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Datasheet for ABIN2748556 PRAS40 ELISA Kit



Overview

Quantity:	96 tests
Target:	PRAS40 (AKT1S1)
Binding Specificity:	pThr246
Reactivity:	Human
Method Type:	Sandwich ELISA
Application:	ELISA
Product Details	
Purpose:	Human Phospho-PRAS40 (T246) ELISA Kit. This assay semi-quantitatively measures phophorylated PRAS40 (Thr246) 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 PRAS40 phosphorylated at site Threonine-246
Characteristics:	 Rapidly measure phosphorylated protein in lysates 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 Microplate Wash Buffer Anti-Phospho Antibody

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	 HRP-Conjugated Secondary Antibody 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:	PRAS40 (AKT1S1)
Alternative Name:	PRAS40 (AKT1S1 Products)
Background:	40 kDa Proline-rich Akt1 Substrate (PRAS40 / AKT1S1 / Lobe) phosphorylated at Threonine-246
Gene ID:	84335
UniProt:	Q96B36
Pathways:	Fc-epsilon Receptor Signaling Pathway, EGFR Signaling Pathway, Neurotrophin Signaling
	Pathway, Regulation of Cell Size, Autophagy, BCR Signaling, Warburg Effect

Application Details

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

Application Details	
	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 µL of TMB One-Step Substrate Reagent to each well.
	Incubate 30 min at RT.
	Add 50 µL 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

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