# Rabbit Anti-Mouse IgG2b (Fc) mAb

Catalog No.: AS128



### **Basic Information**

#### **Observed MW**

55kDa

### **Calculated MW**

60kDa

### Category

Secondary antibody

### **Applications**

WB,FC,ELISA

### **Cross-Reactivity**

Mouse

#### CloneNo number

ARC67783

# **Background**

Antigen binding activity, upstream or within phagocytosis, positive regulation of immune response and positive regulation of phagocytosis. Located in the extracellular space. Partial immunoglobulin complex, circulation. It is derived from several human genes, including IGHG1(immunoglobulin heavy constant  $\gamma1$  (G1m marker)); igg2(immunoglobulin heavy constant  $\gamma2$  (G2m marker)); And IGHG3(immunoglobulin heavy constant  $\gamma3$  (G3m marker)).

# **Recommended Dilutions**

**WB** 1:1000 - 1:5000

FC 1:500 - 1:1000

**ELISA** Recommended starting concentration is 1 µg/mL.

Please optimize the concentration based on your specific assay requirements.

# **Immunogen Information**

**Gene ID Swiss Prot** 16016 P01867-2

### **Immunogen**

A synthetic peptide corresponding to a sequence within amino acids 14-113 of mouse IgG2b (P01867-2) .

### **Synonyms**

IgG2b; Igh-3; gamma2b;

# **Contact**

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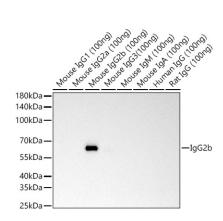
### **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

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

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



 $We stern\ blot\ analysis\ of\ lysates\ from\ Mouse\ lgG2b\ using\ Rabbit\ Anti-Mouse\ lgG2b\ (Fc)\ mAb\ (AS128) at$ 

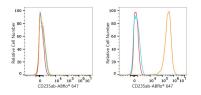
1:1000 dilution incubated overnight at 4°C. Lysates/proteins: 100ng  $\mu g$  per lane.

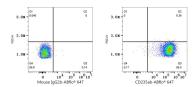
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

 $Negative\ control\ (NC):\ Mouse\ IgG1, Mouse\ IgG2a, Mouse\ IgG3, Mouse\ IgM, Mouse\ IgA, Human\ IgG, Rat\ IgGabber (NC) and Mouse\ IgG1, Mouse\ IgG2a, Mouse\ IgG3, Mouse\ IgM, Mouse\ IgG4, Human\ IgG4, Rat\ IgG4, Mouse\ IgG$ 

Exposure time: 1s.





Flow cytometry:  $1\times10^6$  Jurkat cells (negative control,left) and HEL cells (right) were surface-stained with Mouse anti-Human CD235ab mAb (lgG2b,  $\kappa$ ,orange line) or Mouse lgG2b isotype control (2  $\mu$ g/mL,blue line), followed by Rabbit Anti-Mouse lgG2b (Fc) mAb (AS128,2  $\mu$ g/mL) and Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry:  $1\times10^6$  HEL cells were surface-stained with Mouse 1gG2b isotype control (2  $\mu g/mL$ ,left) or Mouse anti-Human CD235ab mAb (1gG2b,  $\kappa$ ,right). Rabbit Anti-Mouse 1gG2b (Fc) mAb (AS128,2  $\mu g/mL$ ) and Alexa Fluor® 647 conjugated goat anti-rabbit pAb was used as a secondary antibody.