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ABflo® 488 Rabbit anti-Human/Monkey Ki67 mAb

Catalog No.: A23407

Basic Information

Observed MW

Refer to figures

Calculated MW

319kDa/359kDa

Category

Primary antibody

Applications

FC (intra)

Cross-Reactivity

Human, Cynomolgus

CloneNo number

ARC57564

Conjugate

ABflo® 488. Ex:491nm. Em:516nm.

Background

Enables protein C-terminus binding activity. Involved in regulation of chromosome segregation and regulation of mitotic nuclear division. Located in chromosome; nuclear body; and nucleolus. Colocalizes with condensed chromosome. Implicated in Crohn's disease; breast cancer; human immunodeficiency virus infectious disease; and pancreatic cancer. Biomarker of several diseases, including Barrett's esophagus; autoimmune disease of musculoskeletal system (multiple); endocrine gland cancer (multiple); gastrointestinal system cancer (multiple); and interstitial cystitis.

Recommended Dilutions

FC (intra)

5 μl per 10^6 cells in 100 μl volume

Immunogen Information

Gene ID 4288 **Swiss Prot**

P46013

Immunogen

Synthetic peptide. This information is considered to be commercially sensitive.

Synonyms

KIA; MIB-; MIB-1; PPP1R105

Contact

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Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

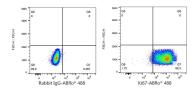
Store at 2-8°C. Avoid freeze.

Buffer: PBS containing 0.2% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

Validation Data



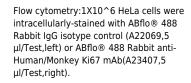


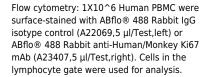


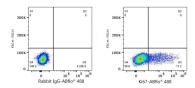




Flow cytometry:1X10^6 knockout (KO) HeLa cells(negative control,left) and HeLa cells (right) were intracellularly-stained with ABflo® 488 Rabbit anti-Human/Monkey Ki67 mAb(A23407,5 µl/Test,orange line) or ABflo® 488 Rabbit IgG isotype control (A22069,5 µl/Test,blue line).Non-fluorescently stained cells were used as blank control (red line).







Flow cytometry: $1X10^6$ Cynomolgus PBMC were surface-stained with ABflo® 488 Rabbit IgG isotype control (A22069,5 μ I/Test,Ieft) or ABflo® 488 Rabbit anti-Human/Monkey Ki67 mAb (A23407,5 μ I/Test,right). Cells in the lymphocyte gate were used for analysis.