anti-IRF2-RAB-S206



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-IRF2-RAB-S206

Description: recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *IRF2* protein under non-denaturing conditions; specificity and affinity tested.

Binder type: rAb Isotype: IgG1 Species: Homo sapiens Produced in: E. coli rAb tags: no tag; Flag-tag

Specificity: reacts with *Homo sapiens* IRF2 **Epitope:** binds to folded domain amino acids 32-144 **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-IRF2-RAB-S206

Antigen information

Protein Name: Interferon regulatory factor 2

HGNC Symbol: IRF2 HGNC ID: 6117 Species: Homo sapiens

UniProt AC: P14316 UniProt KB: IRF2 HUMAN

Protein Sequence:

MSGLNDIFEAQKIEWHEHHHHHHENLYFQSHMPVERMRMRPWLEEQINSNTIPGLKWLNKEKKIFQIPWMHAARHGWDV

EKDAPLFRNWAIHTGKHQPGVDKPDPKTWKANFRCAMNSLPDIEEVKDKSIKKGNNAFRVYRMLP

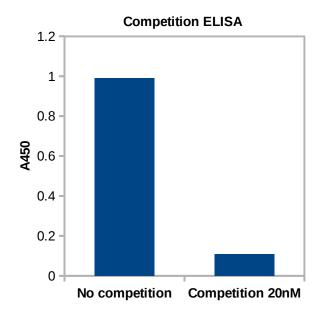
Tag N-terminus: MSGLNDIFEAQKIEWHEHHHHHHENLYFQSHM Tag C-terminus:

Vector Type: RH2.2 Vector Link: http://dnasu.org/DNASU/GetVectorDetail.do?vectorid=612

Protein Sequence Position: 32-144 Antigen source: E. coli Source Lab: RAN Source Lab ID: BL64-Irf2

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:

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: Pass

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

Visit us at http://recombinant-antibodies.org/

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