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# **Myeloperoxidase ELISA Kit**



Image



**Publications** 



#### Overview

Quantity:	96 tests
Target:	Myeloperoxidase (MPO)
Binding Specificity:	AA 49-745
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	312-20000 pg/mL
Minimum Detection Limit:	312 pg/mL
Application:	ELISA

### **Product Details**

Sandwich High Sensitivity ELISA kit for Quantitative Detection of Human MPO
PicoKine™
Cell Culture Supernatant, Cell Lysate, Tissue Homogenate, Serum, Plasma (heparin), Plasma (EDTA), Saliva, Urine
Quantitative
Colorimetric
Expression system for standard: NSO Immunogen sequence: A49-S745
Expression system for standard: NSO Immunogen sequence: A49-S745

#### **Product Details**

Cross-Reactivity (Details):

Sensitivity:	<10pg/mL
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

There is no detectable cross-reactivity with other relevant proteins.

# Target Details

Target:	Myeloperoxidase (MPO)
Alternative Name:	MPO (MPO Products)
Background:	Protein Function: Part of the host defense system of polymorphonuclear leukocytes. It is
	responsible for microbicidal activity against a wide range of organisms. In the stimulated PMN
	MPO catalyzes the production of hypohalous acids, primarily hypochlorous acid in physiologic
	situations, and other toxic intermediates that greatly enhance PMN microbicidal activity.
	Background: Myeloperoxidase(MPO) is a mammalian phagocyte hemoprotein thought to
	primarily mediate host defense reactions. It is abundantly expressed in neutrophils and
	secreted during their activation. Myeloperoxidase is part of the host defense system of human
	polymorphonuclear leukocytes, responsible for microbicidal activity against a wide range of
	organisms. It is located in the nucleus as well as in the cytoplasm. Intranuclear MPO may help
	to protect DNA against damage resulting from oxygen radicals produced during myeloid cell
	maturation and function. The standard product used in this kit is the product of gene
	recombination, consisting of 697(A49-S745) amino acids with the molecular mass of 80KDa.
	Synonyms: Myeloperoxidase,MPO,1.11.2.2,Myeloperoxidase,89 kDa myeloperoxidase,84 kDa
	myeloperoxidase, Myeloperoxidase light chain, Myeloperoxidase heavy chain, MPO,
	Full Gene Name: Myeloperoxidase
	Cellular Localisation: Lysosome.
Gene ID:	4353
UniProt:	P05164
Pathways:	Chromatin Binding
Application Details	

Application Notes: Before using Kit, spin tubes and bring down all components to bottom of tube. Duplicate well

# **Application Details**

	assay was recommended for both standard and sample testing.
Comment:	Sequence similarities: Belongs to the peroxidase family. XPO subfamily.
Plate:	Pre-coated
Protocol:	human MPO ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assa
	technology. A monoclonal antibody from mouse specific for MPO has been precoated onto 96
	well plates. Standards(NSO, A49-S745) and test samples are added to the wells, a biotinylated
	detection polyclonal antibody from goat specific for MPO is added subsequently and then
	followed by washing with PBS or TBS buffer. Avidin-Biotin-Peroxidase Complex was added and
	unbound conjugates were washed away with PBS or TBS buffer. HRP substrate TMB was used
	to visualize HRP enzymatic reaction. TMB was catalyzed by HRP to produce a blue color
	product that changed into yellow after adding acidic stop solution. The density of yellow is
	proportional to the human MPO amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 20000pg/mL, 10000pg/mL, 5000pg/mL, 2500pg/mL,
	1250pg/mL, 625pg/mL, 312pg/mL human MPO standard solutions into the precoated 96-well
	plate. Add 0.1 mL of the sample diluent buffer into the control well (Zero well). Add 0.1 mL of
	each properly diluted sample of human cell culture supernates, cell lysate, tissue lysate, serum
	plasma(heparin, EDTA), saliva or urine to each empty well. See "Sample Dilution Guideline"
	above for details. It is recommended that each human MPO standard solution and each sample
	be measured in duplicate.
Assay Precision:	<ul> <li>Sample 1: n=16, Mean(ng/ml): 2.3, Standard deviation: 0.076, CV(%): 3.3</li> </ul>
	Sample 2: n=16, Mean(ng/ml): 7.4, Standard deviation: 0.4, CV(%): 5.4
	<ul> <li>Sample 3: n=16, Mean(ng/ml): 12.4, Standard deviation: 0.72, CV(%): 5.8,</li> </ul>
	• Sample 1: n=24, Mean(ng/ml): 2.9, Standard deviation: 0.122, CV(%): 4.2
	Sample 2: n=24, Mean(ng/ml): 8, Standard deviation: 0.488, CV(%): 6.1      Sample 2: n=24, Mean(ng/ml): 14.7, Standard deviation: 0.056, CV(%): 6.5
	<ul> <li>Sample 3: n=24, Mean(ng/ml): 14.7, Standard deviation: 0.956, CV(%): 6.5</li> </ul>
Restrictions:	For Research Use only
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

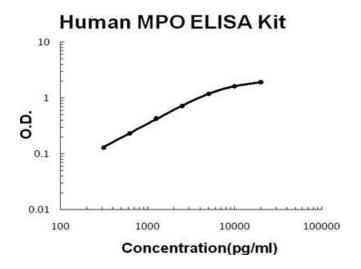
Product cited in:

Fernández, Baldassarro, Sivilia, Giardino, Calzà: "Inflammation severely alters thyroid hormone signaling in the central nervous system during experimental allergic encephalomyelitis in rat: Direct impact on OPCs differentiation failure." in: **Glia**, Vol. 64, Issue 9, pp. 1573-89, (2016) (PubMed).

Vidart, Wajner, Leite, Manica, Schaan, Larsen, Maia: "N-acetylcysteine administration prevents nonthyroidal illness syndrome in patients with acute myocardial infarction: a randomized clinical trial." in: **The Journal of clinical endocrinology and metabolism**, Vol. 99, Issue 12, pp. 4537-45, (2014) (PubMed).

There are more publications referencing this product on: Product page

#### **Images**



#### **ELISA**

Image 1. Human MPO PicoKine ELISA Kit standard curve