# Rabbit IgG isotype control

Catalog No.: AC042 9 Publications



## **Basic Information**

### **Observed MW**

**Calculated MW** 

### Category

Loading control antibody

### **Applications**

WB,IHC-P,IF/ICC,FC,ChIP,ELISA

### **Cross-Reactivity**

### CloneNo number

ARC5105-03

# **Background**

The protein encoded by this gene is a transcriptional regulator and tumor suppressor, serving as an activator of genes involved in both innate and acquired immune responses.

# **Recommended Dilutions**

**WB** 1:5000 - 1:10000

**IHC-P** 1:50 - 1:200

**IF/ICC** 1:50 - 1:200

**ChIP** 3μg antibody for

10μg-15μg of Chromatin

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

### **Contact**

<b>a</b>	400-999-6126
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•	www.abclonal.com.cn

# **Immunogen Information**

Gene ID Swiss Prot

### **Immunogen**

This information is considered to be commercially sensitive.

### **Synonyms**

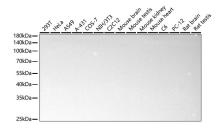
### **Product Information**

SourceIsotypePurificationRabbitIgGProtein A/G purification

### **Storage**

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% sodium azid, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of various lysates using Rabbit IgG isotype control (AC042) at 1:10000 dilution incubated at room temperature for 1.5 hours.

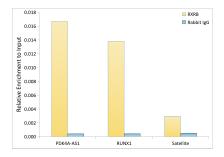
Secondary antibody: () at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

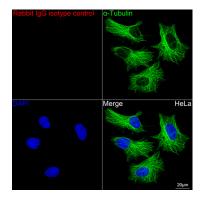
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

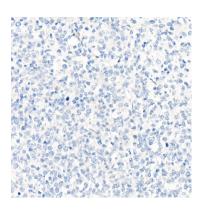
Exposure time: 90s.

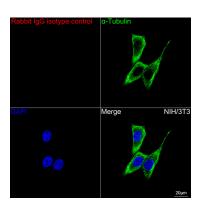


Chromatin immunoprecipitation was performed with 15  $\mu$ g of cross-linked chromatin from HeLa cells transfected with a RXRB expression vector containing a single C-terminal flag-Tag, using 3  $\mu$ g of Rabbit IgG isotype control (AC042) and RXRB Rabbit mAb (A25648). The enrichment of immunoprecipitated DNA at different genomic loci was examined by quantitative PCR. The histogram compares the ratio of the immunoprecipitated DNA to the input at given loci.

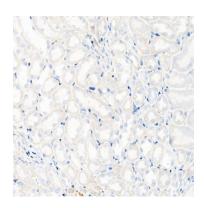


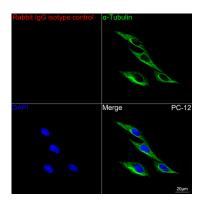
Confocal imaging of HeLa cells using Rabbit IgG isotype control (AC042, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



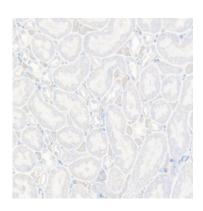


Confocal imaging of NIH/3T3 cells using Rabbit IgG isotype control (AC042, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

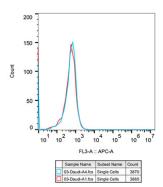




Confocal imaging of PC-12 cells using Rabbit IgG isotype control (AC042, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with  $\alpha$ -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

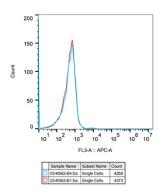


Immunohistochemistry analysis of paraffinembedded Human tonsil using Rabbit IgG isotype control (AC042) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Flow cytometry: Daudi cells were stained with Rabbit IgG isotype control(AC042, 10 µg/mL, blue line) followed by goat anti-Rabbit pAb APC(1:600 dilution) staining. Nonfluorescently stained Daudi cells was used as blank control (red line).

Immunohistochemistry analysis of paraffinembedded Mouse kidney using Rabbit IgG isotype control (AC042) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Flow cytometry: K562 cells were stained with Rabbit IgG isotype control(AC042, 10 µg/mL, blue line) followed by goat anti-Rabbit pAb APC(1:600 dilution) staining. Nonfluorescently stained K562 cells was used as blank control (red line).

Immunohistochemistry analysis of paraffinembedded Rat kidney using Rabbit IgG isotype control (AC042) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.