abcam

Product datasheet

Human Von Willebrand Factor ELISA Kit (VWF) ab108918

23 References 2 图像

概述

检**测方法** Colorimetric

精确度

样品	n	Mean	SD	CV%
Overall				5.6%

批次间

批次内

样品	n	Mean	SD	CV%
Overall				9.6%

样**品**类型 Cell culture supernatant, Serum, Plasma

检测类型 Sandwich (quantitative)

灵敏度 = 0.38 mIU/ml

范围 0.625 mlU/ml - 40 mlU/ml

 回收率
 98 %

 检测时间
 5h 00m

实验**步**骤 Multiple steps standard assay

种属反应性 与反应: Human

产品概述 Human Von Willebrand Factor (VWF) in vitro ELISA (Enzyme-Linked Immunosorbent Assay) kit

is designed for the quantitative measurement of Human Von Willebrand Factor in plasma, serum,

and cell culture supernatants.

A Von Willebrand Factor specific antibody has been precoated onto 96-well plates and blocked. Standards or test samples are added to the wells and subsequently a Von Willebrand Factor specific biotinylated detection antibody is added and then followed by washing with wash buffer. Streptavidin-Peroxidase Conjugate is added and unbound conjugates are washed away with wash buffer. TMB is then used to visualize Streptavidin-Peroxidase enzymatic reaction. TMB is catalyzed by Streptavidin-Peroxidase to produce a blue color product that changes into yellow after adding acidic stop solution. The density of yellow coloration is directly proportional to the amount of Von Willebrand Factor captured in plate.

1

The entire kit may be stored at -20°C for long term storage before reconstitution - Avoid repeated freeze-thaw cycles.

平台

Microplate

性能

存放说明

Store at -20°C. Please refer to protocols.

组件	1 x 96 tests
100X Streptavidin-Peroxidase Conjugate	1 x 80µl
10X Diluent N Concentrate	1 x 30ml
20X Wash Buffer Concentrate	2 x 30ml
100X Biotinylated Human Von Williebrand Factor Antibody	1 x 60µl
Chromogen Substrate	1 x 7ml
Sealing Tapes	3 units
Stop Solution	1 x 11ml
Von Williebrand Factor Microplate (12 x 8 well strips)	1 unit
Von Williebrand Factor Standard	1 vial

功能

Important in the maintenance of hemostasis, it promotes adhesion of platelets to the sites of vascular injury by forming a molecular bridge between sub-endothelial collagen matrix and platelet-surface receptor complex GPIb-IX-V. Also acts as a chaperone for coagulation factor VIII, delivering it to the site of injury, stabilizing its heterodimeric structure and protecting it from premature clearance from plasma.

组织特异性

疾病相关

Plasma.

Defects in VWF are the cause of von Willebrand disease (VWD) [MIM:277480]. VWD defines a group of hemorrhagic disorders in which the von Willebrand factor is either quantitatively or qualitatively abnormal resulting in altered platelet function. Symptoms vary depending on severity and disease type but may include prolonged bleeding time, deficiency of factor VIII and impaired platelet adhesion. Type I von Willebrand disease is the most common form and is characterized by partial quantitative plasmatic deficiency of an otherwise structurally and functionally normal Willebrand factor; type II is associated with a qualitative deficiency and functional anomalies of the Willebrand factor; type III is the most severe form and is characterized by total or near-total absence of Willebrand factor in the plasma and cellular compartments, also leading to a profound deficiency of plasmatic factor VIII.

序列相似性

Contains 1 CTCK (C-terminal cystine knot-like) domain.

Contains 4 TIL (trypsin inhibitory-like) domains.

Contains 3 VWFA domains.
Contains 3 VWFC domains.
Contains 4 VWFD domains.

结**构域** The von Willebrand antigen 2 is required for multimerization of vWF and for its targeting to

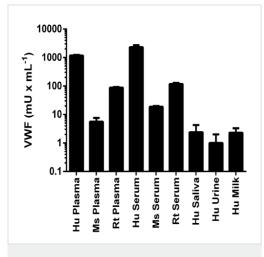
storage granules.

翻译后修饰 All cysteine residues are involved in intrachain or interchain disulfide bonds.

N- and O-glycosylated.

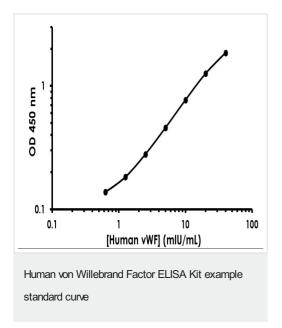
细胞定位 Secreted Secret

图片



VWF measured in biological fluids with background signal subtracted (duplicates +/- SD). There is limited cross-reactivity in mouse (< 2%) and rat samples (< 15%).





Example von Willebrand Factor standard curve – data provided for demonstration purposes only.

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