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Human F9 Antibody Pair Set

Catalog No. E-KAB-0198 Applications ELISA

Synonyms HEMB, P19, PTC, THPH8, Christmas Factor

Kit components & Storage

Title	Specifications	Storage
Human F9 Capture Antibody	1 vial, 100 μ g	Store at -20°C for one year.
		Avoid freeze / thaw cycles.
Human F9 Detection Antibody (Biotin)	1 vial, 50 μL	Store at -20°C for one year.
		Avoid freeze / thaw cycles.

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Product Information

Items		Characteristic (E-KAB-0198)		
		Human F9 Capture Antibody	Human F9 Detection Antibody (Biotin)	
Immunogen	Immunogen	Recombinant Human F9 protein	Recombinant Human F9 protein	
Information	Swissprot	P00740		
Product details	Reactivity	Human	Human	
	Host	Rabbit	Rabbit	
	Conjugation	Unconjugated	Biotin	
	Concentration	0.5mg/mL	/	
	Buffer	PBS with 0.04% Proclin 300, 50%	PBS with 0.04% Proclin 300, 1%	
		glycerol, pH 7.4	protective protein, 50% glycerol, pH	
			7.4	
	Purify	Protein A & Antigen Affinity	Protein A & Antigen Affinity	
	Specificity	Detects Human F9 in ELISAs.		

For Research Use Only

Toll-free: 1-888-852-8623 Tel: 1-832-243-6086 Fax: 1-832-243-6017 Web: www.elabscience.com Email: techsupport@elabscience.com





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Applications

Human F9 Sandwich ELISA Assay:

	Recommended	Reagent	Images
	Concentration/Dilution		
ELISA	0.5-4μg/mL	Human F9 Capture Antibody	
Capture			10 10 10 10 10 10 10 10 10 10 10 10 10 1
ELISA Detection	1:1000-1:10000	Human F9 Detection Antibody (Biotin)	On the state of th

Note: This standard curve is only for demonstration purposes. A standard curve should be generated for each assay!

Background

This gene encodes vitamin K-dependent coagulation factor IX that circulates in the blood as an inactive zymogen. This factor is converted to an active form by factor XIa, which excises the activation peptide and thus generates a heavy chain and a light chain held together by one or more disulfide bonds. The role of this activated factor IX in the blood coagulation cascade is to activate factor X to its active form through interactions with Ca+2 ions, membrane phospholipids, and factor VIII. Alterations of this gene, including point mutations, insertions and deletions, cause factor IX deficiency, which is a recessive X-linked disorder, also called hemophilia B or Christmas disease.

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