 Epitope: binds to folded domain amino acids 2-428 Storage conditions: short term – store at n 4°C (over 6 months), long term - PBS -20°C or -80°C Link: http://recombinant-antibodies.org/binders/anti-SMYD3-RAB-C233

### Antigen information

Protein Name: SET And MYND Domain-Containing Protein 3 HGNC Symbol: SMYD3 HGNC ID: 15513 Species: Homo sapiens UniProt AC: Q9H7B4 UniProt KB: SMYD3\_HUMAN Protein Sequence:

MHHHHHHHHHHHDLGTENLYFQSMEPLKVEKFATAKRGNGLRAVTPLRPGELLFRSDPLAYTVCKGSRGVVCDRCLLGKEK LMRCSQCRVAKYCSAKCQKKAWPDHKRECKCLKSCKPRYPPDSVRLLGRVVFKLMDGAPSESEKLYSFYDLESNINKLTE DKKEGLRQLVMTFQHFMREEIQDASQLPPAFDLFEAFAKVICNSFTICNAEMQEVGVGLYPSISLLNHSCDPNCSIVFNGPH LLLRAVRDIEVGEELTICYLDMLMTSEERRKQLRDQYCFECDCFRCQTQDKDADMLTGDEQVWKEVQESLKKIEELKAHW KWEQVLAMCQAIISSNSERLPDINIYQLKVLDCAMDACINLGLLEEALFYGTRTMEPYRIFFPGSHPVRGVQVMKVGKLQLH QGMFPQAMKNLRLAFDIMRVTHGREHSLIEDLILLLEECDANIRASSSKGGYGLNDIFEAQKIEWHE **Tag N-terminus:** MHHHHHHHHDLGTENLYFQSM **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:** 2-428 **Antigen source:** *E. coli* **Source Lab:** SGC **Source Lab ID:** SMYD3-A003 **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  $1M H_3PO_4$ , 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: Experimental Conditions: Pending

SP Elisa:

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

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Contact: Recombinant Antibody Network admin@recombinant-antibodies.org

#### The University of Chicago

Knapp Center for Biomedical Discovery Rm. 3240G 900 E. 57th St., Chicago, IL 60637 Phone: +1 (773) 834-2776

University of California, San Franciso Byers Hall Rm. 503 1700 4th St., San Francisco, CA 94158 Phone: +1 (530) 341-2371

#### University of Toronto

Best Institute Rm. 117 112 College Avenue, Toronto, Ontario M5G 1L6 Phone: +1 (416) 978-1594