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Datasheet for ABIN7041579 Hemopexin ELISA Kit

Image



Overview

Quantity:	96 tests
Target:	Hemopexin (HPX)
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	6.25 ng/mL - 200 ng/mL
Minimum Detection Limit:	6.25 ng/mL
Application:	ELISA

Product Details

Purpose:	The kit is a sandwich enzyme immunoassay for the in vitro quantitative measurement of hemopexin in human serum, plasma.
Sample Type:	Plasma, Serum, Urine
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Components:	 ELISA Micro Plate antibody coated Enzyme Conjugated Detection Antibody (100X) Calibrator Diluent Concentrate (5X) Wash Solution Concentrate (20X) Chromogen - Substrate Solution STOP Solution

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Material not included:

- 1. Microplate reader with 450 ± 10 nm filter.
- 2. Precision single or multi-channel pipettes and disposable tips.
- 3. Microcentrifuge tubes for diluting samples.
- 4. Squirt bottle or Microtitre washer
- 5. Deionized or distilled water.
- 6. Container for Wash Solution
- 7. Centrifuge for sample collection
- 8. Anticoagulant for plasma collection
- 9. Incubator capable of maintaining 37 °C.
- 10. Microplate shaker

Target Details

Target:	Hemopexin (HPX)
Alternative Name:	Hemopexin (HPX Products)
Pathways:	Transition Metal Ion Homeostasis, Regulation of Leukocyte Mediated Immunity, Positive Regulation of Immune Effector Process, Production of Molecular Mediator of Immune
	Response

Application Details

Sample Volume:	100 µL
Assay Time:	1.5 h
Plate:	Pre-coated
Protocol:	In this assay the hemopexin (HX) present in samples reacts with the anti-HX antibodies, which
	have been adsorbed to the surface of polystyrene microtitre wells. After the removal of
	unbound proteins by washing, anti-HX antibodies conjugated with horseradish peroxidase
	(HRP) are added. These enzyme-labeled antibodies form complexes with the previously bound
	HX. Following another washing step, the enzyme bound to the immunosorbent is assayed by
	the addition of a chromogenic substrate, 3,3',5,5'-tetramethylbenzidine (TMB). The quantity of
	bound enzyme varies directly with the concentration of HX in the sample tested; thus, the
	absorbance, at 450 nm, is a measure of the concentration of HX in the test sample. The
	quantity of HX in the test sample can be interpolated from the standard curve constructed from
	the standards, and corrected for sample dilution.
Reagent Preparation:	 Bring all reagents to room temperature (16°C to 25°C) before use. Diluent Concentrate - The Diluent Solution supplied is a 5X Concentrate and must be diluted

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	 1/5 with distilled or deionized water (1 part buffer concentrate, 4 parts dH2O). Wash Solution Concentrate - The Wash Solution supplied is a 20X Concentrate and must be diluted 1/20 with distilled or deionized water (1 part buffer concentrate, 19 parts dH2O). Crystal formation in the concentrate may occur when storage temperatures are low. Warming of the concentrate to 30-35°Cbefore dilution can dissolve crystals. Enzyme-Antibody Conjugate - Calculate the required amount of working conjugate solution for each microtitre plate test strip by adding 10 µL Enzyme-Antibody Conjugate to 990 µL of 1X Diluent for each test strip to be used for testing. Dilute immediately before use and protect from light. Mix uniformly, but gently. Avoid foaming. Pre-coated ELISA Micro Plate - Ready to use as supplied. Unseal foil pouch and remove plate from pouch. Remove all strips and wells that will not be used in the assay and place back in pouch and re-seal along with desiccant. Calibrator – Prepare according to the lot specific Certificate of Analysis.
Sample Preparation:	 It is recommended to use fresh samples without long storage, otherwise protein degradation and denaturationmay occur in these samples, leading to false results. Samples should therefore be stored for a short periodat 2 - 8 °C or aliquoted at -20 °C (≤1 month) or -80 °C (≤ 3 months). Repeated freeze-thawcycles should be avoided. Prior to assay, the frozen samples should be slowly thawed and centrifuged toremove precipitates. If the sample type is not specified in the instructions, a preliminary test is necessary to determinecompatibility with the kit. If a lysis buffer is used to prepare tissue homogenates or cell culture supernatant, there is a possibility causing a deviation due to the introduced chemical substance. The recommended dilution factor is for reference only. Please estimate the concentration of the samples before performing the test. If the values are not in therange of the standard curve, the optimal sample dilution for the particular experiment has to be determined. Samples should then be diluted with PBS (pH =7.0-7.2).
	Note:
	 The assay requires that each test sample be diluted before use. All samples should be assayed in duplicate each time the assay is performed. The recommended dilutions are only suggestions. Dilutions should be based on the expected concentration of the unknown sample such that the diluted sample falls within the dynamic range of the standard curve. If unsure of sample level, a serial dilution with one or two representative samples before running the entire plate is highly recommended. Serum samples – Recommended starting dilution is 1/40,000. To prepare a 1/40,000 dilution of a sample, transfer 5 µL of sample to 495 µL of 1X diluent. This gives you a 1/100 dilution. Next, dilute the 1/100 by transferring 2 µL into 798 µL of 1X diluent. This gives you a 1/40,000 dilution of a samples – Recommended starting dilution is 1/40,000. To prepare a 1/40,000 dilution of dilution. Mix thoroughly each stage. Plasma samples – Recommended starting dilution is 1/40,000. To prepare a 1/40,000 dilution of a sample, transfer 5 µL of sample to 495 µL of 1X diluent. This gives you a 1/100 dilution.

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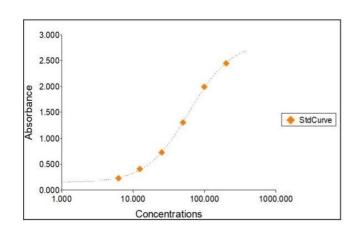
	Next, dilute the 1/100 by transferring 2 μL into 798 μL of 1X diluent. This gives you a 1/40,000
	dilution. Mix thoroughly each stage.
Assay Procedure:	1. All samples and standards should be assayed in duplicates.
	2. The Standards and the test sample(s) should be loaded into the ELISA wells as quickly as
	possible to avoid a shift in OD readings. Using a multichannel pipette would reduce this
	occurrence. Pipette 100 μL of Standard 0 (0.0 ng/mL) in duplicate Standard 1 (6.25 ng/mL) in
	duplicate Standard 2 (12.50 ng/mL) in duplicate Standard 3 (25 ng/mL) in duplicate Standard
	(50 ng/mL) in duplicate Standard 5 (100 ng/mL) in duplicate Standard 6 (200 ng/mL) in
	duplicate
	3. Pipette 100 μ L of sample (in duplicate) into pre designated wells.
	4. Incubate the micro titer plate at room temperature for thirty (30 \pm 2) minutes. Keep plate
	covered and level during incubation.
	5. Following incubation, aspirate the contents of the wells.
	6. Completely fill each well with appropriately diluted Wash Solution and aspirate. Repeat three
	times, for a total of four washes. If washing manually: completely fill wells with wash buffer,
	invert the plate then pour/shake out the contents in a waste container. Follow this by sharply
	striking the wells on absorbent paper to remove residual buffer. Repeat 3 times for a total of
	four washes.
	7. Pipette 100 μ L of appropriately diluted Enzyme-Antibody Conjugate to each well. Incubate a
	room temperature for thirty (30 \pm 2) minutes. Keep plate covered in the dark and level during
	incubation.
	8. Wash and blot the wells as described in Steps 5/6.
	9. Pipette 100 μ L of TMB Substrate Solution into each well.
	10. Incubate in the dark at room temperature for precisely ten (10) minutes.
	11. After ten minutes, add 100 μL of Stop Solution to each well.
	12. Determine the absorbance (450 nm) of the contents of each well within 30 minutes.
	Calibrate the plate reader to manufacturer's specifications.
Restrictions:	For Research Use only
Handling	
Storage:	4 °C
Storage Comment:	1. For unopened kit: All reagents should be stored according to the labels on the vials. All reagents should be stored at 4 °C for long term storage. Keep away from heat or direct sunlight.

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Expiry Date:

6 months

Images



ELISA

Image 1. Standard Curve Graph