

# Rabbit anti-Human IgG Specific Monoclonal Antibody, clone 1C1 (minimum binding to cyno/Ms/rat IgG)

Catalog No.: YR0504

## **Basic Information**

#### **Molecular Weight**

#### **Endotoxin**

<1EU/mg (<0.001EU/ $\mu$ g)Determined by LAL gel clotting assay

#### Sterility

0.2 µm filtration

### **Aggregation**

<5% Determined by SECP

#### Purity

>95% Determined by SDS-PAGE

## **Background**

Rabbit anti-Human IgG Specific Monoclonal Antibody clone 1C1 is a recombinant rabbit Monoclonal antibody, Capture ELISA binding experiment results that recognizes human immunoglobulin isotypes (IgG1, IgG2, IgG3, IgG4), IgG-Fab, total IgG. It does not recognize Cynomolgus macaque total IgG, Rhesus macaque total IgG, Marmoset total IgG, Mouse total IgG, Rat total IgG.

# **Reported Applications**

#### **ELISA**

## **Immunogen Information**

Clone 1C1

Isotype

lgG1

**Immunogen** 

Human IgG Fab

RecommendedIsotype Control(s)

**Recommended Dilution Buffer** 

1×PBS pH 7.0

## **Contact**

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•	www.abclonal.com.cn

## **Product Information**

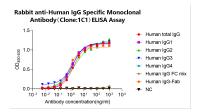
**Production**Purified from cell culture supernatant in an animal-free facility

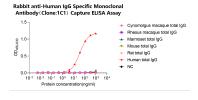
Protein A/G

# Storage

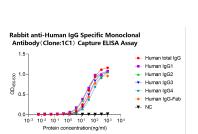
 $2 - 8^{\circ}$ C for up to 4 weeks and -80°C for long term storage (Avoid repeated freezing and thawing)



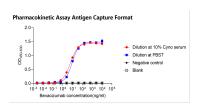




Regular ELISA binding curve demonstrating the recognition of Rabbit anti-Human IgG Specific Monoclonal Antibody. A microtiter plate was coated overnight with various antibodies at a concentration of 2  $\mu$ g/mL. After washing and blocking with PBST + 5% skim milk, increasing concentrations of the Rabbit Anti-Human IgG Fab Antibody (clone 1C1) was added. Horseradish Peroxidase conjugated goat anti-rabbit IgG (Jackson, Cat.: 111-035-045) was used for final detection.



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Capture ELISA binding curve demonstrating the recognition of Rabbit anti-Human IgG Specific Monoclonal Antibody. A microtiter plate was coated overnight with Rabbit anti-Human IgG Specific Monoclonal Antibody (clone 1C1) at a concentration of 2  $\mu$ g/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.

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A microtiter plate was coated overnight with VEGF at a concentration of 2  $\mu g/mL$ . After washing and blocking with PBST + 5% skim milk, 10% cynomolgus macaque serum was added spiked with increasing concentrations of bevacizumab (hlgG1 kappa)(from 10  $\mu g/mL$ ). Clone 1C1 was added. Another unrelated rabbit mAb used for negative control. Detection was performed using HRP conjugated goat anti-rabbit 2nd Ab. No cynomolgus serum added sample is used as control.