anti-SUV39H1-RAB-C489



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-SUV39H1-RAB-C489

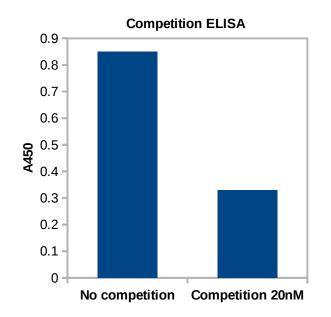
Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *SUV39H1* protein under non-denaturing conditions; specificity and affinity tested.

Binder type: rAb **Isotype:** IgG1 **Species:** *Homo sapiens* **Produced in:** *E. coli* **rAb tags:** Avi-tag; no tag **Specificity:** reacts with *Homo sapiens* SUV39H1 **Epitope:** binds to folded domain amino acids 44-96 **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-SUV39H1-RAB-C489

Antigen information

Protein Name: Suppressor of variegation 3-9 homolog 1 HGNC Symbol: SUV39H1 HGNC ID: 11479 Species: Homo sapiens UniProt AC: O43463 UniProt KB: SUV91_HUMAN Protein Sequence: MSGLNDIFEAQKIEWHEGSAGGSGDFEVEYLCDYKKIREQEYYLVKWRGYPDSESTWEPRQNLKCVRILKQFHKDLEREL LRRGGSGHHHHHH 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: 44-96 Antigen source: *E. coli* Source Lab: SGC Source Lab ID: SUV39H1-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 $1M H_3PO_4$, 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: 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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