anti-C14ORF169-RAB-C350



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-C14ORF169-RAB-C350

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *C14orf169* 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* C14orf169 Epitope: binds to folded domain amino acids 183-641 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-C14ORF169-RAB-C350

Antigen information

Protein Name: chromosome 14 open reading frame 169; NO66 HGNC Symbol: C14orf169 HGNC ID: 20968 Species: *Homo sapiens* UniProt AC: Q9H6W3 UniProt KB: NO66_HUMAN

Protein Sequence:

MSGLNDIFEAQKIEWHEGSAGGSGSPLRRVLAELNRIPSSRRRAARLFEWLIAPMPPDHFYRRLWEREAVLVRRQDHTYY QGLFSTADLDSMLRNEEVQFGQHLDAARYINGRRETLNPPGRALPAAAWSLYQAGCSLRLLCPQAFSTTVWQFLAVLQEQ FGSMAGSNVYLTPPNSQGFAPHYDDIEAFVLQLEGRKLWRVYRPRVPTEELALTSSPNFSQDDLGEPVLQTVLEPGDLLYF PRGFIHQAECQDGVHSLHLTLSTYQRNTWGDFLEAILPLAVQAAMEENVEFRRGLPRDFMDYMGAQHSDSKDPRRTAFM EKVRVLVARLGHFAPVDAVADQRAKDFIHDSLPPVLTDRERALSVYGLPIRWEAGEPVNVGAQLTTETEVHMLQDGIARLVG EGGHLFLYYTVENSRVYHLEEPKCLEIYPQQADAMELLLGSYPEFVRVGDLPCDSVEDQLSLATTLYDKGLLLTKMPLALNG GSGHHHHHH

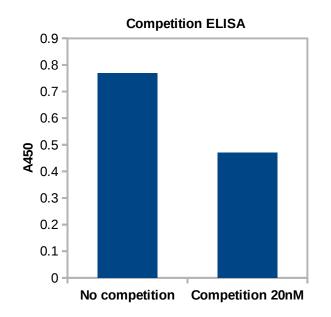
 Tag N-terminus: MSGLNDIFEAQKIEWHEGSAGGSG
 Tag C-terminus: GGSGHHHHHH

 Vector Type:
 p28BIOH-LIC Vector Link: http://www.thesgc.org/sites/default/files/toronto_vectors/p28BIOH-LIC.pdf

 Protein Sequence Position:
 183-641 Antigen source: E. coli Source Lab: SGC Source Lab ID: FLJ21802A-A001

 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: 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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