

Basic Information

Product Name	Anti-CD163 Antibody	
Gene Name	CD163	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, IHC, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN ₃ , 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human CD163 recombinant protein (Position: T47-E201). Human CD163 shares 78.4% and 79.7% amino acid (aa) sequence identity with mouse and rat CD163, respectively.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	125-150 kDa	
Dilution Ratios	Western blot (WB): 1:500-2000 Immunohistochemistry (IHC): 1:50-400 Immunohistochemistry in frozen section: 1:50-400 Immunocytochemistry in fixed cells: 1:50-400 Flow cytometry (FCM): 1-3 µg/1x10 ⁶ cells ELISA: 1:100-1000 (Boiling the paraffin sections in 10mM citrate buffer, pH6.0, or PH8.0 EDTA repair liquid for 20 mins is required for the staining of formalin/paraffin sections.) Optimal working dilutions must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

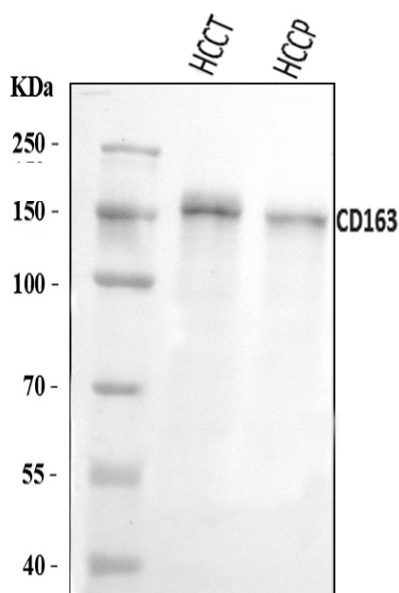
CD163 (Cluster of Differentiation 163) is a protein that in humans is encoded by the CD163 gene. The protein encoded by this gene is a member of the scavenger receptor cysteine-rich (SRCR) superfamily, and is exclusively expressed in monocytes and macrophages. It functions as an acute phase-regulated receptor involved in the clearance and endocytosis of hemoglobin/haptoglobin complexes by macrophages, and may thereby protect tissues from free hemoglobin-mediated oxidative damage. This protein may also function as an innate immune sensor for bacteria and inducer of local

inflammation. Alternatively spliced transcript variants encoding different isoforms have been described for this gene.

Reference

Anti-CD163 Antibody 被引用在3文献中。

Selected Validation Data

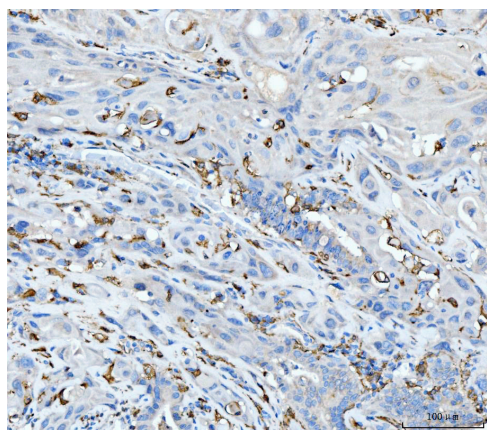


Western blot analysis of CD163 using anti-CD163 antibody (A00812-1). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: HCCT tissue lysates,

Lane 2: HCCP tissue lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-CD163 antigen affinity purified polyclonal antibody (A00812-1) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for CD163 at approximately 125-150 kDa. The expected band size for CD163 is at 125 kDa.



IHC analysis of CD163 using anti-CD163 antibody (A00812-1). CD163 was detected in a paraffin-embedded section of human Gall bladder adenosquamous carcinoma tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-CD163 Antibody (A00812-1) at a dilution of 1:200 and developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.