

ABflo® 610 Rabbit anti-Human/Monkey Ki67 mAb

Catalog No.: A26736

Basic Information

Observed MW

Calculated MW

319kDa/359kDa

Category

Primary antibody

Applications

FC (intra)

Cross-Reactivity

Human, Cynomolgus

CloneNo number

ARC57564

Conjugate

ABflo® 610. Ex:421nm. Em:612nm.

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

2		400-999-6126
\bowtie		cn.market@abclonal.com.cn
•	T	www.abclonal.com.cn

Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at 2-8°C. Avoid freeze.

Buffer: PBS with 0.09% Sodium azide, 0.2% BSA, pH7.3.

Validation Data









Flow cytometry: 1X10^6 knockout (KO) HeLa cells (negative control,left) and HeLa cells (right) were intracellularly-stained with ABflo® 610 Rabbit anti-Human/Monkey Ki67 mAb (A26736,5 µl/Test,orange line) or ABflo® 610 Rabbit IgG isotype control (A25826,5 µl/Test,blue line). Nonfluorescently stained cells were used as blank control (red line).

Flow cytometry: 1X10^6 Human PBMC (untreated,left) and Human PBMC (treated with 5 μ g/mL PHA for 72 hours,right) were intracellularly-stained with ABflo® 450 Rabbit anti-Human cCD3 mAb (A27836,5 μ l/Test) and ABflo® 610 Rabbit anti-Human/Monkey Ki67 mAb (A26736,5 μ l/Test). Cells in the lymphocyte gate were used for analysis.