

Murine Anti-Factor VIII

Clone GMA-8029

Factor VIII (FVIII) is a heterodimer consisting of a heavy chain (ranging in mass from 90 to 200 kDa) bound via metal ions to a light chain (80 kDa). In plasma, FVIII circulates in an inactive form bound to von Willebrand factor. Following activation by factor Xa or thrombin, factor VIIIa can function as cofactor for the enzyme factor IXa in the activation of factor X in the presence of phospholipid and Ca²⁺. Absent or defective FVIII is the cause of the Xlinked recessive bleeding disorder hemophilia A. GMA-8029 (also known as 3D12) recognizes the C2 domain of FVIII, and inhibits the binding of FVIII to phospholipid membranes.¹ It is suitable for ELISA, as well as surface plasmon resonance experiments.²

Description

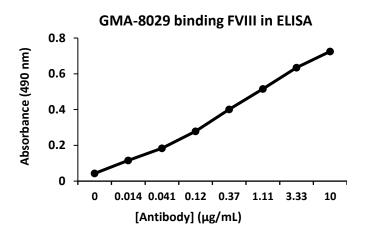
| Antibody Source: | mouse monoclonal, IgG _{2b} |
|------------------------|---|
| Antigen Species Bound: | human |
| Specificity: | FVIII C2 domain |
| Immunogen: | B-domain deleted recombinant human FVIII |

Formulation and Storage

| Purity: | Purified by protein G affinity chromatography from serum-free cell culture supernatant. |
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| Product Formulation: | Lyophilized from a ≥ 1 mg/ml solution in 20 mM NaH ₂ PO ₄ 0.15 M NaCl, 1.0% (w/v) mannitol, pH 7.4. Concentration determined by absorbance measurement at 280 nm using an extinction coefficient of 1.4 ($\varepsilon_{0.1\%}$). |
| Reconstitution: | Reconstitute with deionized water. |
| Storage: | Store lyophilized or reconstituted and aliquoted material at -20° C for prolonged periods. Avoid freeze-thaw cycles. Alternatively, add 0.02% (w/v) sodium azide to reconstituted solution and store at 4° C. |
| Country of Origin: | USA |
| Size Options: | 0.1 mg or 0.5 mg |

| Арр | lications | |
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| Working Concentration: | Approximately 1-5 μg/ml. Researcher should titer antibody in specific assay. |
|------------------------|--|
| ELISA: | Binds immobilized human FVIII. |
| Immunoblotting: | Not recommended. |
| Inhibition: | Inhibitory in aPTT clotting assay. ¹ |



References

[1] S.L. Meeks, J.F. Healey, E.T. Parker, R.T. Barrow, P. Lollar. Antihuman factor VIII C2 domain antibodies in hemophilia A mice recognize a functionally complex continuous spectrum of epitopes dominated by inhibitors of factor VIII activation. (2007). *Blood*. 110(13):4234-4242.

[2] P.T. Nguyen, K.B. Lewis, R.A. Ettinger, J.T. Schuman, J.C. Lin, J.F. Healey, S.L. Meeks, P. Lollar, K.P. Pratt. High-resolution mapping of epitopes on the C2 domain of factor VIII by analysis of point mutants using surface plasmon resonance. (2014). *Blood.* 123(17):2732-2739.

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