

AS039

Leader in Biomolecular Solutions for Life Science



ABflo® 594-conjugated Goat anti-Rabbit IgG (H+L)

Catalog No.: AS039 **95 Publications**

Basic Information

Observed MW

Calculated MW

Category

Secondary Antibody

Applications

IF/ICC,FC

Cross-Reactivity

Rabbit

Conjugate

ABflo® 594. Ex:588nm. Em:604nm.

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 1:50 - 1:200

FC 1:100 - 1:1000

Immunogen Information

Gene ID

Swiss Prot

Immunogen

This information is considered to be commercially sensitive.

Synonyms

Contact

 www.abclonal.com

Product Information

Source

Goat

Isotype

Horseradish peroxidase
conjugated IgG

Purification

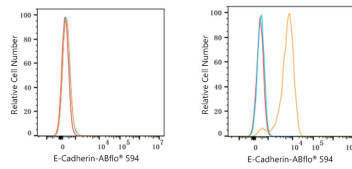
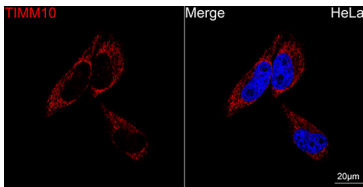
Affinity purification

Storage

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

Buffer: PBS containing 50% glycerol and 1% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

Validation Data



Confocal imaging of HeLa cells using [KD Validated] TIMM10 Rabbit mAb (A24996, dilution 1:200) followed by a further incubation with ABflo® 594-conjugated Goat Anti-Rabbit IgG (H+L) (AS039, dilution 1:400). DAPI was used for nuclear staining (Blue). Objective: 100x.

Flow cytometry: 1×10^6 K-562 cells (negative control, left) and A-431 cells (right) were surface-stained with Purified Rabbit anti-Human E-Cadherin mAb ($5 \mu\text{l}/\text{Test}$, 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® 594-conjugated Goat Anti-Rabbit IgG (H+L) (AS039, 1:1000) was used as a secondary antibody.