

Anti-Coagulation factor IX/F9 Antibody (1B255)

Product Details

Ig Type:	Rabbit IgG
Reactivity:	Human
Conjugation:	Unconjugated
Clone:	1B255
Purification:	Protein A

Applications

Verified Activity:	<p>1. Anti-F9 rabbit monoclonal antibody at 1:500 dilution.</p> <ul style="list-style-type: none">-Lane A: A431 Whole Cell Lysate.-Lane B: Jurkat Whole Cell lysate.-Lysates/proteins at 30 µg per lane.-Secondary-Goat Anti-Rabbit IgG H&L (Dylight800) at 1/10000 dilution.-Developed using the Odyssey technique.-Performed under reducing conditions.-Predicted band size:52 kDa.-Observed band size:55 kDa. <p>2. F9 was immunoprecipitated using:</p> <ul style="list-style-type: none">-Lane A:0.5 mg A431 Whole Cell Lysate.-Lane B:0.5 mg Jurkat Whole Cell Lysate-0.5 µL anti-F9 rabbit monoclonal antibody and 60 µg of Immunomagnetic beads Protein A/G.-Primary antibody:-Anti-F9 rabbit monoclonal antibody, at 1:500 dilution.-Secondary antibody:-Clean-Blot IP Detection Reagent (HRP) at 1:1000 dilution.-Developed using the ECL technique.-Performed under reducing conditions.-Predicted band size: 55 kDa.-Observed band size:55 kDa
Application:	ELISA,IP,WB
Recommended	WB: 1:500-1:1000; ELISA: 1:25000-1:50000; IP: 0.1-0.5 µL/mg of lysate

Properties

Stability & Storage:	Store at 2°C-8°C for 1 month. Store at -20°C or -80°C for 12 months. Avoid repeated freeze-thaw cycles. Preservative-Free.
Shipping:	Shipping with blue ice.

Antigen Details

Immunogen: Recombinant Protein: Human Coagulation Factor IX / FIX / F9 protein (TMPY-02215)

Antigen Species: Human

Synonyms: coagulation factor IX

Research Background

Coagulation factor IX, also known as Christmas factor, Plasma thromboplastin component and PTC, is a secreted protein which belongs to the peptidase S1 family. Coagulation factor IX / F9 contains two EGF-like domains, one Gla (gamma-carboxy-glutamate) domain and one peptidase S1 domain. Coagulation factor IX / F9 is a vitamin K-dependent plasma protein that participates in the intrinsic pathway of blood coagulation by converting factor X to its active form in the presence of Ca^{2+} ions, phospholipids, and factor VIIIa. Defects in Coagulation factor IX / F9 are the cause of thrombophilia due to factor IX defect which is a hemostatic disorder characterized by a tendency to thrombosis. Defects in Coagulation factor IX / F9 are also the cause of recessive X-linked hemophilia B (HEMB) which also known as Christmas disease.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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