

CD36 (PT0434R) PT® Rabbit mAb

Catalog No :	YM8274
Reactivity :	Human; Mouse; Rat;
Applications :	WB;IHC;IF;IP;ELISA
Target :	CD36
Fields :	>>PPAR signaling pathway;>>Phagosome;>>AMPK signaling pathway;>>ECM-receptor interaction;>>Hematopoietic cell lineage;>>Adipocytokine signaling pathway;>>Insulin resistance;>>Fat digestion and absorption;>>Cholesterol metabolism;>>Malaria;>>Diabetic cardiomyopathy;>>Lipid and atherosclerosis
Gene Name :	CD36
Protein Name :	Platelet glycoprotein 4
Human Gene Id :	948
Human Swiss Prot No :	P16671
Mouse Gene Id :	12491
Mouse Swiss Prot No :	Q08857
Rat Swiss Prot No :	Q07969
Specificity :	endogenous
Formulation :	PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA
Source :	Monoclonal, rabbit, IgG, Kappa
Dilution :	IHC 1:1000-1:4000;WB 1:1000-1:5000;IF 1:200-1:1000;ELISA 1:5000-1:20000;IP 1:50-1:200;
Purification :	Protein A

Storage Stability : -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight : 53kD

Observed Band : 90kD

Cell Pathway : PPAR;ECM-receptor interaction;Hematopoietic cell lineage;Adipocytokine;

Background :

The protein encoded by this gene is the fourth major glycoprotein of the platelet surface and serves as a receptor for thrombospondin in platelets and various cell lines. Since thrombospondins are widely distributed proteins involved in a variety of adhesive processes, this protein may have important functions as a cell adhesion molecule. It binds to collagen, thrombospondin, anionic phospholipids and oxidized LDL. It directly mediates cytoadherence of Plasmodium falciparum parasitized erythrocytes and it binds long chain fatty acids and may function in the transport and/or as a regulator of fatty acid transport. Mutations in this gene cause platelet glycoprotein deficiency. Multiple alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Feb 2014],

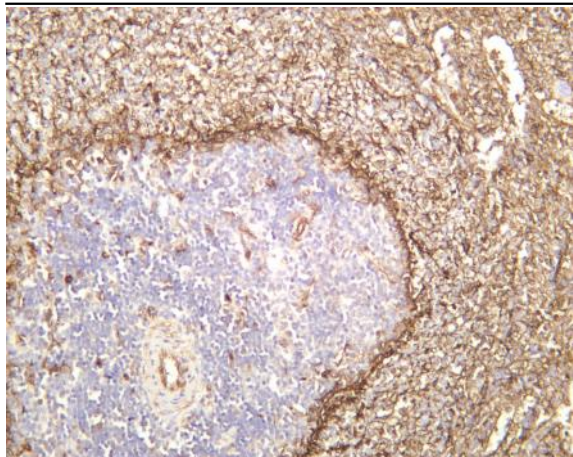
Function :

disease:Defects in CD36 are the cause of platelet glycoprotein IV deficiency [MIM:608404]; also known as CD36 deficiency. Platelet glycoprotein IV deficiency can be divided into 2 subgroups. The type I phenotype is characterized by platelets and monocytes/macrophages exhibiting complete CD36 deficiency. The type II phenotype lacks the surface expression of CD36 in platelets, but expression in monocytes/macrophages is near normal.,disease:Genetic variations in CD36 are associated with susceptibility to coronary heart disease type 7 (CHDS7) [MIM:610938].,function:Seems to have numerous potential physiological functions. Binds to collagen, thrombospondin, anionic phospholipids and oxidized LDL. May function as a cell adhesion molecule. Directly mediates cytoadherence of Plasmodium falciparum parasitized erythrocytes. Binds long chain fatty acids and may function in the transport and/or as a

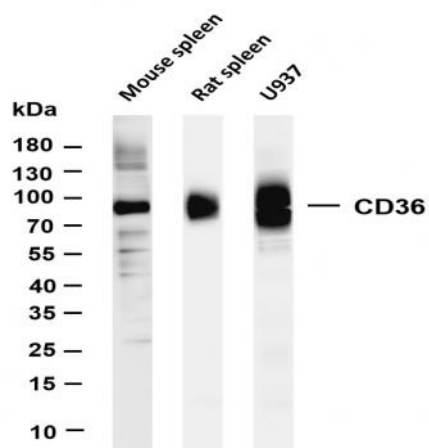
Subcellular Membrane**Location :**

Expression : Adipocyte,Liver,Mammary gland,Milk,Placenta,Platelet,Skeletal muscle,

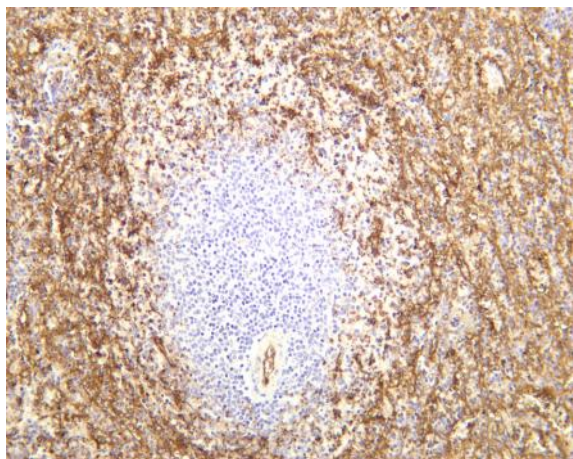
Products Images



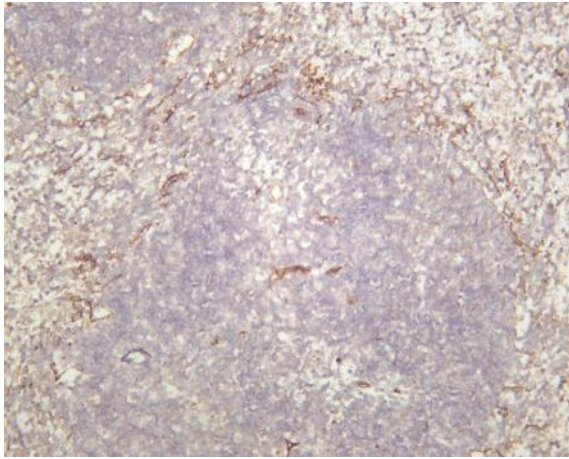
Rat spleen was stained with anti-CD36 (PT0434R) rabbit antibody



Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-CD36 (PT0434R) antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: Mouse spleen Lane 2: Rat spleen Lane 3: U937 Predicted band size: 53kDa Observed band size: 90kDa



Human spleen was stained with anti-CD36 (PT0434R) rabbit antibody



Mouse spleen was stained with anti-CD36 (PT0434R) rabbit antibody