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Monkey IGF-1 Antibody Pair Set

Catalog No. E-KAB-0655 Applications ELISA

Synonyms IGF1;IGF1;IGF-I;IGF1A;IGF-IA;IGF-IB;MGF;Somatomedin C

Kit components & Storage

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

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

Product Information

Items		Characteristic (E-KAB-0655)		
		Monkey IGF-1 Capture Antibody	Monkey IGF-1 Detection Antibody	
			(Biotin)	
Immunogen	Immunogen	Recombinant Monkey IGF-1 protein	Recombinant Monkey IGF-1 protein	
Information	Swissprot	A0A2K5URV9		
Product details	Reactivity	Monkey	Monkey	
	Host	Mouse	Mouse	
	Conjugation	Unconjugated	Biotin	
	Concentration	0.5 mg/mL	/	
	Buffer	PBS with 0.04% Proclin 300; 50%	PBS with 0.04% Proclin 300; 1%	
		glycerol; pH 7.5	protective protein; 50% glycerol; pH	
			7.5	
	Purify	Protein A or G	Protein A or G	
	Specificity	Detects Monkey IGF-1 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

Monkey IGF-1 Sandwich ELISA Assay

	Recommended	Reagent	Images
	Concentration/Dilution		
ELISA	0.5-4 μg/mL	Monkey IGF-1 Capture	
Capture		Antibody	10
			Optical Density
ELISA	1:1000-1:10000	Monkey IGF-1 Detection	Optic
Detection		Antibody (Biotin)	
			0.1 10 100 1000 Monkey IGF-1 Concentration(ng/mL)

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

Background

IGF1, also named as IBP1, MGF, IGF-IA and Somatomedin-C, belongs to the insulin family. IGF1 is structurally and functionally related to insulin but have a much higher growth-promoting activity. Altered expression or mutation of IGF-1 is associated with several human disorders, including type I diabetes and various forms of cancer. Defects in IGF1 are the cause of insulin-like growth factor I deficiency (IGF1 deficiency) which is an autosomal recessive disorder characterized by growth retardation, sensorineural deafness and mental retardation. The antibody is specific to isoform IGF-1A.

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