# antibodies -online.com







## anti-STXBP2 antibody (pSer513)



Overview

Binding Specificity:

Clonality:

**Images** 



Quantity:	100 μL
Target:	STXBP2

pSer513

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Reactivity:	Rat	

Conjugate:	This STXBP2 antibody is un-conjugated

Polyclonal

Application:	Western Blotting (WB), Immunohistochemistry (IH	(C)

## **Product Details**

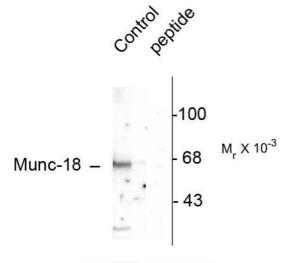
Immunogen:	Synthetic phospho-peptide corresponding to amino acid residues surrounding Ser513 conjugated to KLH
Specificity:	Specific for the ~65k Munc-18 protein phosphorylated at Ser515. Immunolabeling is blocked by the phosphopeptide used as antigen but not by the corresponding dephosphopeptide.
Cross-Reactivity:	Rat (Rattus)
Predicted Reactivity:	bovine, canine, chicken, human, mouse, non-human primate, Xenopus
Purification:	Antigen Affinity Purified from Pooled Serum

## **Target Details**

Target:	STXBP2

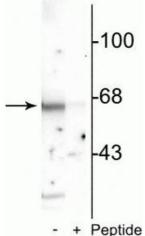
## **Target Details**

rarget Details	
Alternative Name:	STXBP2 (STXBP2 Products)
Background:	Munc-18 (mammalian homologue of Unc-18) is a protein that is thought to be involved in
	regulating exocytosis due, at least in part, to it ability to bind syntaxin (Ciufo et al., 2005).
	Munc18-1 is a neuron-specific member of the Sec1/Munc18 protein family that binds to
	syntaxin1A and is thought to stabilize the complex (Liu et al., 2004). The function of Munc-18 is
	thought to be regulated by PKC phosphorylation of Ser515 on the Munc-18 protein (Sassa et al.
	1996). Anti-Phospho-Ser515 Munc-18 Western blot of rat cortex lysate showing specific
	immunolabeling of the $\sim\!65k$ Munc-18 protein phosphorylated at Ser515. Immunolabeling is
	blocked by the phosphopeptide (peptide) used as antigen but not by the corresponding
	dephosphopeptide (not shown)
Molecular Weight:	'65 kDa
Gene ID:	81804
UniProt:	Q62753
Pathways:	Regulation of Leukocyte Mediated Immunity, Positive Regulation of Immune Effector Process,
	Synaptic Vesicle Exocytosis
Application Details	
Application Notes:	Recommended Dilution: WB: 1:1000 IHC: 1:400 Quality Control: Western blots performed on
	each lot.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	100 $\mu$ L in 10 mM HEPES ( pH 7.5), 150 mM NaCl, 100 $\mu$ g per ml BSA and 50 % glycerol.
Storage:	-20 °C



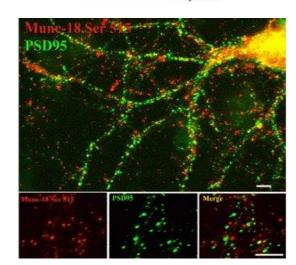
### **Western Blotting**

**Image 1.** Western blots of rat cortex lysate showing specific immunolabeling of the ~65k Munc-18 protein phosphorylated at Ser515. Immunolabeling is blocked by the phosphopeptide (peptide) used as antigen but not by the corresponding dephosphopeptide (not shown).



#### **Western Blotting**

Image 2. Western blot of rat cortical lysate showing specific immunolabeling of the ~65 kDa Munc-18 protein phosphorylated at Ser513 in the first lane (-). Phosphospecificity is shown in the second lane (+) where immunolabeling is blocked by preadsorption of the phosphopeptide used as the antigen, but not by the corresponding non-phosphopeptide (not shown).



#### **Immunostaining**

**Image 3.** Immunostaining of 14 DIV rat cortical neurons showing specific labeling of Munc-18 phosphorylated at Ser513 (red) and PSD95 (green).