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beta 2 Adrenergic Receptor ELISA Kit



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Overview	
Quantity:	96 tests
Target:	beta 2 Adrenergic Receptor (ADRB2)
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	0.312 ng/mL - 20 ng/mL
Minimum Detection Limit:	0.312 ng/mL
Application:	ELISA
Product Details	
Purpose:	The kit is a sandwich enzyme immunoassay for the in vitro quantitative measurement of
	ADRb2 in human serum, plasma, tissue homogenates and other biological fluids.
Sample Type:	Plasma, Serum, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of this index.
Sensitivity:	0.114 ng/mL
Components:	Pre-coated, ready to use 96-well strip plate
	Standard (freeze dried)
	Standard Diluent
	Detection Reagent ADetection Reagent B

Product Details

- · Assay Diluent A
- · Assay Diluent B
- TMB
- Stop Solution
- Wash Buffer (30X)
- Plate sealer for 96 wells
- · Instruction manual

Material not included:

- 1. Microplate reader with 450 ± 10nm filter.
- 2. Precision single or multi-channel pipettes and disposable tips.
- 3. Eppendorf Tubes for diluting samples.
- 4. Deionized or distilled water.
- 5. Absorbent paper for blotting the microtiter plate.
- 6. Container for Wash Solution.

Target Details

Target:	beta 2 Adrenergic Receptor (ADRB2)
Alternative Name:	Adrenergic Receptor Beta 2 (ADRb2) (ADRB2 Products)
Background:	Alternative name: B2AR, Beta-2 adrenoreceptor
Gene ID:	154
UniProt:	P07550
Pathways:	cAMP Metabolic Process, Synaptic Membrane, Regulation of G-Protein Coupled Receptor
	Protein Signaling, Brown Fat Cell Differentiation

Application Details

Sample Volume:	100 μL
Assay Time:	1 - 4.5 h
Plate:	Pre-coated
Protocol:	1. Prepare all reagents, samples and standards
	2. Add 100µL standard or sample to each well. Incubate 2 hours at 37°C
	3. Aspirate and add 100µL prepared Detection Reagent A. Incubate 1 hour at 37°C
	4. Aspirate and wash 3 times
	5. Add 100µL prepared Detection Reagent B. Incubate 1 hour at 37°C
	6. Aspirate and wash 5 times

Application Details

	7. Add 90µL Substrate Solution. Incubate 15-25 minutes at 37°C
	8. Add 50µL Stop Solution. Read at 450nm immediately.
Assay Procedure:	The microtiter plate provided in this kit has been pre-coated with an antibody specific to the
	index. Standards or samples are then added to the appropriate microtiter plate wells with a
	biotin-conjugated antibody preparation specific to the index. Next, Avidin conjugated to
	Horseradish Peroxidase (HRP) is added to each microplate well and incubated. After TMB
	substrate solution is added, only those wells that contain the index, biotin-conjugated antibody
	and enzyme-conjugated Avidin will exhibit a change in color. The enzyme-substrate reaction is
	terminated by the addition of sulphuric acid solution and the color change is measured
	spectrophotometrically at a wavelength of 450nm \pm 10nm. The concentration of the index in
	the samples is then determined by comparing the O.D. of the samples to the standard curve.
Assay Precision:	 Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level the index were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level
	the index were tested on 3 different plates, 8 replicates in each plate.
	CV(%) = SD/meanX100Intra-assay: CV&lt10%
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	Inter-assay: CV<12%
Restrictions:	Inter-assay: CV<12% For Research Use only
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Handling	·
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Handling

provided to minimize exposure to damp air. The test kit may be used throughout the expiration date of the kit (six months from the date of manufacture). Opened test kits will remain stable until the expiring date shown, provided it is stored as prescribed above.

Expiry Date:

12 months