ST6GAL1/CD75 Rabbit mAb

Catalog No.: A24645 Recombinant



Basic Information

Observed MW

50-70kDa

Calculated MW

20kDa/47kDa

Category

SMab Recombinant Monoclonal Antibody

Applications

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

Cross-Reactivity

Human

CloneNo number

ARC63359

Background

This gene encodes a member of glycosyltransferase family 29. The encoded protein is a type II membrane protein that catalyzes the transfer of sialic acid from CMP-sialic acid to galactose-containing substrates. The protein, which is normally found in the Golgi but can be proteolytically processed to a soluble form, is involved in the generation of the cell-surface carbohydrate determinants and differentiation antigens HB-6, CD75, and CD76. This gene has been incorrectly referred to as CD75. Alternative splicing results in multiple transcript variants.

Recommended Dilutions

WB	1:2000 - 1:8000
IHC-P	1:500 - 1:1000
IF/ICC	1:50 - 1:200
FC	1.500 - 1.1000

Immunogen Information

Gene ID	Swiss Prot
6480	P15907

Immunogen

Recombinant fusion protein containing a sequence corresponding to a mino acids 44-406 of human ST6GAL1/CD75 (NP_001340845.1).

Synonyms

ST6N; CDw75; SIAT1; ST6GalI; ST6GAL1/CD75

Contact

vww.abclonal.com

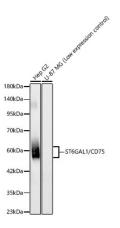
Product Information

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 lysates from Hep G2 cells, using ST6GAL1/CD75 Rabbit mAb (A24645) at 1:7000 dilution.

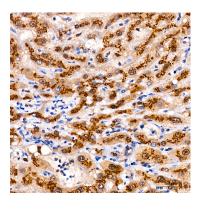
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

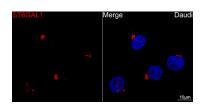
Lysates/proteins: 25ug per lane.

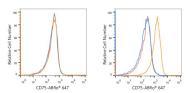
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 90s.



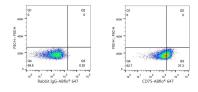




Immunohistochemistry analysis of ST6GAL1/CD75 in paraffinembedded Human liver using ST6GAL1/CD75 Rabbit mAb (A24645) at dilution of 1:800 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.

Confocal imaging of Daudi cells using ST6GAL1/CD75 Rabbit mAb (A24645,dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007,dilution 1:500)(Red).DAPI was used for nuclear staining (Blue). Objective: 100x.

Flow cytometry: 1X10^6 HeLa cells (negative control,left) and Daudi cells (right) were surface-stained with ST6GAL1/CD75 Rabbit mAb (A24645,2 µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,2 µg/mL,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 (A22070,2 µg/mL,left) or ST6GAL1/CD75 Rabbit mAb (A24645,2 µg/mL,right).