

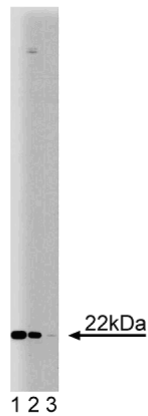
## Technical Data Sheet

**Purified Mouse Anti-Ninjurin****Product Information**

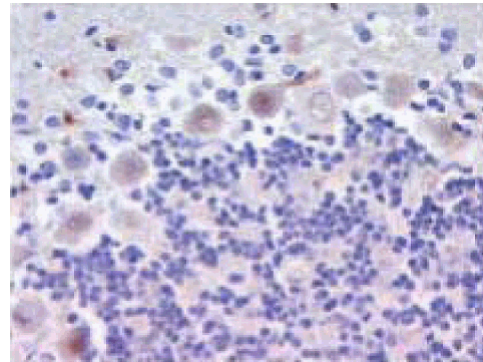
<b>Material Number:</b>	<b>610776</b>
<b>Size:</b>	50 µg
<b>Concentration:</b>	250 µg/ml
<b>Clone:</b>	50/Ninjurin
<b>Immunogen:</b>	Human Ninjurin aa. 1-152
<b>Isotype:</b>	Mouse IgG2a
<b>Reactivity:</b>	QC Testing: Human Tested in Development: Dog, Rat, Mouse
<b>Target MW:</b>	18-22 kDa
<b>Storage Buffer:</b>	Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium azide.

**Description**

Ninjurin is a protein whose expression is dramatically increased after sciatic nerve transection and crush injuries. The *Ninjurin* gene encodes for a protein of 152 amino acids with two putative transmembrane domains. Ninjurin was iodinated in vivo and promoted the aggregation of Jurkat cells expressing Ninjurin, indicating a portion of Ninjurin is exposed to the cell surface. The adhesive properties of Ninjurin were energy and temperature dependent, required Ca<sup>2+</sup> and Mg<sup>2+</sup>, and the integrity of the cytoskeleton. Also, the adhesion domain of Ninjurin is located at the extracellular NH<sub>2</sub>-terminal domain. Although its predicted mass is 16 kDa, Ninjurin migrates as a 18-22 kDa protein in SDS-PAGE, depending on the cell line or tissue. mRNA analysis revealed that Ninjurin is widely expressed with the highest levels in liver, thymus, heart, and the lowest level in brain.



**Western blot analysis of Ninjurin on HepG2 lysate.**  
Lane 1: 1:250, lane 2: 1:500, lane 3: 1:1000 dilution of anti-Ninjurin.



**Immunofluorescent staining on Rat Brain section.**

**Preparation and Storage**

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at -20°C.

**BD Biosciences**

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## Application Notes

### Application

Western blot	Routinely Tested
Immunofluorescence	Tested During Development
Immunohistochemistry	Tested During Development
Immunoprecipitation	Not Recommended

### Recommended Assay Procedure:

Western blot: Please refer to [http://www.bdbiosciences.com/pharming/en/protocols/Western\\_Blotting.shtml](http://www.bdbiosciences.com/pharming/en/protocols/Western_Blotting.shtml).

### Suggested Companion Products

Catalog Number	Name	Size	Clone
611555	HepG2 Cell Lysate	500 µg	(none)
554002	HRP Goat Anti-Mouse Ig	1.0 ml	(none)
554001	FITC Goat Anti-Mouse Ig	0.5 mg	Polyclonal

### Product Notices

1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
2. Please refer to [www.bdbiosciences.com/pharming/en/protocols](http://www.bdbiosciences.com/pharming/en/protocols) for technical protocols.
3. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
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

Araki T, Milbrandt J. Ninjurin, a novel adhesion molecule, is induced by nerve injury and promotes axonal growth. *Neuron*. 1996; 17(2):353-361.(Biology)  
Araki T, Zimonjic DB, Popescu NC, Milbrandt J. Mechanism of homophilic binding mediated by ninjurin, a novel widely expressed adhesion molecule. *J Biol Chem*. 1997; 272(34):21373-21380.(Biology)  
Chen JS, Coustan-Smith E, Suzuki T. Identification of novel markers for monitoring minimal residual disease in acute lymphoblastic leukemia. *Blood*. 2001; 97(7):2115.(Clone-specific: Flow cytometry)