

Datasheet for ABIN5654920

GDA ELISA Kit



Overview

Quantity:	96 tests
Target:	GDA
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

Sample Type:	Plasma, Serum, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of Guanine Deaminase (GDA). No significant cross-reactivity or interference between Guanine Deaminase (GDA) and analogues was observed.
Sensitivity:	0.129 ng/mL

Target Details

Target:	GDA
Alternative Name:	Guanine Deaminase (GDA Products)

Target Details

Background:	Gene Name: Guanine Deaminase
	Gene Aliases: CYPIN, Guanase, Guanine aminase, Guanine aminohydrolase, p51-nedasin
Gene ID:	9615
UniProt:	Q9Y2T3
Application Details	
Comment:	The stability of kit is determined by the loss rate of activity. The loss rate of this kit is less than
	5 % within the expiration date under appropriate storage condition. To minimize extra influence
	on the performance, operation procedures and lab conditions, especially room temperature, ai
	humidity, incubator temperature should be strictly controlled. It is also strongly suggested that
	the whole assay is performed by the same operator from the beginning to the end.
Assay Time:	3 h
Plate:	Pre-coated
Protocol:	The test principle applied in this kit is Sandwich enzyme immunoassay. The microtiter plate
	provided in this kit has been pre-coated with an antibody specific to Guanine Deaminase (GDA
	Standards or samples are then added to the appropriate microtiter plate wells with a biotin-
	conjugated antibody specific to Guanine Deaminase (GDA). 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 Guanine Deaminase (GDA), 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
	Guanine Deaminase (GDA) 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
	Guanine Deaminase (GDA) were tested 20 times on one plate, respectively
	Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level
	Guanine Deaminase (GDA) were tested on 3 different plates, 8 replicates in each plate. CV(%) =
	SD/meanX100
	Intra-Assay: CV<10%
	Inter-Assay: CV<12%
Restrictions:	For Research Use only

Handling

Handling Advice:	The Stop Solution is acidic. Do not allow to contact skin or eyes. Calibrators, controls and
	specimen samples should be assayed in duplicate. Once the procedure has been started, all
	steps should be completed without interruption.
Storage:	4 °C,-20 °C
Storage Comment:	-20°C. Bring all reagents to room temperature before beginning test. The kit may be stored at 4°C for immediate use within two days upon arrival. Reseal any unused strips with desiccant pack. Minimize freeze/thaw cycles.
Expiry Date:	4-8 months