

APC/Cyanine7 Rabbit anti-Human/Monkey IgD mAb

Catalog No.: A27285

Basic Information

Observed MW

Calculated MW

47kDa

Category

Primary antibody

Applications

FC

Cross-Reactivity

Human, Rhesus

CloneNo number

ARC60051

Conjugate

APC-Cy7. Ex:651nm. Em:779nm.

Background

Predicted to enable antigen binding activity and immunoglobulin receptor binding activity. Involved in positive regulation of interleukin-1 production. Located in blood microparticle and extracellular exosome.

Recommended Dilutions

FC 5 μ l per 10^6 cells in
100 μ l volume

Immunogen Information

Gene ID

3495

Swiss Prot

P01880

Immunogen

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

Synonyms

Immunoglobulin heavy constant delta

Contact

☎ | 400-999-6126

✉ | cn.market@abclonal.com.cn

🌐 | www.abclonal.com.cn

Product Information

Source

Rabbit

Isotype

IgG

Purification

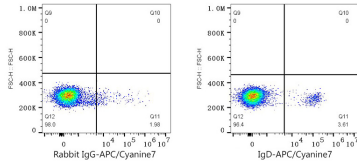
Affinity purification

Storage

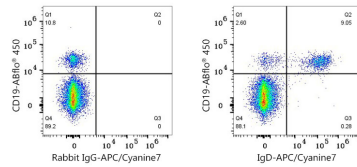
Store at 2-8°C. Avoid freeze.

Buffer: PBS with 0.09% Sodium azide, 0.2% BSA, pH7.3.

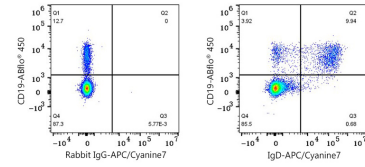
Validation Data



Flow cytometry: 1×10^6 Human PBMC were surface-stained with APC/Cyanine7 Rabbit IgG isotype control ($5 \mu\text{l}/\text{Test}$, left) or APC/Cyanine7 Rabbit anti-Human IgD mAb ($5 \mu\text{l}/\text{Test}$, right).



Flow cytometry: 1×10^6 Human PBMC were surface-stained with ABflo® 450 Rabbit anti-Human/Monkey CD19 mAb ($5 \mu\text{l}/\text{Test}$) and APC/Cyanine7 Rabbit IgG isotype control ($5 \mu\text{l}/\text{Test}$, left) or APC/Cyanine7 Rabbit anti-Human/Monkey IgD mAb ($5 \mu\text{l}/\text{Test}$, right). Cells in the lymphocyte gate were used for analysis.



Flow cytometry: 1×10^6 Rhesus PBMC were surface-stained with ABflo® 488 Mouse anti-Human CD11b mAb ($5 \mu\text{l}/\text{Test}$), ABflo® 450 Rabbit anti-Human/Monkey CD19 mAb ($5 \mu\text{l}/\text{Test}$) and APC/Cyanine7 Rabbit IgG isotype control ($5 \mu\text{l}/\text{Test}$, left) or APC/Cyanine7 Rabbit anti-Human/Monkey IgD mAb ($5 \mu\text{l}/\text{Test}$, right). Cells in the CD11b⁻ gate were used for analysis.