

À des fins de recherche uniquement

# Anticorps Monoclonal anti-Phospho-AKT (Ser473)

Numéro de catalogue: 66444-1-Ig 763 Publications



## Informations de base

Numéro de catalogue:	66444-1-Ig	Numéro d'acquisition GenBank:	NM_005163	Méthode de purification:	Purification par protéine A
Taille:	100μl, Concentration: 1500 μg/ml by 207	Identification du gène (NCBI):	v-akt murine thymoma viral oncogene homolog 1	CloneNo.:	1C10B8
Nanodrop:	Nom complet:			Dilutions recommandées:	WB 1:5000-1:50000 IHC 1:100-1:400
Hôte:	Mouse	MW observés:	60-62 kDa		
Isotype:	IgG1				

## Applications

Applications testées:  
FC, IHC, WB, ELISA

Demandes citées:  
FC, IF, IHC, WB

Spécificité de l'espèce:  
Humain, rat, souris

Espèces citées:  
canin, Humain, Lapin, poisson-zèbre, porc, poulet, rat, souris

**Remarque-IHC:** il est suggéré de démasquer l'antigène avec un tampon de TE buffer pH 9,0; (\*) À défaut, le démasquage de l'antigène peut être effectué avec un tampon citrate pH 6,0.

Contrôles positifs:

WB : cellules Jurkat traitées à la calyculine A, cellules HEK-293 traitées à la calyculine A, cellules HEK-293 traitées à la calyculine A, cellules HSC-T6 traitées à la calyculine A, cellules Jurkat traitées au TPA, cellules NIH/3T3 traitées à la calyculine A, cellules PC-3 traitées à la calyculine A

IHC : tissu de cancer du sein humain, cellules Jurkat traitées à la calyculine A, tissu de cancer du côlon humain

## Informations générales

The serine-threonine protein kinase AKT1 is catalytically inactive in serum-starved primary and immortalized fibroblasts. Survival factors can suppress apoptosis in a transcription-independent manner by activating the serine/threonine kinase AKT1, which then phosphorylates and inactivates components of the apoptotic machinery. This antibody detects all the members of AKT with phospho-modification at Ser473.

## Publications notables

Autrice	Pubmed ID	Journal	Application
Wenzhong Peng	36274350	Tissue Cell	WB
Tong Li	33152931	Biomed Pharmacother	WB
Di Cui	36175877	BMC Cancer	WB

## Stockage

Stockage:

Stocker à -20°C. Stable pendant un an après l'expédition.

Tampon de stockage:

PBS avec azoture de sodium à 0,02 % et glycérol à 50 % pH 7,3

L'aliquotage n'est pas nécessaire pour le stockage à -20°C

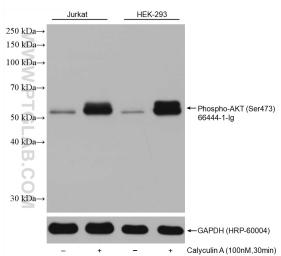
\*\*\* Les 20ul contiennent 0,1% de BSA.

For technical support and original validation data for this product please contact:  
T: 1(888) 4PTGLAB (1-888-478-4522) (toll free  
in USA), or 1(312) 455-8498 (outside USA)

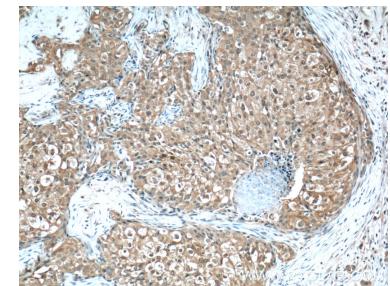
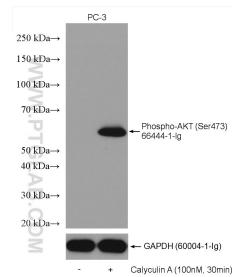
E: proteintech@ptglab.com  
W: ptglab.com

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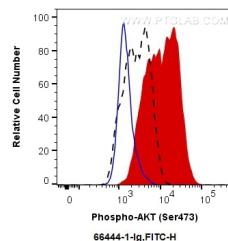
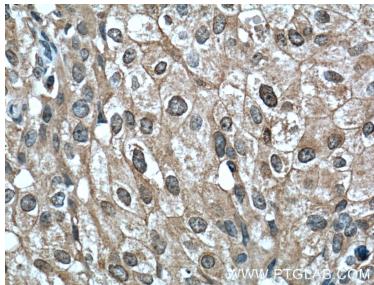
## Données de validation sélectionnées



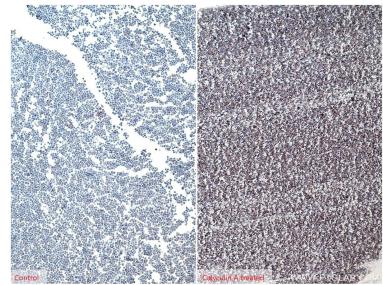
Non-treated Jurkat and HEK-293 cells and Calyculin A treated Jurkat and HEK-293 cells were subjected to SDS PAGE followed by western blot with 66444-1-Ig (Phospho-AKT (Ser473) antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours.



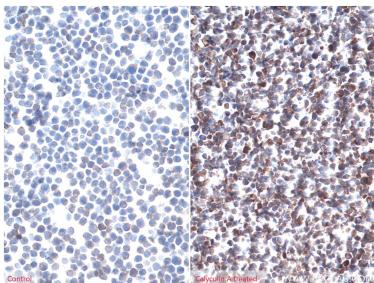
Immunohistochemical analysis of paraffin-embedded human breast cancer tissue slide using 66444-1-Ig (AKT-phospho-S473 antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



1X10<sup>6</sup> PC-3 cells untreated (dashed line) or treated with Calyculin A (red) were intracellularly stained with 0.5 ug Anti-Human Phospho-AKT (Ser473) (66444-1-Ig, Clone:1C10B8) and Coralite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000, or 0.5 ug Control Antibody (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.



Immunohistochemical analysis of paraffin-embedded untreated (left) or Calyculin A treated (right) Jurkat cells slide using 66444-1-Ig (Phospho-AKT (Ser473) antibody) at dilution of 1:8000 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunohistochemical analysis of paraffin-embedded untreated (left) or Calyculin A treated (right) Jurkat cells slide using 66444-1-Ig (Phospho-AKT (Ser473) antibody) at dilution of 1:8000 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).