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Datasheet for ABIN367449 Motilin ELISA Kit

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Overview

Quantity:	96 tests
Target:	Motilin (MLN)
Reactivity:	Rat
Method Type:	Sandwich ELISA
Detection Range:	31.25-2000 pg/mL
Minimum Detection Limit:	31.25 pg/mL
Application:	ELISA

Product Details

Purpose:	For the quantitative determination of rat motilin(MTL) concentrations in serum, plasma, tissue homogenates.
Sample Type:	Serum, Plasma, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of rat MTL.
Cross-Reactivity (Details):	Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between the target antigen and all analogues for other species. Therefore, cross reaction may still exist.
Sensitivity:	7.8 pg/mL
Components:	Assay plate (12 × 8 coated Microwells)

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- Standard (freeze dried)
- Biotin-antibody (100 × concentrate)
- HRP-avidin (100 × concentrate)
- Biotin-antibody Diluent
- HRP-avidin Diluent
- Sample Diluent
- Wash Buffer (25 × concentrate)
- TMB Substrate
- Stop Solution
- Adhesive Strip (for 96 wells)
- Instruction manual

Target Details

Target:	Motilin (MLN)
Alternative Name:	Motilin (MTL) (MLN Products)
Pathways:	Hormone Activity
Application Details	
Application Notes:	 The supplier is only responsible for the kit itself, but not for the samples consumed during the assay. The user should calculate the possible amount of the samples used in the whole test. Please reserve sufficient samples in advance. Samples to be used within 5 days may be stored at 2-8°C, otherwise samples must be stored at -20°C (≤ 1 month) or -80°C (≤ 2 months) to avoid loss of bioactivity and contamination. Grossly hemolyzed samples are not suitable for use in this assay. If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary. Please predict the concentration before assaying. If values for these are not within the range of the standard curve, users must determine the optimal sample dilutions for their particular experiments. Tissue or cell extraction samples prepared by chemical lysis buffer may cause unexpected ELISA results due to the impacts of certain chemicals. Owing to the possibility of mismatching between antigens from another resource and antibodies used in this supplier's kits (e.g., antibody targets conformational epitope rather than linear epitope), some native or recombinant proteins from other manufacturers may nor be recognized by this supplier's products. Influenced by factors including cell viability, cell number and cell sampling time, samples from cell culture supernatant may not be recognized by the kit. Fresh samples without long time storage are recommended for the test. Otherwise, protein
	degradation and denaturalization may occur in those samples and finally lead to wrong

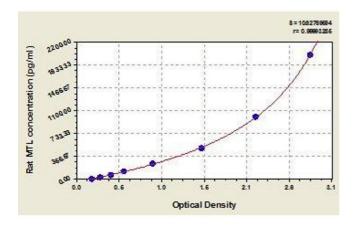
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Application Details

	results.
Comment:	Detection wavelength: 450 nm
	Information on standard material:
	Depending on the antigen to be detected, standards can be either native or recombinant
	protein. The recombinant proteins are being expressed in CHO cells in most cases. Please
	inquire for more information. The formulation of auxiliary material in the standard is considered
	proprietary information, however it does not contain any poisonous substance. Proclin 300
	(1:3000) is used as preservative.
	Information on reagents:
	In most cases the stop solution provided is 1 N H2SO4. The formulation of wash solution is
	proprietary information. None of the components contain (sodium) azide, thimerosal, 2-
	mercaptoethanol (2-ME) or any other poisonous materials. For the sandwich method kits, the
	sample diluent, antibody diluent, enzyme diluent and standard all contain BSA.
	Information on antibodies:
	The antibodies provided in different kits vary in regards to clonality and host. Some antibodies
	are affinity purified, some are Protein A
Sample Volume:	100 μL
Assay Time:	1 - 4.5 h
Plate:	Pre-coated
Protocol:	This assay employs the quantitative sandwich enzyme immunoassay technique. Antibody
	specific for MTL has been pre-coated onto a microplate. Standards and samples are pipetted
	into the wells and any MTL present is bound by the immobilized antibody. After removing any
	unbound substances, a Horseradish Peroxidase (HRP)-conjugated antibody specific for MTL is
	added to the wells. Following a wash to remove any unbound reagent, a substrate solution is
	added to the wells and color develops in proportion to the amount of MTL bound in the initial
	step. The color development is stopped and the intensity of the color is measured.
Assay Precision:	Intra-assay precision (precision within an assay): Three samples of known concentration were
	tested twenty times on one plate to assess precision.
	Inter-assay precision (precision between assays): Three samples of known concentration were
	tested in twenty assays to assess precision.

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Application Details	
	Intra-assay: CV% less than 8%
	Inter-assay: CV% less than 10%
Restrictions:	For Research Use only
Handling	
Precaution of Use:	The Stop Solution provided with this kit is an acid solution. Wear eye, hand, face and clothing
	protection when using this material.
Handling Advice:	The kit should not be used beyond the expiration date on the kit label.
	 Do not mix or substitute reagents with those from other lots or sources.
	 If samples generate values higher than the highest standard, dilute the samples with Sample Diluent and repeat the assay.
	 Any variation in Sample Diluent, operator, pipetting technique, washing technique, incubation
	time/temperature and kit age can cause variation in binding.
	• This assay is designed to eliminate interference by soluble receptors, binding proteins and
	other factors present in biological samples. Until all factors have been tested in the
	Immunoassay, the possibility of interference cannot be excluded.
Storage:	4 °C/-20 °C
Storage Comment:	For unopened kit: All the reagents should be kept according to the labels on vials.
Expiry Date:	6 months
Publications	
Product cited in:	Zhang, Lin, Luo, Zhang, Guo, Wu, Shi, Deng, Chen, Zhang, Bharucha, Huang, Sutton, Windsor,
	Xue, Xia: "Chai-Qin-Cheng-Qi Decoction and Carbachol Improve Intestinal Motility by Regulating
	Protein Kinase C-Mediated Ca2+ Release in Colonic Smooth Muscle Cells in Rats with Acute
	Necrotising Pancreatitis." in: Evidence-based complementary and alternative medicine :
	eCAM , Vol. 2017, pp. 5864945, (2017) (PubMed).
	Feng, Li, Chen, Lin, Yang, Cai, Lv, Cao, Li, Xu, Li, Jia: "Inhibiting roles of berberine in gut
	movement of rodents are related to activation of the endogenous opioid system." in:



ELISA

Image 1. Typical standard curve

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