

Rabbit anti-human IgG FC (YTE mutation) Specific Monoclonal Antibody, Clone 6H2

Catalog No.: YR0510

Basic Information

Molecular Weight

Endotoxin

<1EU/mg (<0.001EU/μg) Determined by LAL gel clotting assay

Sterility

Aggregation

Purity

Background

Related studies have shown that M252Y/S254T/T256E (YTE) mutations in the Fc region of IgG antibodies can prolong the half-life of therapeutic antibodies, optimize the pharmacokinetic characteristics, and reduce the frequency of administration. We developed Rabbit Anti-human IgG-FC (YTE mutation) Specific Monoclonal Antibody that effectively captures and stably binds to IgG Fc segment YTE mutants. The YTE mutant for therapeutic IgG has important application value in pharmacodynamic/pharmacokinetic studies and detection technology.

Reported Applications

ELISA

Immunogen Information

Clone

Isotype

Immunogen


Recommended Isotype Control(s)

Recommended Dilution Buffer

Contact

 | 400-999-6126

 | cn.market@abclonal.com.cn

 | www.abclonal.com.cn

Product Information

Production

Purified from cell culture supernatant in an animal-free facility

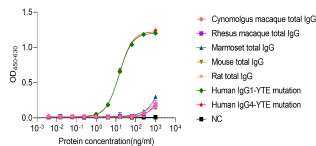
Purification

Storage

Store at 2 - 8°C. 2 - 8°C for up to 4 weeks and -80°C for long term storage (Avoid repeated freezing and thawing)

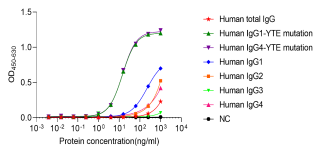
Validation Data

Rabbit Anti-human IgG-FC(YTE mutation) Specific Monoclonal Antibody (Clone 6H2)

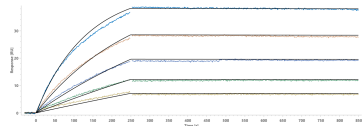


A microtiter plate was coated overnight with Rabbit Anti-human IgG-FC (YTE mutation) Specific Monoclonal Antibody (Clone 6H2) at a concentration of 1 µg/mL. After washing and blocking with PBST + 5% skim milk, increasing concentrations of various biotinylated antibodies was added. Horseradish Peroxidase conjugated Neutravidin, (Thermo Fisher, 31001) was used for final detection.

Rabbit Anti-human IgG-FC(YTE mutation) Specific Monoclonal Antibody (Clone 6H2)



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Rabbit Anti-human IgG-FC (YTE mutation) Specific Monoclonal Antibody (Clone 6H2) captured on Protein A Chip can bind Human IgG1-FC (YTE) mutation with an affinity constant of 0.2 nM as determined in SPR assay.