

ABflo® 488-conjugated Donkey anti-Rabbit IgG (H+L)

Catalog No.: AS035

13 Publications

Basic Information

Observed MW

Calculated MW

Category

Secondary antibody

Applications

IF/ICC,FC

Cross-Reactivity

Conjugate

ABflo® 488. Ex:491nm. Em:516nm.

Background

Secondary antibodies are affinity-purified antibodies which will work with target-specific primary antibody in the detection, sorting or purification of its specified target. Secondary antibodies offer increased versatility enabling users to use many detection systems (e.g. HRP, AP, fluorescence). They can also provide greater sensitivity through signal amplification as multiple secondary antibodies. Most commonly, secondary antibodies are generated by immunizing the host animal (different from host species of primary antibody) with a pooled population of normal immunoglobulins from the host species of primary antibody and can be further purified and modified (i.e. antibody fragmentation, label conjugation, etc.) to ensure well-characterized specificity to corresponding normal immunoglobulins.

Recommended Dilutions

IF/ICC 1:100 - 1:800

FC 1:100 - 1:800

Immunogen Information

Gene ID

Swiss Prot

Immunogen

This information is considered to be commercially sensitive.

Synonyms

Contact

 | 400-999-6126 | cn.market@abclonal.com.cn | www.abclonal.com.cn

Product Information

Source

Donkey

Isotype

ABflo™ 488 conjugated IgG

Purification

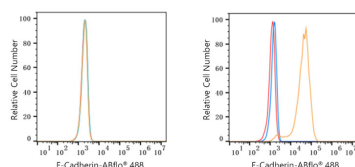
Affinity purification

Storage

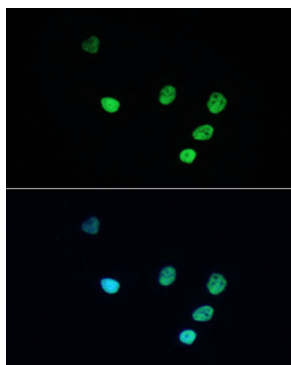
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.025% Sodium Azide, 0.75% BSA, 50% glycerol, pH7.3.

Validation Data



Flow cytometry: 1X10⁶ K-562 cells (negative control, left) and A-431 cells (right) were surface-stained with Purified Rabbit anti-Human E-Cadherin mAb (5 µl/Test, orange line) or secondary antibody only (blue line). Non-fluorescently stained K-562 and A-431 cells were used as blank control (red line). ABflo® 488-conjugated Donkey Anti-Rabbit IgG (H+L) (AS035, 1:800) was used as a secondary antibody.



Immunofluorescence analysis of HeLa cells, using PARP1 antibody (A0942) as the primary antibody at dilution of 1:100. The cells were incubated with the primary antibody overnight at 4°C. Secondary antibody: ABflo® 488-conjugated Donkey Anti-Rabbit IgG (H+L) (AS035) at 1:100 dilution. Blue: DAPI for nuclear staining.