## **CD163 Recombinant Antibody**



## **RACO0045**

Source:

Human

## **Product Information**

Size: Protein Background:

50ul Acute phase-regulated receptor involved in clearance and endocytosis of

**Reactivity:**hemoglobin/haptoglobin complexes by macrophages and may thereby protect tissues from free hemoglobin-mediated oxidative damage. May play a role in the uptake and

Human recycling of iron, via endocytosis of hemoglobin/haptoglobin and subsequent

breakdown of heme. Binds hemoglobin/haptoglobin complexes in a calcium-

dependent and pH-dependent manner. Exhibits a higher affinity for complexes of

 $he moglobin \ and \ multimeric \ haptoglobin \ of \ HP*1F \ phenotype \ than \ for \ complexes \ of$ 

hemoglobin and dimeric haptoglobin of HP\*1S phenotype.

Isotype: Gene ID:

Rabbit IgG CD163

Applications: Uniprot

ELISA, IHC, FC Q86VB7

Recommended dilutions: Synonyms:

IHC:1:50-1:500 Scavenger receptor cysteine-rich type 1 protein M130, Hemoglobin scavenger receptor,

CD163, Soluble CD163, sCD163, CD163, M130

Immunogen:

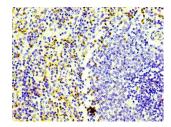
A synthesized peptide.

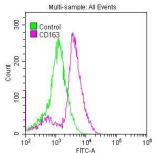
Storage:

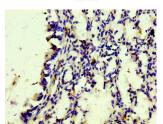
Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and

50% glycerol.

## **Product Images**







IHC image of RACO0045 diluted at 1:100 and staining in paraffinembedded human spleen tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

Overlay histogram showing Raw264.7 cells stained with RACO0045 (red line) at for 4h). The cells were fixed with 70% Ethylalcohol (18h) and then permeabilized with 0.3% Triton X-100 for 2 min. The cells were then incubated in 1x PBS /10% normal goat serum to block non-specific protein-protein interactions followed by primary antibody for 1 h at 4°C. The secondary antibody used was FITC goat anti-rabbit IgG (H+L) at 1/200 dilution for 1 h at 4°C. Control antibody (green line) was used under the same conditions. Acquisition of >10,000 events was performed.

IHC image of RACO0045 diluted at 1:100 and staining in paraffinembedded human lung tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.