anti-ZSCAN12-RAB-S103



Applications

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

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

Antibody information

rAb ID: anti-ZSCAN12-RAB-S103

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *ZSCAN12* 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; Flag-tag

Specificity: reacts with *Homo sapiens* ZSCAN12 **Epitope:** binds to folded domain amino acids 37-132

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-ZSCAN12-RAB-S103

Antigen information

Protein Name: Zinc finger and SCAN domain-containing protein 12 **HGNC Symbol:** ZSCAN12 **HGNC ID:** 13172 **Species:** *Homo sapiens*

UniProt AC: O43309 UniProt KB: ZSC12 HUMAN

Protein Sequence:

MSGLNDIFEAQKIEWHEHHHHHHENLYFQSHMNNTHSREVFRQYFRQFCYQETSGPREALSRLRELCHQWLRPETHTKE QILELLVLEQFLTILPEELQAWVQEQHPESGEEVVTVLEDLERELDEPG

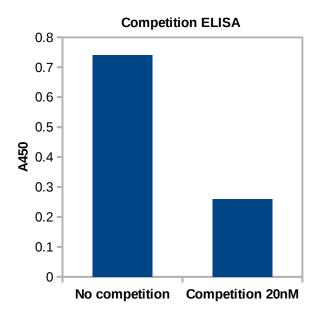
Tag N-terminus: MSGLNDIFEAQKIEWHEHHHHHHHENLYFQSHM Tag C-terminus:

Vector Type: pET15Avi6HT_NESG Vector Link: http://beta.labgeni.us/registries/DNASU/pET15Avi6HT_NESG/

Protein Sequence Position: 37-132 Antigen source: E. coli Source Lab: Rutgers Source Lab ID: HR8495A.005

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: Pass

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

Immunofluorescence:

Status: Pass

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

IP-MS – immunoprecipitation for mass spectrometric analysis:

Status:

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

ChIP – chromatin immunoprecipitation:

Status:

Experimental Conditions: Pending

IP - immunoprecipitation:

Status:

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/

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