

À des fins de recherche uniquement

Anticorps Recombinant de lapin anti- Phospho-STAT5A (Tyr694)

Numéro de catalogue: 80115-1-RR

1 Publications



Informations de base

Numéro de catalogue:	BC027036	Méthode de purification:
80115-1-RR	Purification par protéine A	
Taille:	Identification du gène (NCBI):	CloneNo.:
100ul , Concentration: 500 µg/ml by Nanodrop;	6776	2P21
Hôte:	Nom complet:	Dilutions recommandées:
Lapin	signal transducer and activator of transcription 5A	WB 1:2000-1:10000
Isotype:	MW calculé	
IgG	794 aa, 92 kDa	
	MW observés:	
	90-95 kDa	

Applications

Applications testées:	Contrôles positifs:
WB, ELISA	WB : cellules TF-1 traitées au GM-CSF,
Demandes citées:	
WB	
Spécificité de l'espèce:	
Humain	
Espèces citées:	
Humain	

Informations générales

STAT5A, also named as STAT5, belongs to the transcription factor STAT family. In response to cytokines and growth factors, STAT family members are phosphorylated by the receptor associated kinases, and then form homo- or heterodimers that translocate to the cell nucleus where they act as transcription activators. STAT5A is activated by, and mediates the responses of many cell ligands, such as IL2, IL3, IL7 GM-CSF, erythropoietin, thrombopoietin, and different GHs. Phosphorylation at Tyr694 is obligatory for Stat5 activation. Activation of STAT5A in myeloma and lymphoma associated with a TEL/JAK2 gene fusion is independent of cell stimulus and has been shown to be essential for the tumorigenesis. The mouse counterpart of human STAT5A is found to induce the expression of BCL2L1/BCL-X(L), which suggests the antiapoptotic function of STAT5A in cells.

Publications notables

Autrice	Pubmed ID	Journal	Application
Guanyang Chen	35502613	Bioengineered	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

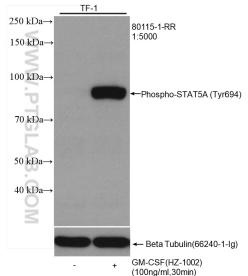
L'aliquotage n'est pas nécessaire pour le stockage à -20°C

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

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Données de validation sélectionnées



Non-treated TF-1 and GM-CSF (HZ-1002) treated TF-1 cells were subjected to SDS PAGE followed by western blot with 80115-1-RR (Phospho-STAT5A (Tyr694) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Tubulin antibody as loading control.