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Adalimumab Antibody ELISA Kit



Images



Overview

Quantity:	96 tests
Target:	Adalimumab Antibody
Binding Specificity:	Free Chain
Reactivity:	Chemical, Human
Method Type:	Sandwich ELISA
Application:	ELISA
Product Details	
Purpose:	Enzyme immunoassay for the semi-quantitative determination of free antibodies to
	Adalimumab in human serum and plasma samples.
Brand:	ImmunoGuide®
Sample Type:	Serum, Plasma (EDTA - heparin)
Analytical Method:	Semi-Quantitative
Detection Method:	Colorimetric
Specificity:	Free antibodies against Adalimumab (Humira®)S creening test was performed with 90
	different native and RF-negative human sera and additionally 2 different ATI positive sera. All
	produced OD450nm values (ranged from 0.053 to 0.113) less than the cut-off value (3x0.086).
Sensitivity:	10 ng/mL
Characteristics:	This test does not measure the antibodies if they already are bound to the drug Adalimumab.
Components:	• 1 x 12 x 8 Microtiter Plate Break apart strips pre-coated with the drug Adalimumab.

- 1 x 2 mL Positive Control Ready to use. Contains reactive antibody, proteins, stabilizer and
 15 mM NaN3.
- 1 x 2 mL Negative Control Ready to use. Contains human serum, stabilizer and <15 mM NaN3.
- 1 x 60 mL Assay Buffer Blue colored. Ready to use. Contains proteins and <15 mM NaN3.
- 1 x 12 mL Enzyme Conjugate Red colored. Ready to use. Contains horseradish peroxidase(HRP)-conjugated Adalimumab, Proclin® and stabilizers.
- 1 x 12 mL TMB Substrate Solution Ready to use. Contains 3,3',5,5'-Tetramethylbenzidine (TMB).
- 1 x 12 mL Stop Solution Ready to use. 1 N Hydrochloric acid (HCl).
- 1 x 50 mL Wash Buffer, Concentrate (20x) Contains buffer, Tween® 20 and KathonTM.
- 2 x 1 Adhesive Seal For sealing microtiter plate during incubation.

Material not included:

- Micropipettes (< 3 % CV) and tips to deliver 5-1000 μL.
- Bidistilled or deionised water and calibrated glasswares (e.g. flasks or cylinders).
- · Wash bottle, automated or semi-automated microtiter plate washing system.
- Microtiter plate reader capable of reading absorbance at 450 nm (reference wavelength at 600-650 nm is optional).
- Absorbent paper towels, standard laboratory glass or plastic vials, and a timer.

Target Details

Target:	Adalimumab Antibody
Target Type:	Antibody
Background:	Adalimumab is a tumor necrosis factor (TNF) blocker indicated for treatment of Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, Ankylosing Spondylitis, Crohn's Disease, Plaque Psoriasis. Accoring to the prescribing information, the use of Adalimumab was associated to the development of anti- Adalimumab antibodies, even some were reported to be neutralizing, in various percentages of patients during therapy with the drug Adalimumab. This might lead to severe complications. The Antibody to Adalimumab ELISA Kit can be efficiently used for monitoring anti- Adalimumab antibodies during therapy and offers the clinician a tool
	for decision on possible preventive measures to reduce anti-Adalimumab antibodies.
Molecular Weight:	144 kDa
Application Details	
Application Notes:	 Before performing the assay, samples and assay kit should be brought to room temperature (about 30 minutes beforehand) and ensure the homogeneity of the solution. All Standards should be run with each series of unknown samples.

· Standards should be subject to the same manipulations and incubation times as the samples being tested. • All steps of the test should be completed without interruption. · Use new disposable plastic pipette tips for each reagent, standard or specimen in order to avoid cross contamination. Comment: Antibody to Adalimumab ELISA is suitable also for using by an automated ELISA processor. Sample Volume: 5 μL Assay Time: 25h Plate: Pre-coated Protocol: The Antibody to Adalimumab ELISA is a sandwich type ELISA for the determination of free antibodies against Adalimumab in serum and plasma samples. During the first incubation period, antibodies to Adalimumab in patient serum/plasma samples are captured by the drug Adalimumab coated on the wall of the microtiter wells. After washing away the unbound components from samples, a peroxidase-labelled Adalimumab conjugate is added and then incubated. Antibody to Adalimumab, if present in sample, will make a bridge, with its two identical Fab arms, between the Adalimumab coated on the well and the other Adalimumab labeled with peroxidase. After a second washing step, the bound enzymatic activity is detected by addition of tetramethylbenzidine (TMB) chromogen-substrate. Finally, the reaction is terminated with stop solution. The positive reaction is related to the presence of antibodies to Adalimumab in the sample. Reagent Preparation: Wash Buffer: Dilute 10 mL Wash Buffer (up to 200 mL) at the ratio of 1:20 with distilled water. Warm up at 37 °C to dissolve crystals. Mix vigorously. Store at 2-8 °C for up to 4 weeks. Prepare Wash Buffer before starting the assay procedure. Sample Collection: Normal serum or plasma collection Sample Preparation: Serum/ Plasma: Dilute Serum/ Plasma (Sample) at the ratio of 1:101 with Assay Buffer. For dilution at 1:101, 5 µL Sample + 500 µL Assay Buffer Negative and Positive Controls are ready-to-use and should NOT be diluted with the assay buffer. Serum, Plasma (EDTA, Heparin): The usual precautions for venipuncture should be observed. It is important to preserve the chemical integrity of a blood specimen from the moment it is collected until it is assayed. Do not use grossly hemolytic, icteric or grossly lipemic specimens.

Samples appearing turbid should be centrifuged before testing to remove any particulate material.

Storage: 2-8 °C &leq,-20 °C (Aliquots)

Keep away from heat or direct sun light.

Avoid repeated freeze-thaw cycles.

Stability: 3 days at 2-8 °C, 6 months at -20 °C

Assay Procedure:

- 1. Pipette 100 µL of each Ready-to Use Negative Control, Positive Control, and 1:101 Diluted Samples (as described in section 10.2) into the respective wells of microtiter plate. Wells A1: Negative Control B1: Negative Control C1: Positive Control D1 and so on: Diluted samples (Serum/Plasma)
- 2. Cover the plate with adhesive seal. Shake plate carefully. Incubate 60 min at room temperature (RT) (18-25 °C).
- 3. Remove adhesive seal. Aspirate or decant the incubation solution. Wash the plate 3 \times 300 μ L of Diluted Wash Buffer per well. Remove excess solution by tapping the inverted plate on a paper towel.
- 4. Pipette 100 μL of Enzyme Conjugate (HRP-Adalimumab) into each well.
- 5. Cover plate with adhesive seal. Shake plate carefully. Incubate 60 min at RT.
- 6. Remove adhesive seal. Aspirate or decant the incubation solution. Wash the plate 3 \times 300 μ L of Diluted Wash Buffer per well. Remove excess solution by tapping the inverted plate on a paper towel.
- 7. Pipette 100 µL of Ready-to-Use TMB Substrate Solution into each well.
- 8. Incubate 20 min at RT. Avoid exposure to direct sunlight...
- 9. Stop the substrate reaction by adding 100 μ L of Stop Solution into each well. Briefly mix contents by gently shaking the plate. Color changes from blue to yellow.
- 10. Measure optical density (OD) with a photometer at 450 nm (Reference at OD620nm is optional) within 15 min after pipetting the Stop Solution.

Calculation of Results:

For the run to be valid, the OD450 nm of the Positive Control should be \geq 1.000 and the OD450 nm of each Negative Control should be \leq 0.150. If not, improper technique or reagent deterioration may be suspected and the run should be repeated. The results are evaluated by dividing all individual results by the mean OD450nm of the Negative Controls. The results are expressed in arbitrary units (AU/mL).

Range: > 3 AU/mL (Positive)

Range: &leq, 3 AU/mL (Negative)

The results themselves should not be the only reason for any therapeutical consequences.

They have to be correlated to other clinical observations.

	Sample calculation for a positive sample:
	OD of sample = 0.740 Cut-off = 3×10^{-2}
	Average OD of Negative Controls = 3 x 0.100 = 0.300 = 3 AU/mL
	Concentration of sample = 0.740/0.100 = 7.4 AU/mL (positive)
Assay Precision:	Intra-assay CV: <10%.
	Inter-assay CV: <10%
Restrictions:	For Research Use only
Handling	
Buffer:	< 15mM NaN3
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
	should be handled by trained staff only.
Storage:	4 °C
Storage Comment:	The kit is shipped at ambient temperature and should be stored at 2-8°C.
	Keep away from heat or direct sun light.
	The storage and stability of specimen and prepared reagents is stated in the corresponding
	chapters.
	The microtiter strips are stable up to the expiry date of the kit in the broken, but tightly closed
	bag when stored at 2-8°C.
Expiry Date:	24 months

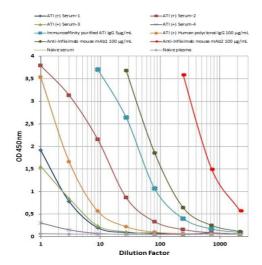


Image 1.

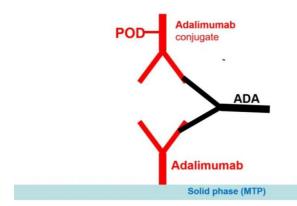


Image 2.

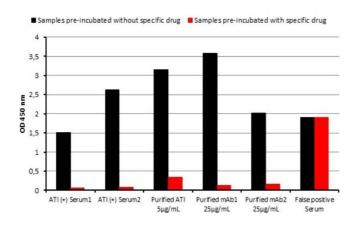


Image 3.