anti-HDAC7-RAB-C321



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-HDAC7-RAB-C321

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *HDAC7* 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* HDAC7 **Epitope:** binds to folded domain amino acids 483-903

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-HDAC7-RAB-C321

Antigen information

Protein Name: Histone deacetylase 7

HGNC Symbol: HDAC7 HGNC ID: 14067 Species: Homo sapiens

UniProt AC: Q8WUI4 UniProt KB: HDAC7 HUMAN

Protein Sequence:

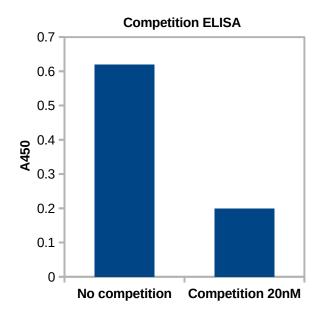
MSGLNDIFEAQKIEWHEGSAGGSGRAQSSPAAPASLSAPEPASQARVLSSSETPARTLPFTTGLIYDSVMLKHQCSCGDN SRHPEHAGRIQSIWSRLQERGLRSQCECLRGRKASLEELQSVHSERHVLLYGTNPLSRLKLDNGKLAGLLAQRMFVMLPC GGVGVDTDTIWNELHSSNAARWAAGSVTDLAFKVASRELKNGFAVVRPPGHHADHSTAMGFCFFNSVAIACRQLQQQSK ASKILIVDWDVHHGNGTQQTFYQDPSVLYISLHRHDDGNFFPGSGAVDEVGAGSGEGFNVNVAWAGGLDPPMGDPEYLA AFRIVVMPIAREFSPDLVLVSAGFDAAEGHPAPLGGYHVSAKCFGYMTQQLMNLAGGAVVLALEGGHDLTAICDASEACVA ALLGNRVDPLSEEGWKQKPNLNAIRSLEAVIRVHSKYWGCMQRLASGGSGHHHHHH

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: 483-903 Antigen source: *E. coli* Source Lab: SGC Source Lab ID: HDAC7-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 3,3',5,5'-Tetramethyl-benzidine/H₂O₂ peroxidase substrate (Thermo Scientific), quenched with 1M H₃PO₄, and the absorbance at 450 nm (A450) was determined.

Spiked IP: Status:

Experimental Conditions: Pending

Immunofluorescence:

Status:

Experimental Conditions: https://recombinant-antibodies.org/protocols/#uagb-tabs tab1

IP-MS – immunoprecipitation for mass spectrometric analysis:

Status: Pass

Experimental Conditions: https://recombinant-antibodies.org/protocols/#uagb-tabs tab1

ChIP – chromatin immunoprecipitation:

Status:

Experimental Conditions: Pending

IP - immunoprecipitation:

Status:

Experimental Conditions: Pending

SP Elisa: Status:

Experimental Conditions: https://recombinant-antibodies.org/protocols/#uagb-tabs tab1

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