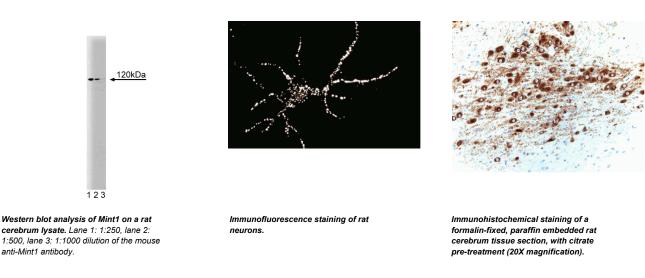
# Technical Data Sheet Purified Mouse Anti-Mint1

Product Information	
Material Number:	611028
Size:	50 µg
Concentration:	250 μg/ml
Clone:	23/Mint1
Immunogen:	Rat Mint 1 aa. 268-377
Isotype:	Mouse IgG1
Reactivity: Target MW:	QC Testing: Rat Tested in Development: Mouse 120 kDa
Storage Buffer:	Aqueous buffered solution containing BSA, glycerol, and $\leq 0.09\%$ sodium azide.

## Description

Neuronal communication via neurotransmitter release is mediated by the synaptic vesicle cycle. The initial step in exocytosis is the docking of the vesicle in the active zone of the plasma membrane. This step is followed by fusion of the vesicle and plasma membrane and exocytosis. Munc18-1, a major brain protein, is essential for exocytosis. It binds the vesicle fusion protein syntaxin, along with Doc2a and 2b, two proteins that associate peripherally with the synaptic vesicle. Munc18-1 is a family member of membrane trafficking proteins. Its function is thought to be mediated by two Munc18-1-interacting proteins termed Mint 1 and Mint 2, which are 50% homologous. They are expressed exclusively in brain and bind Munc18-1 (MID) with high affinity. They contain an N-terminal Munc18-1 interacting domain and C-terminal PTB (pTyr/PIP interaction) and PDZ (membrane protein interaction) domains, suggesting the ability of Mint proteins to link vesicle exocytosis to Tyr phosphorylation and/or localization at synaptic intercellular junctions. Thus, the Mint proteins, along with Munc18-1 and syntaxin, form a multimeric complex that mediates appropriate docking/fusion of synaptic vesicles.

This antibody is routinely tested by western blot analysis. Other applications were tested at BD Biosciences Pharmingen during antibody development only or reported in the literature.



## **Preparation and Storage**

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at  $-20^{\circ}$  C.

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# **Application Notes**

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F	Appreation				
	Western blot	Routinely Tested			
	Immunofluorescence	Tested During Development			
	Immunohistochemistry	Tested During Development			

## **Recommended Assay Procedure:**

Western blot: Please refer to http://www.bdbiosciences.com/pharmingen/protocols/Western\_Blotting.shtml

#### Suggested Companion Products

Catalog Number	Name	Size	Clone
611463	Rat Cerebrum Lysate	500 μg	(none)
554002	HRP Goat Anti-Mouse Igs	1.0 ml	(none)
554001	FITC Goat Anti-Mouse Igs (Multiple Adsorption)	0.5 mg	Polyclonal

## **Product Notices**

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols. 2.
- Source of all serum proteins is from USDA inspected abattoirs located in the United States. 3.
- 4. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.

#### References

Biederer T, Sudhof TC. Mints as adaptors. Direct binding to neurexins and recruitment of munc18. J Biol Chem. 2000; 275(51):39803-39806. (Biology: Immunofluorescence, Western blot)

Fisher RJ, Pevsner J, Burgoyne RD. Control of fusion pore dynamics during exocytosis by Munc18. Science. 2001; 291(5505):875-878.(Biology: Immunoprecipitation)

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Wong RW, Setou M, Teng J, Takei Y, Hirokawa N. Overexpression of motor protein KIF17 enhances spatial and working memory in transgenic mice. Proc Natl Acad Sci U S A. 2002; 99(22):14500-14505.(Biology: Western blot)

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