

AS007

Leader in Biomolecular Solutions for Life Science



Cy3-conjugated Goat anti-Rabbit IgG (H+L)

Catalog No.: AS007 **207 Publications**

Basic Information

Observed MW

Calculated MW

Category

Secondary Antibody

Applications

IF/ICC,FC

Cross-Reactivity

Rabbit

Conjugate

Cy3. Ex:548nm. Em:562nm.

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

 www.abclonal.com

Product Information

Source

Goat

Isotype

Cy3 conjugated IgG

Purification

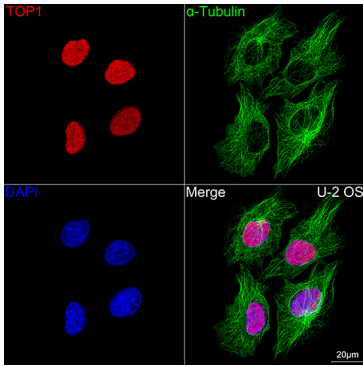
Affinity purification

Storage

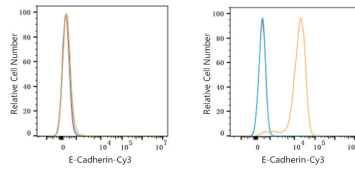
Store at -20°C. Avoid freeze / thaw cycles.

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

Validation Data



Confocal imaging of U-2 OS cells using DNA topoisomerase I (TOP1) Rabbit mAb(A12409,dilution 1:100)(Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012,dilution 1:400) (Green). DAPI was used for nuclear staining (blue). Objective: 100x.Secondary antibody: Cy3 Goat Anti-Rabbit IgG (H+L) (AS007,dilution 1:100)(Red),ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L)(AS076,dilution 1:200) (Green)



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 μ l/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). Cy3 Goat Anti-Rabbit IgG (H+L)(AS007, 1:800) was used as a secondary antibody.