

# Datasheet for ABIN2859305

# **HMOX1 ELISA Kit**

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Publication



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### Overview

Quantity:	96 tests
Target:	HMOX1
Binding Specificity:	AA 1-288
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	93.7-6000 pg/mL
Minimum Detection Limit:	93.7 pg/mL
Application:	ELISA

## **Product Details**

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Human HO-1/HMOX1
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin), Plasma (EDTA)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: E.coli
	Immunogen sequence: M1-M288
Specificity:	E.coli, M1-M288
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:	HMOX1
Alternative Name:	HMOX1 (HMOX1 Products)
Background:	Protein Function: Heme oxygenase cleaves the heme ring at the alpha methene bridge to form
	biliverdin. Biliverdin is subsequently converted to bilirubin by biliverdin reductase. Under
	physiological conditions, the activity of heme oxygenase is highest in the spleen, where
	senescent erythrocytes are sequestrated and destroyed. Exhibits cytoprotective effects since
	excess of free heme sensitizes cells to undergo apoptosis.
	Background: HMOX1 (heme oxygenase (decycling) 1) is a human gene that encodes for the
	enzyme heme oxygenase 1. It is localized to chromosome 22. Heme oxygenase, an essential
	enzyme in heme catabolism, cleaves heme to form biliverdin, which is subsequently converted
	to bilirubin by biliverdin reductase, and carbon monoxide, a putative neurotransmitter. Heme
	oxygenase activity is induced by its substrate heme and by various nonheme substances.
	Heme oxygenase occurs as 2 isozymes, an inducible heme oxygenase-1 and a constitutive
	heme oxygenase-2. HMOX1 and HMOX2 belong to the heme oxygenase family.
	Synonyms: Heme oxygenase 1,HO-1,1.14.99.3,HMOX1,HO, HO1,
	Full Gene Name: Heme oxygenase 1
	Cellular Localisation: Microsome . Endoplasmic reticulum membrane, Peripheral membrane
	protein, Cytoplasmic side.
Gene ID:	3162
UniProt:	P09601
Pathways:	Transition Metal Ion Homeostasis, Regulation of Leukocyte Mediated Immunity, Positive
	Regulation of Immune Effector Process, Production of Molecular Mediator of Immune
	Response, SARS-CoV-2 Protein Interactome
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.

# **Application Details**

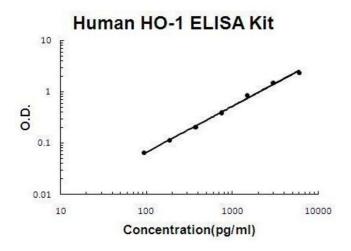
Comment:	Tissue Specificity: Expressed at higher levels in renal cancer tissue than in normal tissue (at
Comment.	protein level).
Plate:	Pre-coated
Protocol:	human HO-1 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assa
	technology. A monoclonal antibody from mouse specific for HO-1 has been precoated onto 96
	well plates. Standards(E.coli, M1-M288) and test samples are added to the wells, a biotinylated
	detection polyclonal antibody from goat specific for HO-1 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 HO-1 amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 6000pg/mL, 3000pg/mL, 1500pg/mL, 750pg/mL, 375pg/mL,
	187.5pg/mL, 93.7pg/m human HO-1 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, serum or plasma(heparin, EDTA) to
	each empty well. See "Sample Dilution Guideline" above for details. It is recommended that
	each human HO-1 standard solution and each sample be measured in duplicate.
Assay Precision:	<ul> <li>Sample 1: n=16, Mean(pg/ml): 527, Standard deviation: 19, CV(%): 3.6</li> </ul>
	• Sample 2: n=16, Mean(pg/ml): 2085, Standard deviation: 91.74, CV(%): 4.4
	• Sample 3: n=16, Mean(pg/ml): 3561, Standard deviation: 185.2, CV(%): 5.2,
	<ul> <li>Sample 1: n=24, Mean(pg/ml): 686, Standard deviation: 31.6, CV(%): 4.6</li> </ul>
	Sample 2: n=24, Mean(pg/ml): 2664, Standard deviation: 151.8, CV(%): 5.7
	<ul> <li>Sample 3: n=24, Mean(pg/ml): 3724, Standard deviation: 242.1, 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

### **Publications**

Product cited in:

Huang, Zhu, Li, Sui, Min: "Effect of Blueberry Anthocyanins Malvidin and Glycosides on the Antioxidant Properties in Endothelial Cells." in: **Oxidative medicine and cellular longevity**, Vol. 2016, pp. 1591803, (2017) (PubMed).

#### **Images**



#### **ELISA**

**Image 1.** Human HO-1/HMOX1 PicoKine ELISA Kit standard curve