Anti-Factor VIII:c light chain [RFF-VIII:c/5]

Catalogue number: 153359
Sub-type:
Images:

Contributor

Inventor: Alison Goodall
Institute: University College London (UCL) ; Absolute Antibody
Images:

Tool details

***FOR RESEARCH USE ONLY**

Name: Anti-Factor VIII:c light chain [RFF-VIII:c/5]

Alternate name:

Class: Recombinant
Conjugate: Unconjugated
Description: The commonest severe congenital bleeding disorder in all races is haemophilia A. The characteristic defect is a lack of coagulation factor VIII:C. Factor VIII:C is a glycoprotein that functions as a cofactor for factor IXa which, in the presence of calcium and phospholipids, converts factor X to the activated form Xa. This antibody may be used for the purification of Factor VIII from normal plasma and cryoprecipitate.
Purpose:
Parental cell:
Organism:
Tissue:
Model:
Gender:
Isotype: IgG1 kappa
Reactivity: Human
Selectivity:
Host: Mouse
Immunogen: Purified Human FVIII
Immunogen UNIPROT ID:
Sequence:
Growth properties:
Production details:
Formulation:
Recommended controls:

Bacterial resistance:
Selectable markers:
Additional notes:

Target details

Target: Factor VIII:c light chain

Target alternate names:

Target background: The commonest severe congenital bleeding disorder in all races is haemophilia A. The characteristic defect is a lack of coagulation factor VIII:C. Factor VIII:C is a glycoprotein that functions as a cofactor for factor IXa which, in the presence of calcium and phospholipids, converts factor X to the activated form Xa. This antibody may be used for the purification of Factor VIII from normal plasma and cryoprecipitate.

Molecular weight:

Ic50:

Applications

Application: IF ; WB
Application notes:

Handling

Format: Liquid
Concentration: 1mg/ml
Passage number:
Growth medium:
Temperature:
Atmosphere:
Volume:
Storage medium:
Storage buffer: RPMI + 10% FCS; non-adherent; subculture every 2-3 days; split 1:5
Storage conditions:
Shipping conditions: Shipping at 4° C

Related tools

Related tools:

References

References: Goodall AH et al. 1985. Thromb Haemost. 54(4):878-91. PMID: 3937279