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### MMP12 ELISA Kit

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**Publications** 



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#### Overview

Quantity:	96 tests
Target:	MMP12
Binding Specificity:	AA 18-462
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	62.5-4000 pg/mL
Minimum Detection Limit:	62.5 pg/mL
Application:	ELISA

#### **Product Details**

Sample Type: Cell Culture Supernatant, Serum, Plasma (heparin)		
Sample Type: Cell Culture Supernatant, Serum, Plasma (heparin)  Analytical Method: Quantitative  Detection Method: Colorimetric  Immunogen: Expression system for standard: NSO Immunogen sequence: A18-C462  Specificity: Expression system for standard: NSO Immunogen sequence: A18-C462	Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Mouse MMP-12
Analytical Method:  Detection Method:  Colorimetric  Expression system for standard: NSO  Immunogen sequence: A18-C462  Specificity:  Expression system for standard: NSO  Immunogen sequence: A18-C462	Brand:	PicoKine™
Detection Method:  Colorimetric  Expression system for standard: NSO  Immunogen sequence: A18-C462  Specificity:  Expression system for standard: NSO  Immunogen sequence: A18-C462	Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin)
Immunogen: Expression system for standard: NSO Immunogen sequence: A18-C462  Specificity: Expression system for standard: NSO Immunogen sequence: A18-C462	Analytical Method:	Quantitative
Immunogen sequence: A18-C462  Specificity: Expression system for standard: NSO  Immunogen sequence: A18-C462	Detection Method:	Colorimetric
Specificity: Expression system for standard: NSO  Immunogen sequence: A18-C462	Immunogen:	Expression system for standard: NSO
Immunogen sequence: A18-C462		Immunogen sequence: A18-C462
	Specificity:	Expression system for standard: NSO
Cross-Reactivity (Details): There is no detectable cross-reactivity with other relevant proteins.		Immunogen sequence: A18-C462
	Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.

#### **Product 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	
Target Details		
Target:	MMP12	
Alternative Name:	MMP12 (MMP12 Products)	
Background:	Protein Function: May be involved in tissue injury and remodeling. Has significant elastolytic	
	activity. Can accept large and small amino acids at the P1' site, but has a preference for leucine	
	Aromatic or hydrophobic residues are preferred at the P1 site, with small hydrophobic residues	
	(preferably alanine) occupying P3 (By similarity)	
	Background: Matrix metalloproteinase-12(MMP12), also known as MME or ME, is an enzyme	
	that in humans is encoded by the MMP12 gene. The gene is part of a cluster of MMP genes	
	which localize to chromosome 11q22.2. It is thought that the protein encoded by this gene is	
	cleaved at both ends to yield the active enzyme, but this processing has not been fully	
	described. The enzyme degrades soluble and insoluble elastin. It may play a role in aneurysm	
	formation and studies in mice suggest a role in the development of emphysema. This gene	
	may involved in tissue injury and remodeling.	
	Synonyms: Macrophage metalloelastase,MME,3.4.24.65,Matrix metalloproteinase-12,MMP-	
	12,Mmp12,Mme, Mmel,	
	Full Gene Name: Macrophage metalloelastase	
	Cellular Localisation: Secreted, extracellular space, extracellular matrix.	
Gene ID:	17381	
UniProt:	P34960	
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.	
Comment:	Sequence similarities: Belongs to the peptidase M10A family.	
Plate:	Pre-coated	

## Application Details

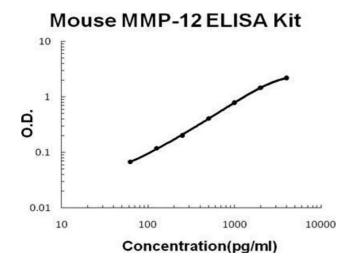
Protocol:	mouse MMP-12 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent
	assay technology. A monoclonal antibody from rat specific for MMP-12 has been precoated
	onto 96-well plates. Standards(NSO, A18-C462) and test samples are added to the wells, a
	biotinylated detection polyclonal antibody from goat specific for MMP-12 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 mouse MMP-12 amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 4000pg/mL, 2000pg/mL, 1000pg/mL, 500pg/mL, 250pg/mL,
	125pg/mL, 62.5pg/mL mouse MMP-12 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 mouse cell culture supernates, serum or plasma(heparin) to each
	empty well. See "Sample Dilution Guideline" above for details. It is recommended that each
	mouse MMP-12 standard solution and each sample be measured in duplicate.
Assay Precision:	• Sample 1: n=16, Mean(pg/ml): 337, Standard deviation: 18.2, CV(%): 5.4
	<ul> <li>Sample 2: n=16, Mean(pg/ml): 1472, Standard deviation: 92.74, CV(%): 6.3</li> </ul>
	• Sample 3: n=16, Mean(pg/ml): 2522, Standard deviation: 121.1, CV(%): 4.8,
	• Sample 1: n=24, Mean(pg/ml): 426, Standard deviation: 27.69, CV(%): 6.5
	<ul> <li>Sample 2: n=24, Mean(pg/ml): 1538, Standard deviation: 109.2, CV(%): 7.1</li> <li>Sample 3: n=24, Mean(pg/ml): 2768, Standard deviation: 157.8, CV(%): 5.7</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:	Papakonstantinou, Karakiulakis, Batzios, Savic, Roth, Tamm, Stolz: "Acute exacerbations of
	COPD are associated with significant activation of matrix metalloproteinase 9 irrespectively of
	airway obstruction, emphysema and infection." in: Respiratory research, Vol. 16, pp. 78, (2016)

#### (PubMed).

Zang, Zhuang, Deng, Yang, Ye, Xie, Ren, Fu, Luo, Xu, Liu: "Pulmonary C Fibers Modulate MMP-12 Production via PAR2 and Are Involved in the Long-Term Airway Inflammation and Airway Hyperresponsiveness Induced by Respiratory Syncytial Virus Infection." in: **Journal of virology**, Vol. 90, Issue 5, pp. 2536-43, (2016) (PubMed).

Vaz, Rajasekaran, Potteti, Reddy: "Myeloid-specific Fos-related antigen-1 regulates cigarette smoke-induced lung inflammation, not emphysema, in mice." in: **American journal of respiratory cell and molecular biology**, Vol. 53, Issue 1, pp. 125-34, (2015) (PubMed).

#### **Images**



#### **ELISA**

**Image 1.** Mouse MMP-12 PicoKine ELISA Kit standard curve