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Datasheet for ABIN4955901 Human/Mouse MAPK Phosphorylation Array

1	Image	1	Publication
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Overview

Quantity:	4 samples				
Reactivity:	Human, Mouse				
Method Type:	Sandwich ELISA				
Application:	Antibody Array (AA)				
Product Details					
Purpose:	C-Series Human/Mouse MAPK Phosphorylation Antibody Array 1 Kit. Detects 17				
	Human/mouse proteins. Suitable for all liquid sample types but intended for use with cell and				
	tissue lysates.				
Sample Type:	Plasma, Cell Culture Supernatant, Serum, Cell Lysate, Tissue Lysate				
Analytical Method:	Semi-Quantitative				
Detection Method:	Chemiluminescent				
Specificity:	Akt (P-Ser473), CREB (P-Ser133), ERK1 (P-T202/Y204)/ERK2 (P-Y185/Y187), GSK3a (P-Ser21),				
	GSK3b (P-Ser9), HSP27 (P-Ser82), JNK (P-Thr183), MEK (P-Ser217/Ser221), MKK3 (P-Ser189),				
	MKK6 (P-Ser207), MSK2 (P-Ser360), mTOR (P-Ser2448), p38 (P-Thr180/Tyr182), P53 (P-Ser15),				
	P70S6K (P-Thr421/Ser424), RSK1 (P-Ser380), RSK2 (P-Ser386)				
Characteristics:	Easy to use				
	No specialized equipment needed				
	Compatible with nearly any liquid sample				
	Proven technology (many publications)Highly sensitive (pg/mL)				

- Sandwich ELISA specificity
- Higher density than ELISA, Western blot or bead-based multiplex

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Product Details

Components:	Antibody Array Membranes
	Biotinylated Detection Antibody Cocktail
	Blocking Buffer
	Wash Buffers 1 and 2
	Cell & Tissue Lysis Buffer
	Detection Buffers C and D
	Plastic Incubation Tray
	Protease Inhibitor Cocktail (in select kits)
Material not included:	Pipettors, pipet tips and other common lab consumables
Material not included:	Pipettors, pipet tips and other common lab consumables Orbital shaker or oscillating rocker
Material not included:	
Material not included:	Orbital shaker or oscillating rocker
Material not included:	Orbital shaker or oscillating rocker Tissue Paper, blotting paper or chromatography paper
Material not included:	Orbital shaker or oscillating rocker Tissue Paper, blotting paper or chromatography paper Adhesive tape or Saran Wrap
Material not included:	Orbital shaker or oscillating rocker Tissue Paper, blotting paper or chromatography paper Adhesive tape or Saran Wrap Distilled or de-ionized water

Application Details

Application Notes:	Perform ALL incubation and wash steps under gentle rotation or rocking motion (~0.5 to 1
	cycle/sec) using an orbital shaker or oscillating rocker to ensure complete and even
	reagent/sample coverage. Rocking/rotating too vigorously may cause foaming or bubbles to
	appear on the membrane surface which, should be avoided. All washes and incubations should
	be performed in the Incubation Tray (ITEM 10) provided in the kit. Cover the Incubation Tray
	with the lid provided during all incubation steps to avoid evaporation and outside debris
	contamination. Ensure the membranes are completely covered with sufficient sample or
	reagent volume during each incubation. Avoid forceful pipetting directly onto the membrane,
	instead, gently pipette samples and reagents into a corner of each well. Aspirate samples and
	reagents completely after each step by suctioning off excess liquid with a pipette. Tilting the
	tray so the liquid moves to a corner and then pipetting is an effective method. Optional
	overnight incubations may be performed for the following step to increase overall spot signal
	intensities:
	- Sample Incubation
	- Biotinylated Antibody Cocktail Incubation
	- HRP-Streptavidin Incubation
Comment:	The C-Series arrays feature chemiluminescent signal detection. The antibodies are spotted on

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	nitrocellulose membrane solid supports and are handled in a very similar manner to Western				
	blots.				
	All C-Series arrays work on the sandwich ELISA principle, utilizing a matched pair of antibodies				
	an immobilized capture antibody and a corresponding biotinylated detection antibody.				
Sample Volume:	1 mL				
Plate:	Membrane				
Protocol:	1. Block membranes				
	2. Incubate with Sample				
	3. Incubate with Biotinylated Detection Antibody Cocktail				
	4. Incubate with HRP-Conjugated Streptavidin				
	5. Incubate with Detection Buffers				
	6. Image with chemiluminescent imaging system				
	7. Perform densitometry and analysis				
Restrictions:	For Research Use only				
Handling					
<u> </u>	The antibody printed side of each membrane is marked by a dash (-) or number (#) in the uppe				
<u> </u>	The antibody printed side of each membrane is marked by a dash (-) or number (#) in the upper left corner. Do not allow membranes to dry out during the experiment or they may become				
Handling Handling Advice:	left corner. Do not allow membranes to dry out during the experiment or they may become				
<u> </u>	left corner. Do not allow membranes to dry out during the experiment or they may become fragile and break OR high and/or uneven background may occur. Grasp membranes by the				
<u> </u>	left corner. Do not allow membranes to dry out during the experiment or they may become fragile and break OR high and/or uneven background may occur. Grasp membranes by the corners or edges only using forceps. DO NOT touch printed antibody spots.				
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Handling Advice: Storage:	left corner. Do not allow membranes to dry out during the experiment or they may become fragile and break OR high and/or uneven background may occur. Grasp membranes by the corners or edges only using forceps. DO NOT touch printed antibody spots.				
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Image 1.

	A	В	с	D	E	F	G	н
1	POS	POS	NEG	NEG	Akt (P-S473)	CREB (P-S133)	ERK1 (P-T202/Y204) ERK2 (P-Y185/Y187)	GSK3a (P-S21)
3	GSK3b (P-S9)	HSP27 (P-S82)	JNK (P-T183)	MEK (P-S217/221)	MKK3 (P-S189)	MKK6 (P-S207)	MSK2 (P-S360)	mTOR (P-52448)
5	p38 (P-T180/Y182)	P53 (P-S15)	P7056K (P-T421/S424)	RSK1 (P-S380)	RSK2 (P-S386)	NEG	NEG	POS

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