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anti-PRRX1 antibody (Internal Region)

2 Images



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Overview

Quantity:	100 μL
Target:	PRRX1
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)

Product Details

Immunogen:	A synthesized peptide derived from human PRRX1, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	PRRX1 Antibody detects endogenous levels of total PRRX1.
Predicted Reactivity:	Pig,Bovine,Horse,Rabbit,Dog,Chicken
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).

Target Details

Target:	PRRX1
Alternative Name:	PRRX1 (PRRX1 Products)

Target Details

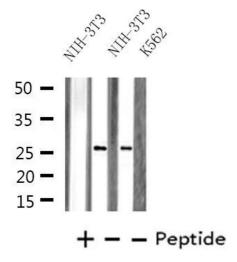
Background:	Description: Acts as a transcriptional regulator of muscle creatine kinase (MCK) and so has a
	role in the establishment of diverse mesodermal muscle types. The protein binds to an A/T-rich
	element in the muscle creatine enhancer (By similarity).
	Gene: PRRX1
Molecular Weight:	27 kDa
Gene ID:	5396
UniProt:	P54821

Application Details

Application Notes:	WB 1:500-1:1000, IF/ICC 1:100-1:500, IHC 1:50-1:200, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

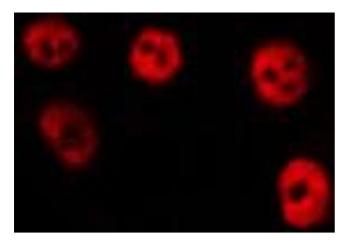
Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 $\%$ sodium azide and 50 $\%$ glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



Western Blotting

Image 1. Western blot analysis of extracts from NIH-3T3/K562 cells, using PRRX1 antibody.



Immunofluorescence (fixed cells)

Image 2. ABIN6275339 staining HepG2 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibod