

ABflo® 647 Rabbit anti-Human IgM mAb

Catalog No.: A24695

Basic Information

Observed MW

Calculated MW
49kDa

Category
Primary antibody

Applications
IF/ICC,FC

Cross-Reactivity
Human

CloneNo number
ARC63173-ABflo647

Conjugate
ABflo® 647. Ex:648nm. Em:664nm.

Recommended Dilutions

IF/ICC	1:50-1:200
FC	5 µl per 10 ⁶ cells in 100 µl volume

Contact

		400-999-6126
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		www.abclonal.com.cn

Background

Immunoglobulins (Ig) are the antigen recognition molecules of B cells. An Ig molecule is made up of 2 identical heavy chains and 2 identical light chains (see MIM 147200) joined by disulfide bonds so that each heavy chain is linked to a light chain and the 2 heavy chains are linked together. Each Ig heavy chain has an N-terminal variable (V) region containing the antigen-binding site and a C-terminal constant (C) region, encoded by an individual C region gene, that determines the isotype of the antibody and provides effector or signaling functions. The heavy chain V region is encoded by 1 each of 3 types of genes: V genes (see MIM 147070), joining (J) genes (see MIM 147010), and diversity (D) genes (see MIM 146910). The C region genes are clustered downstream of the V region genes within the heavy chain locus on chromosome 14. The IGHM gene encodes the C region of the mu heavy chain, which defines the IgM isotype. Naive B cells express the transmembrane forms of IgM and IgD (see IGHD; MIM 1471770) on their surface. During an antibody response, activated B cells can switch to the expression of individual downstream heavy chain C region genes by a process of somatic recombination known as isotype switching. In addition, secreted Ig forms that act as antibodies can be produced by alternative RNA processing of the heavy chain C region sequences. Although the membrane forms of all Ig isotypes are monomeric, secreted IgM forms pentamers, and occasionally hexamers, in plasma (summary by Janeway et al., 2005).[supplied by OMIM, Aug 2010]

Immunogen Information

Gene ID	Swiss Prot
3507	P01871-1

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 218-453 of human IgM(P01871).

Synonyms

IGHM; AGM1; MU; VH; immunoglobulin heavy constant mu

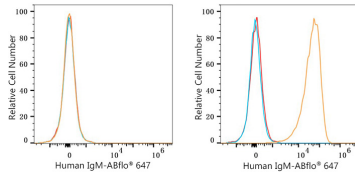
Product Information

Source	Isotype	Purification
Rabbit	IgM	Affinity purification

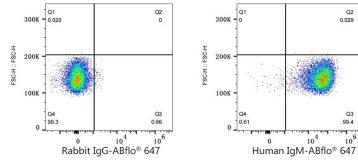
Storage

Store at 2-8°C. Avoid freeze.
Buffer: PBS with 0.03% proclin300,0.2% BSA,pH7.3.

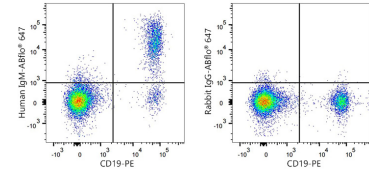
Validation Data



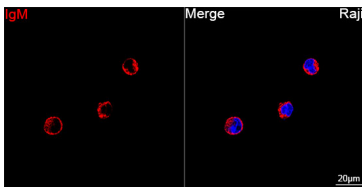
Flow cytometry: 1×10^6 Jurkat cells (negative control, left) and Daudi cells (right) were surface-stained with ABflo® 647 Rabbit anti-Human IgM mAb (A24695, 5 μ l/Test, orange line) or ABflo® 647 Rabbit IgG isotype control (A22070, 5 μ l/Test, blue line). Non-fluorescently stained cells were used as blank control (red line).



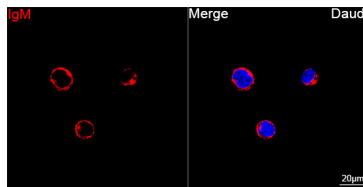
Flow cytometry: 1×10^6 Daudi cells were surface-stained with ABflo® 647 Rabbit IgG isotype control (A22070, 5 μ l/Test, left) or ABflo® 647 Rabbit anti-Human IgM mAb (A24695, 5 μ l/Test, right).



Flow cytometry: 1×10^6 Human PBMC were surface-stained with PE Mouse anti-Human CD19 mAb (A22816, 5 μ l/Test) and ABflo® 647 Rabbit anti-Human IgM mAb (A24695, 5 μ l/Test, left) or ABflo® 647 Rabbit IgG isotype control (A22070, 5 μ l/Test, right).



Confocal imaging of Raji cells using ABflo® 647 Rabbit anti-Human IgM mAb (A24695, dilution 1:100). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal imaging of Daudi cells using ABflo® 647 Rabbit anti-Human IgM mAb (A24695, dilution 1:100). DAPI was used for nuclear staining (Blue). Objective: 100x.