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Human Aβ1-42 Antibody Pair Set

Catalog No. E-KAB-0447 Applications ELISA

Synonyms Aβ (1-42); Amyloid Beta 42; Amyloid Beta 42

Kit components & Storage

Title	Specifications	Storage
Human Aβ1-42 Capture Antibody	1 vial, 100 μ g	Store at -20℃ for one year. Avoid
		freeze/thaw cycles.
Human Aβ1-42 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-0447)	
		Hymnon AR1 42 Continue Antibody	Human Aβ1-42 Detection Antibody
		Human Aβ1-42 Capture Antibody	(Biotin)
Immunogen	Immunogen	Recombinant Human Aβ1-42 protien	Recombinant Human Aβ1-42 protien
Information	Swissprot	P05067	
Product details	Reactivity	Human	Human
	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 Human Aβ1-42 in ELISAs.	

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Applications

Human Aβ1-42 Sandwich ELISA Assay

	Recommended	Reagent	Images
	Concentration/Dilution		
ELISA	0.5-4 μg/mL	Human Aβ1-42 Capture	
Capture		Antibody	10
			o opical Density
ELISA	1:1000-1:10000	Human Aβ1-42 Detection	O O Dig
Detection		Antibody (Biotin)	0.01
			10 100 1000 10000
			Human Aβ1-42 Concentration(pg/mL)

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

Background

A β derives from APP via proteolytic cleavage by proteases called α -, β - and γ -secretase. The α -secretase cleavage precludes the formation of A β , while the β - and γ -cleavages generate APP components with amyloidogenic features. Amyloid beta A4 precursor protein (APP), encoded by APP gene which locate on human chromosome 21q, is a cell surface receptor and performs physiological functions on the surface of neurons relevant to neurite growth, neuronal adhesion and axonogenesis. APP expressed in all fetal tissues and is pronounced in brain, kidney, heart and spleen, but weak in liver. Defects in APP are the cause of Alzheimer disease type 1 (AD1). This antibody can recogniaze the N-terminus of human APP: Soluble APPalpha and Soluble APP-beta.

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