.-online.com antibodies

Datasheet for ABIN433206 Sortilin 1 ELISA Kit



Overview

Quantity:	96 tests
Target:	Sortilin 1 (SORT1)
Reactivity:	Rat
Method Type:	Sandwich ELISA
Detection Range:	0.15 ng/mL - 10 ng/mL
Minimum Detection Limit:	0.15 ng/mL
Application:	ELISA

Product Details

Sample Type:	Plasma, Serum
Detection Method:	Colorimetric
Sensitivity:	0.056 ng/mL
Material not included:	 Microplate reader with 450nm filter. Precision single or multi-channel pipettes and disposable tips. Eppendorf Tubes for diluting samples. Deionized or distilled water. Absorbent paper for blotting the microtiter plate. Container for Wash Solution

Target Details

Target:

Sortilin 1 (SORT1)

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/6 | Product datasheet for ABIN433206 | 09/12/2023 | Copyright antibodies-online. All rights reserved.

Target Details

Products
rophin Signaling Pathway
ed by the current condition and scientific technology, we cannot completely conduct the prehensive identification and analysis on the raw material provided by suppliers. So emight be some qualitative and technical risks to use the kit. final experimental results will be closely related to validity of the products, operation of the end users and the experimental environments. Please make sure that sufficient poles are available. from different batches may be a little different in detection range, sensitivity and color loping time. ot mix or substitute reagents from one kit lot to another. Use only the reagents supplied anufacturer. act all reagents from strong light during storage and incubation. All the bottle caps of ents should be covered tightly to prevent the evaporation and contamination of poorganism. e may be some foggy substance in the wells when the plate is opened at the first time. It to thave any effect on the final assay results. Do not remove microtiter plate from the ige bag until needed. go operations during the reagents preparation and loading, as well as incorrect meter setting for the plate reader may lead to incorrect results. A microplate plate reader a bandwidth of 10mm or less and an optical density range of 0-3 0.D. or greater at 450 ± n wavelength is acceptable for use in absorbance measurement. Please read the action carefully and adjust the instrument prior to the experiment. the same operator might get different results in two separate experiments. In order to etter reproducible results, the operation of every step in the assay should be controlled. kit has been strictly passed Q.C test. However, results from end users might be asistent with our in-house data due to some unexpected transportation conditions or rent lab equipments. Intra-assay variance among kits from different batches might arise above factors, too. from different manufacturers for the same item might produce different results, since ave not compared our products with other manufacturers.
ation on standard material: Indard might be recombinant protein or natural protein, that will depend on the specific
reover, the expression system is E.coli or yeast or mammal cell. There is 0.05% proclin the standard as preservative.
f na a

Information on reagents:

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 2/6 | Product datasheet for ABIN433206 | 09/12/2023 | Copyright antibodies-online. All rights reserved.

	The stop solution used in the kit is sulfuric acid with concentration of 1 mol/L. And the wash
	solution is TBS. The standard diluent contains 0.02 % sodium azide, assay diluent A and assay
	diluent B contain 0.01% sodium azide. Some kits can contain is BSA in them.
	Information on antibodies:
	The provided antibodies and their host vary in different kits.
Sample Volume:	100 µL
Plate:	Pre-coated
Sample Collection:	Serum: Allow samples to clot for two hours at room temperature or overnight at 4°C before
	centrifugation for 20 minutes at approximately 1000 × g. Assay immediately or store samples
	in aliquot at -20°C or -80°C. Avoid repeated freeze/thaw cycles.
	Plasma: Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge samples for 15
	minutes at 1000 × g within 30 minutes of collection. Remove plasma and assay immediately or
	store samples in aliquot at -20°C or -80°C. Avoid repeated freeze/thaw cycles.
	Biological Fluids: Centrifuge samples for 20 minutes at 1000 × g. Remove particulates and
	assay immediately or store samples in aliquot at -20 °C or -80 °C for later use. Avoid repeated
	freeze/thaw cycles.
Sample Preparation:	Notes:
	• The user should calculate the possible amount of the samples used in the whole test. Please reserve sufficient samples in advance.
	• Please predict the concentration before assaying. If values for these are not within the range
	of the standard curve, users must determine the optimal sample dilutions for their particular experiments.
	• If the samples are not indicated in the manual, a preliminary experiment to determine the
	validity of the kit is necessary.
	 Tissue or cell extraction samples prepared by chemical lysis buffer may cause unexpected ELISA results due to the impacts from certain chemicals.
	 Due to the possibility of mismatching between antigen from other origin and antibody used in
	our kits (e.g., antibody targets conformational epitope rather than linear epitope), some native
	or recombinant proteins from other manufacturers may not be recognized by our products.
	Influenced by the factors including cell viability, cell number or sampling time, samples from
	cell culture supernatant may not be detected by the kit.
	Fresh samples without long time storage is recommended for the test. Otherwise, protein
	degradation and denaturalization may occur in those samples and finally lead to wrong

Assay Procedure:

- 1. Determine wells for diluted standard, blank and sample. Prepare wells for standard points, 1 well for blank. Add dilutions of standard (read Reagent Preparation), blank and samples into the appropriate wells, respectively. You might now need to incubate at 37 °C. And then add Detection Reagent A to each well. You might need to incubate at 37 °C again.
 - 2. Aspirate the solution and wash with 350 µL of 1X Wash Solution to each well using a squirt bottle, multi-channel pipette, manifold dispenser or autowasher, and let it sit for 1-2 minutes. Remove the remaining liquid from all wells completely by snapping the plate onto absorbent paper. Repeat 3 times. After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against absorbent paper.
 - 3. Add 100 μL of Detection Reagent B working solution to each well. Incubate for 30 minutes at 37 °C after covering it with the Plate sealer.
- 4. Repeat the aspiration/wash process for total 5 times as conducted in step 2.
- 5. Add 90 µL of Substrate Solution to each well. Cover with a new Plate sealer. Incubate for 15 - 25 minutes at 37 °C (Don't exceed 30 minutes). Protect from light. The liquid will turn blue by the addition of Substrate Solution.
- 6. Add 50 µL of Stop Solution to each well. The liquid will turn yellow by the addition of Stop solution. Mix the liquid by tapping the side of the plate. If color change does not appear uniform, gently tap the plate to ensure thorough mixing.
- 7. Remove any drop of water and fingerprint on the bottom of the plate and confirm there is no bubble on the surface of the liquid. Then, run the microplate reader and conduct measurement at 450 nm immediately.

Note:

- 1. Assay preparation: Keep appropriate numbers of wells for each experiment and remove extra wells from microplate. Rest wells should be resealed and stored at -20 °C.
 2. Samples or reagents addition: Please use the freshly prepared Standard. Please carefully add samples to wells and mix gently to avoid foaming. Do not touch the well wall. For each step in the procedure, total dispensing time for addition of reagents or samples to the assay plate should not exceed 10 minutes. This will ensure equal elapsed time for each pipetting step, without interruption. To avoid cross-contamination, change pipette tips between additions of standards, samples, and reagents. Also, use separated reservoirs for each reagent.
- 3. Incubation: To ensure accurate results, proper adhesion of plate sealers during incubation steps is necessary. Do not allow wells to sit uncovered for extended periods between incubation steps. Once reagents are added to the well strips, DO NOT let the strips DRY at any time during the assay. Incubation time and temperature must be controlled.
- 4. Washing: The wash procedure is critical. Complete removal of liquid at each step is essential for good performance. After the last wash, remove any remaining Wash Solution by aspirating or decanting and remove any drop of water and fingerprint on the bottom of the plate. Insufficient washing will result in poor precision and false elevated absorbance reading.
- 5. Controlling of reaction time: Observe the change of color after adding TMB Substrate (e.g. observation once every 10 minutes), if the color is too deep, add Stop Solution in advance to avoid excessively strong reaction which will result in inaccurate absorbance reading.

Application Details	
	 6. TMB Substrate is easily contaminated. Please protect it from light. 7. The environment humidity which is less than 60 % might have some effects on the final performance, therefore, a humidifier is recommended to be used at that condition.
Calculation of Results:	Average the duplicate readings for each standard, control, and samples and subtract the
	average zero standard optical density. Construct a standard curve by plotting the mean O.D.
	and concentration for each standard and draw a best fit curve through the points on the graph
	or create a standard curve on log-log graph paper with the target concentration on the y-axis
	and absorbance on the x-axis. Draw the best fit straight line through the standard points and it
	can be determined by regression analysis.Using some plot software, for instance, curve expert
	1.30, is also recommended. If samples have been diluted, the concentration read from the
	standard curve must be multiplied by the dilution factor.
Assay Precision:	Intra-assay Precision (precision within an assay): Three samples with low, medium and high
	levels of the target antigen were tested twenty times on one plate, respectively.
	Inter-assay Precision (precision between assays): Three samples with low, medium and high
	levels of the target antigen were tested on three different plates, eight replicates in each plate.
	CV (%) = SD/mean X 100
	 Intra-assay: CV less than 10 %
	Inter-assay: CV less than 12 %
Restrictions:	For Research Use only
Handling	
Precaution of Use:	The Stop Solution suggested for use with this kit is an acid solution. Wear eye, hand, face, and
	clothing protection when using this material.
Handling Advice:	To minimize extra influence on the performance, operation procedures and lab conditions,
	especially room temperature, air 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.
Storage:	4 °C/-20 °C
Storage Comment:	• For unopened kit: All the reagents should be kept according to the labels on vials. The
	Standard, Detection Reagent A, Detection Reagent B and the 96-well strip plate should be stored at -20° C upon receipt while the others should be at 4° C
	 stored at -20°C upon receipt while the others should be at 4°C. For opened kit: When the kit is opened, the remaining reagents still need to be stored
	according to the above storage condition. Besides, please return the unused wells to the foil
	pouch containing the desiccant pack, and reseal along entire edge of zip-seal.

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 5/6 | Product datasheet for ABIN433206 | 09/12/2023 | Copyright antibodies-online. All rights reserved. Note: It is highly recommended to use the remaining reagents within 1 month provided this is within the expiration date of the kit.

• For ELISA kit, 1 day storage at 37°C can be considered as 2 months at 4°C, which means 3 days at 37°C equaling 6 months at 4°C.

Expiry Date:

6 months