# anti-WDR12-RAB-C528



### 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-WDR12-RAB-C528

**Description:** recombinant Fab fragment obtained by recombinant antibody (rAb) phage display recognizing *WDR12* 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* WDR12 **Epitope:** binds to folded domain amino acids 1-423 **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-WDR12-RAB-C528

### **Antigen information**

Protein Name: WD repeat domain protein 12 HGNC Symbol: WDR12 HGNC ID: 14098 Species: Homo sapiens UniProt AC: Q9GZL7 UniProt KB: WDR12\_HUMAN

#### **Protein Sequence:**

MSGLNDIFEAQKIEWHEGSAGGSGMAQLQTRFYTDNKKYAVDDVPFSIPAASEIADLSNIINKLLKDKNEFHKHVEFDFLIKG QFLRMPLDKHMEMENISSEEVVEIEYVEKYTAPQPEQCMFHDDWISSIKGAEEWILTGSYDKTSRIWSLEGKSIMTIVGHTD VVKDVAWVKKDSLSCLLLSASMDQTILLWEWNVERNKVKALHCCRGHAGSVDSIAVDGSGTKFCSGSWDKMLKIWSTVP TDEEDEMEESTNRPRKKQKTEQLGLTRTPIVTLSGHMEAVSSVLWSDAEEICSASWDHTIRVWDVESGSLKSTLTGNKVFN CISYSPLCKRLASGSTDRHIRLWDPRTKDGSLVSLSLTSHTGWVTSVKWSPTHEQQLISGSLDNIVKLWDTRSCKAPLYDLA AHEDKVLSVDWTDTGLLLSGGADNKLYSYRYSPTTSHVGAGGSGHHHHHH

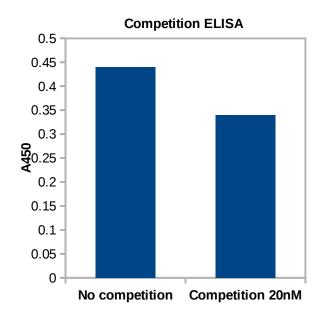
 Tag N-terminus:
 MSGLNDIFEAQKIEWHEGSAGGSG
 Tag C-terminus:
 GGSGHHHHHH

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

 Protein Sequence Position:
 1-423
 Antigen source:
 E. coli
 Source Lab:
 SGC
 Source Lab ID:
 WDR12-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/H<sub>2</sub>O<sub>2</sub> substrate (Thermo Scientific), guenched with 1M H<sub>3</sub>PO<sub>4</sub>, and the absorbance at 450 nm (A450) was determined.

Spiked IP: Status: Pass Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/spiked-IP</u>

#### Immunofluorescence:

Status: Pass Experimental Conditions: <u>http://recombinant-antibodies.org/protocols/immunofluorescence</u>

# 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: <u>http://recombinant-antibodies.org/protocols/ELISA-IC50-EC50-direct-coating</u>

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