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# **NBL1 ELISA Kit**





### Overview

Quantity:	96 tests
Target:	NBL1
Binding Specificity:	AA 17-178
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	62.5-4000 pg/mL
Minimum Detection Limit:	62.5 pg/mL
Application:	ELISA

## **Product Details**

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Mouse DAN/NBL1
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin), Plasma (EDTA)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: NSO
	Immunogen sequence: A17-D178
Specificity:	NSO, A17-D178
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.
Sensitivity:	<10pg/mL

### **Product Details**

### Material not included:

Microplate reader in standard size. Automated plate washer. Adjustable pipettes and pipette tips. Multichannel pipettes are recommended in the condition of large amount of samples in the detection. Clean tubes and Eppendorf tubes. Washing buffer (neutral PBS or TBS). Preparation of 0.01M TBS: Add 1.2g Tris, 8.5g Nacl

## **Target Details**

Target Details	
Target:	NBL1
Alternative Name:	NBL1 (NBL1 Products)
Background:	Protein Function: Possible candidate as a tumor suppressor gene of neuroblastoma. May play
	an important role in preventing cells from entering the final stage (G1/S) of the transformation
	process.
	Background: Differential screening-selected gene aberrative in neuroblastoma (DAN) is a
	member of the DAN family of secreted glycoproteins that are putative BMP antagonists. The
	NBL1 gene, also known as DAN, is originally cloned from a normal rat fibroblast cDNA library by
	a differential screening method. The human DAN gene is mapped to chromosome 1p36.13-
	p36. It is found that the DAN gene possesses a tumor suppressive activity when overexpressed
	in v-src transformed cells.
	Synonyms: Neuroblastoma suppressor of tumorigenicity 1,N03,Zinc finger protein
	DAN,Nbl1,Dan, Dana,
	Full Gene Name: Neuroblastoma suppressor of tumorigenicity 1
	Cellular Localisation: Secreted.
Gene ID:	17965
UniProt:	Q61477
Application Details	
Application Notes:	Before using Kit, spin tubes and bring down all components to bottom of tube. Duplicate well
	assay was recommended for both standard and sample testing.
Plate:	Pre-coated
Protocol:	mouse DAN ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assay

Application Notes.	before doing fait, opin tabes and bring down an compenents to bottom of tabe. Duplicate wen
	assay was recommended for both standard and sample testing.
Plate:	Pre-coated
Protocol:	mouse DAN ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assay
	technology. A monoclonal antibody from rat specific for DAN has been precoated onto 96-well
	plates. Standards (NSO, A17-D178) and test samples are added to the wells, a biotinylated
	detection polyclonal antibody from goat specific for DAN is added subsequently and then
	followed by washing with PBS or TBS buffer. Avidin-Biotin-Peroxidase Complex was added and

# **Application Details**

Restrictions:	For Research Use only
	Sample 3: n=24, Mean(pg/ml): 3107, Standard deviation: 242.34, CV(%): 7.8
	<ul> <li>Sample 2: n=24, Mean(pg/ml): 1948, Standard deviation: 146.10, CV(%): 7.5</li> </ul>
	<ul> <li>Sample 1: n=24, Mean(pg/ml): 792, Standard deviation: 50.68, CV(%): 6.4</li> </ul>
	• Sample 3: n=16, Mean(pg/ml): 2944, Standard deviation: 197.24, CV(%): 6.7,
	Sample 2: n=16, Mean(pg/ml): 1810, Standard deviation: 112.22, CV(%): 6.2
Assay Precision:	<ul> <li>Sample 1: n=16, Mean(pg/ml): 675, Standard deviation: 37.12, CV(%): 5.5</li> </ul>
	mouse DAN standard solution and each sample is measured in duplicate.
	each empty well. See "Sample Dilution Guideline" above for details. We recommend that each
	properly diluted sample of mouse cell culture supernates, serum or plasma(heparin, EDTA) to
	0.1 mL of the sample diluent buffer into the control well (Zero well). Add 0.1 mL of each
	125pg/mL, 62.5pg/mL mouse DAN standard solutions into the precoated 96-well plate. Add
Assay Procedure:	Aliquot 0.1 mL per well of the 4000pg/mL, 2000pg/mL, 1000pg/mL, 500pg/mL, 250pg/mL,
A D I	
	proportional to the mouse DAN amount of sample captured in plate.
	product that changed into yellow after adding acidic stop solution. The density of yellow is
	to visualize HRP enzymatic reaction. TMB was catalyzed by HRP to produce a blue color
	unbound conjugates were washed away with PBS or TBS buffer. HRP substrate TMB was use

# Handling

Handling Advice:	Avoid multiple freeze-thaw cycles.
Storage:	-20 °C,4 °C
Storage Comment:	Store at 4°C for 6 months, at -20°C for 12 months. Avoid multiple freeze-thaw cycles
Expiry Date:	12 months

# Mouse DAN ELISA Kit 10 10 0.1 10 100 1000 10000 Concentration(pg/ml)

### **ELISA**

**Image 1.** Mouse DAN/NBL1 PicoKine ELISA Kit standard curve