

AKT (pan) (Phospho Ser473) Rabbit mAb (AR1249)

Key Features

Host Species:	Rabbit		
Reactivity:	Human, Mouse, Rat		
Applications:	WB,IHC,IF,IP,ELISA		
Isotype:	IgG,Карра		
MW:	56kD (Calculated) 60kD (Observed)		
Recommended Dilution Ratios			
IHC:	1:200-500		
WB:	1:1000-5000		
IF:	1:200-1000		
ELISA:	1:5000-20000		
IP:	1:50-200		
Storage	-15°C to -25°C/1 year (Do not lower than -25°C)		
Basic Information			
Clonality	Monoclonal		
Immunogen Information			
Specificity	Endogenous		
Target Information			
Gene name	AKT1/AKT2/AKT3		
Protein Name	RAC-alpha serine/ threonine-protein kinase; RAC-beta serine/ threonine-protein kinase; RAC gamma serine/ threonine-protein kinase		

Organism	Gene ID	UniProt ID
Human	207; 208; 10000	P31749; P31751;
		Q9Y243
Mouse	11651; 11652; 23797	P31750
Rat	24185; 25233; 29414	P47196; P47197;
		Q63484

Cellular Localization

Tissue specificity

Validation Data



 $\begin{array}{c} kDa \\ kDa \\ 180 \\ -130 \\ 100 \\ -70 \\ -55 \\ -70 \\ -55 \\ -35 \\ 25 \\ 25 \\ 15 \\ 10 \\ -10 \end{array} + \begin{array}{c} AKT (Phospho \\ Ser473) \\ -AKT (Phospho \\ Ser473) \\ -10$



Cytoplasm. Nucleus. Cell membrane. Nucleus after activation by integrin-linked protein kinase 1 (ILK1). Nuclear translocation is enhanced by interaction with TCL1A. Phosphorylation on Tyr-176 by TNK2 results in its localization to the cell membrane where it is targeted for further phosphorylations on Thr-308 and Ser-473 leading to its activation and the activated form translocates to the nucleus. Colocalizes with WDFY2 in intracellular vesicles (PubMed:16792529).

Expressed in prostate cancer and levels increase from the normal to the malignant state (at protein level). Expressed in all human cell types so far analyzed. The Tyr-176 phosphorylated form shows a significant increase in expression in breast cancers during the progressive stages i.e. normal to hyperplasia (ADH), ductal carcinoma in situ (DCIS), invasive ductal carcinoma (IDC) and lymph node metastatic (LNMM) stages.

Western Blot analysis using HepG2 whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-PI3-Kinase p85 α rabbit mAb diluted at 1:2000. anti-AKT (Phospho Ser473) Rabbit mAb diluted at 1:2000. Loading control: Mouse anti GAPDH Secondary: Dylight 800, Goat Anti Rabbit IgG Dylight 680, Goat Anti Rabbit IgG

Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-AKT (Phospho Ser473) antibody. The HRP conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody.

Lane 1: NIH-3T3

Lane 2: PC-12

Predicted band size: 56kDa

Observed band size: 60kDa

Mouse lung was stained with anti-AKT (Phospho Ser473) rabbit antibody



Rat lung was stained with anti-AKT (Phospho Ser473) rabbit antibody

Human lung carcinoma was stained with anti-AKT (Phospho Ser473) rabbit antibody

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