# CD38 Rabbit mAb

Catalog No.: A25398 Recombinant



## **Basic Information**

#### **Observed MW**

45kDa

### **Calculated MW**

14kDa/34kDa

### Category

Primary antibody

### **Applications**

WB,IHC-P,FC,ELISA

### **Cross-Reactivity**

Human

#### CloneNo number

ARC66212

## **Background**

The protein encoded by this gene is a non-lineage-restricted, type II transmembrane glycoprotein that synthesizes and hydrolyzes cyclic adenosine 5'-diphosphate-ribose, an intracellular calcium ion mobilizing messenger. The release of soluble protein and the ability of membrane-bound protein to become internalized indicate both extracellular and intracellular functions for the protein. This protein has an N-terminal cytoplasmic tail, a single membrane-spanning domain, and a C-terminal extracellular region with four N-glycosylation sites. Crystal structure analysis demonstrates that the functional molecule is a dimer, with the central portion containing the catalytic site. It is used as a prognostic marker for patients with chronic lymphocytic leukemia. Alternative splicing results in multiple transcript variants.

## **Recommended Dilutions**

**WB** 1:3000 - 1:18000

IHC-P 1:500 - 1:2000

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**952

Swiss Prot
P28907

### **Immunogen**

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

### **Synonyms**

ADPRC1; cADPR1; ADPRC 1

## **Contact**

<u>a</u>		400-999-6126
$\bowtie$		cn.market@abclonal.com.cn
$\odot$	П	www.abclonal.com.cn

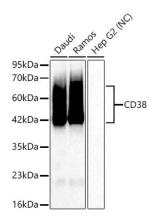
### **Product Information**

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.

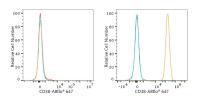


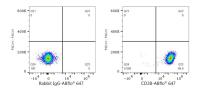
Western blot analysis of various lysates using CD38 Rabbit mAb (A25398) at 1:3000 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

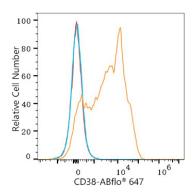
Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Negative control (NC): Hep G2.

Exposure time: 30s.



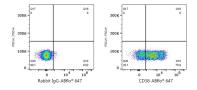


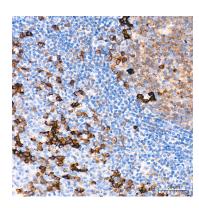


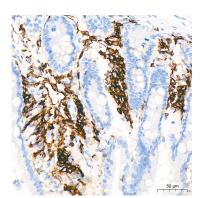
Flow cytometry: 1X10^6 Hep G2 cells (negative control,left) and Daudi cells (right) were surface-stained with CD38 Rabbit mAb (A25398,2 µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,5 µl/Test,blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry:  $1X10^6$  Daudi cells were surface-stained with ABflo® 647 Rabbit IgG isotype control (A22070,5  $\mu$ I/Test,left) or CD38 Rabbit mAb (A25398,2  $\mu$ g/mL,right).

Flow cytometry: 1X10^6 Human PBMC were surface-stained with CD38 Rabbit mAb (A25398,2 µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,5 µl/Test,blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained Human PBMC were used as blank control (red line).







Flow cytometry: 1X10^6 Human PBMC were surface-stained with ABflo® 647 Rabbit IgG isotype control (A22070,5 µl/Test,left) or CD38 Rabbit mAb (A25398,2 µg/mL,right).

Immunohistochemistry analysis of paraffinembedded Human tonsil tissue using CD38 Rabbit mAb (A25398) at a dilution of 1:800 (40x lens). High pressure antigen retrieval was performed with 0.01 M Tris-EDTA buffer (pH 9.0) prior to IHC staining.

Immunohistochemistry analysis of paraffinembedded Human colon tissue using CD38 Rabbit mAb (A25398) at a dilution of 1:800 (40x lens). High pressure antigen retrieval was performed with 0.01 M Tris-EDTA buffer (pH 9.0) prior to IHC staining.