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Datasheet for ABIN2747914 WNK1 ELISA Kit



Overview

Quantity:	96 tests
Target:	WNK1
Binding Specificity:	pThr60, total
Reactivity:	Human
Method Type:	Sandwich ELISA
Application:	ELISA

Product Details

Purpose:	Human Phospho-WNK1 (T60) & Total WNK1 ELISA Kit. This assay semi-quantitatively measures phophorylated WNK1 (Thr60) & Total WNK1 in lysate samples.
Sample Type:	Cell Lysate, Tissue Lysate
Analytical Method:	Semi-Quantitative
Detection Method:	Colorimetric
Specificity:	The antibody pair provided in this kit recognizes human WNK1 phosphorylated at site Threonine-60 and total WNK1.
Characteristics:	 Simultaneously measure Phosphorylated protein and pan protein in one experiment (for normalization purpose) Screen numerous different cell lysates without performing a Western Blot analysis Minimal hands-on time, convenient, and non-radioactive material
Components:	Pre-Coated 96-well Strip MicroplateWash Buffer

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	 Anti-Phospho Antibody Anti-Pan Antibody HRP-Conjugated Secondary Antibody Streptavidin-Conjugated HRP Assay Diluent TMB One-Step Substrate Stop Solution Lysis Buffer Positive Control Sample
Material not included:	 Distilled or deionized water 100 mL and 1 liter graduated cylinders Tubes to prepare sample dilutions Protease and Phosphatase inhibitors Precision pipettes to deliver 2 µL to 1 mL volumes Adjustable 1-25 mL pipettes for reagent preparation Benchtop rocker or shaker Microplate reader capable of measuring absorbance at 450 nm

Target Details

Target:	WNK1
Alternative Name:	WNK1 (WNK1 Products)
Background:	WNK Lysine-deficient Protein Kinase 1 (WNK1, KDP, PHA2C, PRKWNK1) phosphorylated at Threonine-60 & total WNK1
Gene ID:	65125
UniProt:	Q9H4A3

Application Details

Sample Volume:	100 μL
Plate:	Pre-coated
Protocol:	1. Prepare all reagents and samples as instructed in the manual.
	2. Add 100 µL of sample or positive control to each well.
	3. Incubate 2.5 h at RT or O/N at 4 °C.
	4. Add 100 µL of prepared primary antibody to each well.
	5. Incubate 1 h at RT.
	6. Add 100 μL of prepared 1X HRP-Streptavidin to each well.

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	 7. Incubate 1 h at RT. 8. Add 100 μL of TMB One-Step Substrate Reagent to each well. 9. Incubate 30 min at RT. 10. Add 50 μL of Stop Solution to each well. 11. Read at 450 nm immediately.
Assay Procedure:	Prepare all reagents and samples as instructed in the manual.
	Add 100 μ L of sample or positive control to each well.
	Incubate 2.5 h at RT or O/N at 4 °C.
	Add 100 μ L of prepared primary antibody to each well.
	Incubate 1 h at RT.
	Add 100 μ L of prepared 1X HRP-Streptavidin to each well.
	Incubate 1 h at RT.
	Add 100 μ L of TMB One-Step Substrate Reagent to each well.
	Incubate 30 min at RT.
	Add 50 μ L of Stop Solution to each well.
	Read at 450 nm immediately.
Restrictions:	For Research Use only

Handling

Storage:	-20 °C
Storage Comment:	Upon receipt, the kit should be stored at -20 °C. Please use within 6 months from the date of
	shipment. After initial use, Wash Buffer Concentrate (Item B), Assay Diluent (Item E), TMB One-
	Step Substrate Reagent (Item H), HRP-Streptavidin (Item G), Stop Solution (Item I) and Cell
	Lysate Buffer (Item J) should be stored at 4 °C to avoid repeated freeze-thaw cycles. Return
	unused wells to the pouch containing desiccant pack, reseal along entire edge and store at -20
	°C. Reconstituted Positive Control (Item K) should be stored at -70 °C.
Expiry Date:	6 months