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ORDIN DE PLATA NR.: 2919                                TIP.DOC. 1 :
                                DATA EMITERII:vineri, 14 iunie 2:
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PLATITI: 4800-00                                LEI: Patru Mii Opt Sute lei 00 bani :
:
:
=====:
PLATITOR: (R) "BIOSISTEM          CONTUL DE PLATI/CODUL IBAN :
MLD" S.R.L.                                MD95ML000000002251429243 :
                                CODUL FISCAL :1010600028048 / :
:
:
=====:
PRESTATORUL PLATITOR                                CODUL BANCII:
BC"Moldindconbank"S.A. suc."Invest" Chisinau        :MOLDMD2X329:
=====:
BENEFICIAR (R) Institutul          CONTUL DE PLATI/CODUL IBAN :
de Medicina Urgenta IMSP            MD55VI022510300000002MDL :
                                CODUL FISCAL :1003600152606 / :
:
:
=====:
PRESTATORUL BENEFICIAR                                CODUL BANCII:
B.C."VICTORIABANK"S.A.                :VICBMD2X :
=====:
DESTINATIA PLATII: Pentru garantia pentru:          TIPUL TRANSFERULUI :
oferta la procedura de achizitie public:            NORMAL/URGENT :N:
a nr. ocds-b3wdpl-MD-1717752817958 din :           :
17.06.2024 :                                       :
:                                       :
:                                       :
:                                       L.S. :
=====:
                                CODUL TRANZACTIEI:001: :
                                DATA PRIMIRII:14/06/2024 : SEMNATURILE :
                                DATA EXECUTARII: : EMITENTULUI :
:-----:
CONducATOR:Web Poiata Vitalie :
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SIB3DQEBCwUAMCIxIDAeBgNVBAMTF0NFULQxLUNBLUL1vbGRpbmRjb25iYW5rMB4X :
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:
: (semnatura electronica) :
CONTABIL-SEF:Web Nasedchin Alexandr :
MIIGZwYJKoZIhvcNAQcCoIIGWDCcBlQCAQExCzAJBgUrDgMCGGUAMAsGCSqGSIB3 :
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:
: L.S. (semnatura electronica) :
CONducATOR: :
: (semnatura manuala) :
CONTABIL-SEF: :
: (semnatura manuala) :
SEMnATURA PRESTATORUL L.S. :
:-----:
MOTIVUL REFUZULUI : L.S. :
-----:

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BC "MOLDINDCONBANK" S.A. Filiala "Invest"

Republica Moldova, MD-2068
mun. Chişinău, bd. Moscovei, 14/1
Tel. : (373-22) 43-44-81, 43-46-24
Fax : (373-22) 43-44-22
cod: MOLDM2X329

Data 14. IAN. 2016
Nr. 03/2 - 19/23

Республика Молдова, MD-2068
мун. Кишинэу, бул. Московей, 14/1
Тел. : (373-22) 43-44-81, 43-46-24
Факс : (373-22) 43-44-22
код: MOLDM2X329

Filiala „Invest” BC „Moldindconbank” SA confirmă existența contului curent
in moneda nationala al **“BIOSISTEM MLD” S.R.L. (c/f 1010600028048)**, cu
IBAN MD95ML000000002251429243.

Codul băncii MOLDM2X329.

Director

Nina Turcan

Director financiar



Nina Balmuş

Ex. Diana Brinza
Tel. 43-45-96

REPUBLICA



MOLDOVA

CERTIFICAT DE ÎNREGISTRARE

Societatea cu Răspundere Limitată "BIOSISTEM MLD"
— ESTE ÎNREGISTRATĂ LA CAMERA ÎNREGISTRĂRII DE STAT —

Numărul de identificare de stat - codul fiscal
1010600028048

Data înregistrării

12.08.2010

Data eliberării

12.08.2010

Svirepova Ludmila, registrator

*Funcția, numele, prenumele persoanei
care a eliberat certificatul*

L. Svirepova
semnătura

MD 0101250





AGENȚIA SERVICII PUBLICE

Departamentul înregistrare și licențiere a unităților de drept

EXTRAS

din Registrul de stat al persoanelor juridice

Nr. 531522 data 15.09.2023

Denumirea completă: **Societatea cu Răspundere Limitată "BIOSISTEM MLD"**

Denumirea prescurtată: **"BIOSISTEM MLD" S.R.L.**

Forma juridică de organizare: **Societate cu răspundere limitată,**

Numărul de identificare de stat și codul fiscal (IDNO): **1010600028048**

Data înregistrării de stat: **12.08.2010**

Sediul: **MD-2001, str. Albișoara, 16/1, ap. 7, mun. Chișinău, Republica Moldova.**

Obiectul principal de activitate:

- 1. Activitatea farmaceutică; importul și (sau) producerea articolelor de parfumerie și cosmetică**
- 2. Fabricarea, comercializarea, asistența tehnică, repararea și verificarea articolelor de tehnică și optică medicală**
- 3. Acordarea asistenței medicale de către instituțiile medico-sanitare private**
- 4. Comerțul cu ridicata al calculatoarelor, echipamentelor periferice și software-ului**
- 5. Întreținerea și repararea mașinilor de birou și a tehnicii de calcul**
- 6. Consultații în domeniul sistemelor de calcul**

Capitalul social: **5400 lei.**

Administrator: **POIATA VITALIE, IDNP 0983103892591,**

Asociații:

1. **POIATA VITALIE, IDNP 0983103892591, cota 1803,60 lei, ce constituie 33,4%**

Beneficiar efectiv:

1.1. **POIATA VITALIE, IDNP 0983103892591,**

2. **NASEDCHIN ALEXANDR, IDNP 2002001070747, cota 1798,20 lei, ce constituie 33,3%**

Beneficiar efectiv:

2.1. **NASEDCHIN ALEXANDR, IDNP 2002001070747,**

3. **KOJEVNIKOV DMITRII, IDNP 0972305012362, cota 1798,20 lei, ce constituie 33,3%**

Beneficiar efectiv:

3.1. **KOJEVNIKOV DMITRII, IDNP 0972305012362**

Prezentul extras este eliberat în temeiul art.34 al Legii nr.220-XVI din 19 octombrie 2007 privind înregistrarea de stat a persoanelor juridice și a întreprinzătorilor individuali și confirmă datele din Registrul de stat la data de: **15.09.2023.**

**Registrator în domeniul
înregistrării de stat**

Digitally signed by Rusu Diana
Date: 2023.09.15 16:44:17 EEST
Reason: MoldSign Signature
Location: Moldova



Rusu Diana



EB 0461494

Lista fondatorilor Biosistem-mld SRL

Nr.	Nume, Prenume	IDNP
1.	Vitalie Poiata	0983103892591
2.	Alexandr Nasedchin	2002001070747
3.	Dmitrii Kojevnikov	0972305012362

[QinPot Main Body]

Product Number	Product Name	Quantity	Note
CQR10-Y-EU	QinPot Yellow	1 pc	Pipe Connector is separately sold as an option. *Pipe Connector is not included.
CQR10-P-EU	QinPot Pink	1 pc	
CQR10-G-EU	QinPot Green	1 pc	
CQR10-B-EU	QinPot Blue	1 pc	

• For different type of pipe connector, consult your COOPDECH representative.

[QinPot Liner]

Product Number	Product Name	Quantity	Note
CQD10-H-EU	QinPot Liner	50 pcs	
CQD10-G-EU	QinPot Liner Solidifying	50 pcs	

• Specification and design may be changed for improvement without prior notice.

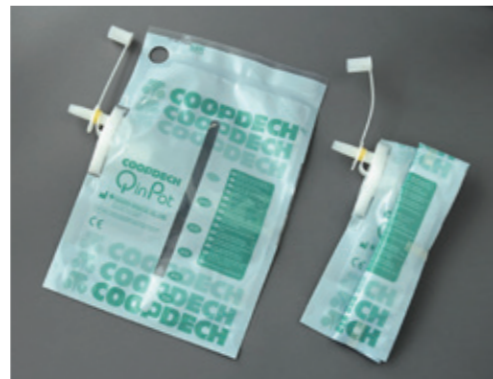
[Accessories]

Product Number	Product Name	Quantity	Note
RD822KC	Pipe Connector	1 pc	Pin type(Kawaju,Uni type)
RD822A	Pipe Connector	1 pc	Schroeder(Amco type)

QinPot Main Body



QinPot Liner



QinPot Connecting Tube



Daiken Medical Co., Ltd. has obtained ISO 13485 certification, an international standard on quality management systems related to medical devices.

Pioneering the future of medical society
DAIKEN MEDICAL CO., LTD.
International Department: 1-6-6, Funakoshi-cho, Chuo-ku, Osaka-city, 540-0036, Japan
COOPDECH's product information is available at the corporate website.
<http://www.daiken-iki.co.jp/>

[Agency]

2016.10

Contents of this catalog are as of October 2016.



COOPDECH

COOPDECH QinPot

(The Wall-mounted Type Closed 1000mL Suction System)

Simple

It has only one connection port, it is easy to set and very unlikely to misconnect.

Safe

It is the alternative to glass jars & other disposable suction system. Solidifying substance is preinstalled inside the liner. Fluids are stored inside without the risk of direct exposure.

Compact

Thanks to its compact size, it is well suited for installations under limited space, and 4 colors (Blue, Green, Pink and Yellow) are available. It is suitably designed for hospital wards and ICU.

DAIKEN MEDICAL CO., LTD.



Cleaner and easier to use. The optimum shape for safety and usability.



Simple Control

The cover lock, ON-OFF lever and pressure controller operating parts are simple in look and colored gray.

Hygiene

For high hygienic standard, hydrophobic filter are acting as antibacterial and over flow protection.

Dual Safety

For higher safety standard, built in hydrophobic filter is equipped as standard.



The liner can be attached facing toward either left or right.



The liner can be removed with the suction tube.

Speedy Setting and Disposal

The liner can be ready for use in QinPot with one hand only.

Liner with Built-in Check Valve

The suction tube connection port of the liner incorporates an one-way check valve to prevent accidental overflow of drained substances. Also the cap effectively seals the contents of the liner.

Autoclavable Transparent and Rigid Canister

Highly transparent and rigid canister. The canister is made of high impact polycarbonate with high transparency. The fluid flow and general operation can be checked via the slit window of the liner.

(OPTION) Pressure Controller with Lock System

The pressure controller is positioned at the bottom of the device for better viewing and operation. Once the controller is pushed in, lock system fixes its pressure.

Independent ON-OFF Lever

The ON-OFF lever is independent and convenient for frequent use.



One connection only

QinPot Liner requires only one connection. Just connect the patient tubing and the liner is ready for use.

Compact design

QinPot is small enough to be placed at the bedside. The liner is thin and foldable, allowing compact storage from packing to disposal. (Dimensions of outer carton box of 50 pieces: 420 x 270 x 310 mm.)

Cost-effective

The liner's cost has been greatly reduced while still maintaining an extremely safe design. It remains compact even in disposal, helping to reduce costs. (15~20% less disposal cost, for more information please contact COOPDECH representatives.)

Safe

The device is a sealed unit with the liner containing a coagulant for the drained substances, so further processes are unnecessary. The coagulated material can then be incinerated or disposed safely and cleanly.

Size Comparison



QinPot Liner Bag Box (50Pcs) 1L
Size:W420mm×L310mm×H270mm
Weight:2kg

FITFIX Liner Bag Box (50Pcs) 1L
Size:W710mm×L280mm×H420mm
Weight:7.4kg

The liner can be replaced easily without removing the unit from the wall.

▶ How to setup a liner



(Set the liner in the canister.)



(Close the lid.)



(Connect the suction tube to the liner adaptor port before start suctioning.)

▶ Disposal



(Put the cap on the adaptor port of the liner while suctioning. (It is not necessary to stop suctioning.))



(Stop suction and opened the lid.)



(To remove the liner, put a finger through the hole at the topside of the liner and pull up.)

It's so easy ♪



EC DECLARATION OF CONFORMITY

Manufacturer: DAIKEN MEDICAL CO., LTD.
2-6-2, Ayumino, Izumi-city, Osaka 594-1157, Japan

European Representative: Medical Technology Promedt Consulting GmbH
Altenhofstrasse 80, D-66386 St. Ingbert, Germany

We, DAIKEN MEDICAL CO.LTD, do hereby declare that the medical devices, cited below, conform of the requirements of the Medical Devices Directive "93/42/EEC" and Annex 1.

Name: Coopdech Qin Pot
Product group: Canisters, Aspirator Collection (10-211)
Model(s)/Type(s): CQR10-Y-EU, CQR10-P-EU, CQR10-G-EU, CQR10-B-EU, CQR10-PY-EU, CQR10-PP-EU, CQR10-PG-EU, CQR10-PB-EU, CQR10-SY-EU, CQR10-SP-EU, CQR10-SG-EU, CQR10-SB-EU, CQR-Y-EU, CQR-P-EU, CQR-G-EU, CQR-B-EU, CQD10-H-EU, CQD10-G-EU
Accessories: Pipe Connector
Class: Class I (Annex IX Rule 1)
Applied approach: Annex VII of Medical Devices Directive "93/42/EEC"

Products covered :

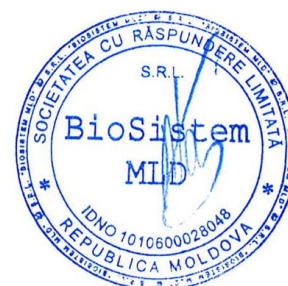
Document Number	title	Number of products
F-EN03-01-00 (QP)	Quality Standard (Product Standard) (Model: QINPOT & QINPOT LINER)	Quality Standard (Product Standard)
SHJS-SB-001	Quality Standard (Product Standard)	Quality Standard (Product Standard)
QQD000-067	COOPDECH Qin Pot Quality records	each Lot Number

Lot Number:

Main Body: XXXXXX (six digits serial number)

Liner: XYMMDDBZ (X: H (without solidifying agent) or G (with solidifying agent), YY-year, MM-month, DD-day, B-Batch No. if necessary, Z-specification if necessary)

This Declaration of conformity is valid in connection with the released document for the respective batch of produced device.



Standards applied:

EN ISO 15223-1:2016

Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements (ISO 15223-1:2016, Corrected version 2017-03)

EN 1041:2008

Information supplied by the manufacturer with medical devices

EN ISO 10079-3:2009

Medical suction equipment – Part 3: Suction equipment powered from a vacuum or pressure source (ISO 10079-3:1999)

EN ISO 13485:2016

Medical devices – Quality management systems – Requirements for regulatory purposes (ISO 13485: 2016)

EN ISO 13485:2016/AC:2018

Medical devices – Quality management systems – Requirements for regulatory purposes

EN ISO 14971:2012

Medical devices – Application of risk management to medical devices (ISO 14971:2007, Corrected version 2007-10-01)

EN 62366:2008

Medical devices – Application of usability engineering to medical devices (IEC 62366:2007)

MEDDEV2.7.1 Rev.3

Clinical Evaluation – A Guide for Manufacturers and Notified Bodies (December 2009)

Modification to the devices without prior approval from the undersigned will render this declaration null and void. This declaration of conformity and the "Technical Documentation" required by the directive will be on file at the manufacturer.

Place of declaration:DAIKEN MEDICAL CO., LTD.
2-6-2, Ayumino, Izumi-city, Osaka 594-1157, Japan**Date of declaration:**

17 Nov 2020

Signature:

(Authorized person)


Hiroaki Takimoto
Quality Assurance Manager