# CD38 Rabbit mAb

Catalog No.: A25398 Recombinant



## **Basic Information**

### **Observed MW**

45kDa

### **Calculated MW**

14kDa/34kDa

### Category

Primary antibody

### **Applications**

ELISA,WB,IHC-P,FC

#### **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	1:1000 - 1:5000	
IHC-P	1:500 - 1:1000	
FC	1:500 - 1:1000	

## **Immunogen Information**

Gene ID	Swiss Prot
952	P28907

### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 43-300 of human CD38 (NP\_001766.2).

## **Synonyms**

ADPRC1; cADPR1; ADPRC 1

## Contact

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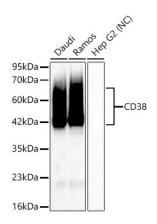
### **Product Information**

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

#### Storage

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

Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.



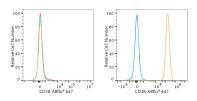
Western blot analysis of various lysates using CD38 Rabbit mAb (A25398) at 1:3000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

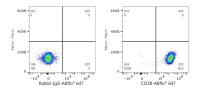
Lysates/proteins: 25 µg per lane.

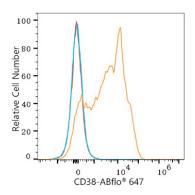
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020). Negative control (NC): Hep G2.

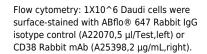
Exposure time: 30s.



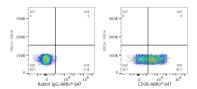


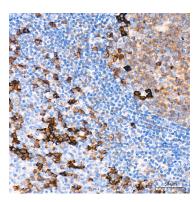


Flow cytometry: 1X10^6 Hep G2 cells (negative control,left) and Daudi cells (right) were surface-stained with CD38 Rabbit mAb (A25398,2 µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,5 µl/Test,blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently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



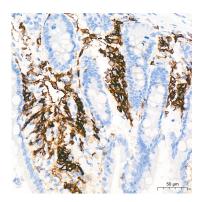
Flow cytometry: 1X10^6 Human PBMC were surface-stained with CD38 Rabbit mAb (A25398,2 µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,5 µl/Test,blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained Human PBMC were used as blank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





Immunohistochemistry analysis of CD38 in paraffin-embedded 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 CD38 in paraffin-embedded 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.

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