# antibodies - online.com





# Datasheet for ABIN4886389

# **Adalimumab ELISA Kit**



## Overview

Quantity:	96 tests
Target:	Adalimumab
Reactivity:	Human, Mouse, Rat
Method Type:	Sandwich ELISA
Detection Range:	1.56-50 ng/mL
Minimum Detection Limit:	1.56 ng/mL
Application:	ELISA

## **Product Details**

Purpose:	Quantification of Adalimumab in biological matrices
Sample Type:	Plasma, Serum
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	Adalimumab (Humira)
Cross-Reactivity (Details):	hlgG1, rituximab, and infliximab prepared at 250 ng/mL were assayed and exhibited no crossreactivity or interference.
Sensitivity:	1.5 ng/mL
Characteristics:	The Adalimumab ELISA kit is designed to measure free Adalimumab with high specificity and sensitivity. The assay design uses a pair of antibodies allowing detection of whole Adalimumab molecules in biological matrices

## **Product Details**

Components:	Coated microtiter plate, 96 wells
	Calibrator diluent 1.8ml
	Calibrator 12ul
	10X wash buffer - 25ml
	Assay buffer - 50ml
	1000X detection reagent - 17ul
	TMB - 12ml
	TMB stop solution - 12ml
	Plate sealers - 3
Material not included:	Precision pipettes calibrated to deliver 5-1000µL
	Multi-channel pipette calibrated to deliver 50-200μL
	Plate shaker
	Disposable tips
	Vortex-Mixer
	Distilled or de-ionized water
	Microplate reader capable of reading 450nm with background subtrac

# **Target Details**

Target:	Adalimumab
Abstract:	Adalimumab Products
Target Type:	Antibody
Background:	Adalimumab (Trade name Humira®) is a human monoclonal antibody used to block the action of tumor necrosis factor alpha (TNF-). TNF- $\alpha$ is a strong proinflammatory cytokine involved in normal immunity and also various autoimmune diseases.
Gene ID:	7124

## **Application Details**

Application Notes:	Optimal working dilution should be determined by the investigator.
Sample Volume:	15 µL
Assay Time:	2.5 h
Plate:	Pre-coated
Protocol:	The Adalimumab ELISA kit is designed to measure free Adalimumab with high specificity and

sensitivity. This assay employs the sandwich enzyme immunoassay technique. A precoated anti-Adalimumab 96 well plate is provided. Calibrator, quality control samples and test samples are pipetted into the appropriate wells. Adalimumab present in biological matrices is bound by the immobilized capture antibody. After washing away any unbound substances, enzyme linked detection antibody is added to the wells. The plate is washed to remove any unbound antibody-enzyme reagent and a substrate solution is added to the wells for color development. The color development is proportional to the amount of Adaliumumab present in test samples and the concentration is calculated from the standard series.

#### Reagent Preparation:

Prepare only the appropriate amount of required reagent on the day of use. Store all reagents as per instructions stated on the label. 1. Wash Buffer (1X) Preparation: Dilute wash buffer concentrate with ultra-pure water 1/10 before use (for example add 20 mL concentrate to 180 mL ultra-pure water). Mix well. 2. Detection Reagent (1X) Preparation: Dilute detection reagent with assay buffer 1/1000 before use (for example add 11  $\mu$ L concentrate to 11 mL of assay buffer). Mix well. 3. Preparation of Calibrators: Prepare calibrators with concentrations ranging from 2,500 ng/mL to 78 ng/mL. The following is an example calibrator curve.

#### Sample Collection:

This kit is compatible with EDTA-plasma, heparinplasma and serum samples. Samples can be stored at or below -20 °C for up to 1 year.

#### Sample Preparation:

Dilute calibrators and test samples 1/50 with assay buffer (for example add 5µL of prepared calibrator or sample to 245µL of assay buffer). Mix well. Do not store diluted samples.

### Assay Procedure:

This assay employs the sandwich enzyme immunoassay technique. Anti- Adalimumab is coated onto a 96 well microplate. Calibrator, quality control samples and test samples are pipetted into the appropriate wells. Adalimumab present in biological matrices is bound by the immobilized capture antibody. After washing away any unbound substances, enzyme linked detection antibody is added to the wells. The plate is washed to remove any unbound antibody-enzyme reagent and a substrate solution is added to the wells for color development. The color development is proportional to the amount of Adalimumab present in test samples.

#### Calculation of Results:

1. Construct a standard curve by plotting the absorbance obtained from each standard against concentration. Use a 4 or 5 parameter curve fit. Alternatively a log-log curve fit may be used. 2. The concentration of the unknowns can be read directly from this standard curve using the absorbance value for each sample. 3. Any sample undiluted or diluted still reading greater than the highest standard should be diluted appropriately with calibrator diluent and retested. If the samples have been diluted, the concentration determined from the standard curve must be multiplied by the dilution factor.

#### Assay Precision:

Precision: The precision was determined by analyzing samples prepared at 313 ng/mL in 6

# **Application Details**

Expiry Date:

	replicates on 6 different occasions. Intra-assay coefficient of variation (CV) $<$ 10%. Inter-assay CV $<$ 10%.
	Recovery: 250 ng/mL of Adalimumab was spiked in 10 lots of human serum. Recovery ranges are from 90-109% with an average recovery of 104%.
Restrictions:	For Research Use only
Handling	
Preservative:	Without preservative
Precaution of Use:	Read manual completely before beginning
Storage:	-20 °C
Storage Comment:	Store kit components at -20°C unless specified otherwise. DO NOT USE past kit expiration date. Some vials contain a small amount of reagents. Spin tubes on pulse setting prior to opening.

12 months