Formularul ofertei (F3.1)

Data depunerii ofertei: 02 12 2019

Digitally signed by Bunic Gheorghe
Date: 2019.12.02 21:49:16 EET
Reason: MoldSign Signature
Location: Moldova

Către:

IMSP Institutul Mamei și Copilului (IMC)

Ericon SRL declară că:

_Ericon SRL se angajează să furnizeze/presteze, în conformitate cu documentele de atribuire şi condiţiile stipulate în specificaţiile tehnice şi preţ, următoarele bunuri şi/sau servicii-

privind achiziționarea Reactivi și consumabile pentru laboratoarele CSRGM

Suma totală a ofertei fără TVA constituie:
100231,66 lei (Una suta mii doua sute treizeci si unu lei 66 bani)

a) Suma totală a ofertei cu TVA constituie: **108250,00 lei** (Una suta opt mii doua sute cincizeci lei 00 bani)
b) Prezenta ofertă va rămîne valabilă pentru perioada de timp specificată în **FDA4.8.**, începînd cu data-limită pentru depunerea ofertei, în conformitate cu **FDA5.2.**, va rămîne obligatorie şi va putea fi acceptată în orice moment pînă la expirarea acestei perioade;
c) În cazul acceptării prezentei oferte, Ericon SRL se angajează să obţină o Garanţie de bună execuţie în conformitate cu **FDA7**, pentru executarea corespunzătoare a contractului de achiziţie publică.
d) Nu sîntem în nici un conflict de interese, în conformitate cu punctul **IPO5.4**.
e) Compania semnatară, afiliaţii sau sucursalele sale, inclusiv fiecare partener sau subcontractor ce fac parte din contract, nu au fost declarate neeligibile în baza prevederilor legislaţiei în vigoare sau a regulamentelor cu incidenţă în domeniul achiziţiilor publice, în conformitate cu punctul **IPO5.5**.

Semnat:_____ L.Ș.
[semnătura persoanei autorizate pentru semnarea ofertei]

Nume:_Bunic Gheorghe

În calitate de: director
Ofertantul: _Ericon SRL

Adresa: Chisinau, Durlesti str.V. Lupu 6

Data: 02 12 2019

ORDIN DE PLATA NR.: 324 TIP.DOC. 1
DATA EMITERII:3 decembrie 2019

PLATITI: 1003-00 LEI: Una Mie Trei lei 00 bani

PLATITOR: (R) ERICON SRL CODUL IBAN:MD60VI222400008100401MDL
CODUL FISCAL :1003600000316 /

PRESTATORUL PLATITOR CODUL BANCII:
B.C."VICTORIABANK"S.A. fil.nr.8 Chisinau :VICBMD2X802:

BENEFICIAR (R) IMSP Institut CODUL IBAN:MD91ML000000000225152859:
ul Mamei si Copilului CODUL FISCAL :1003600151643

PRESTATORUL BENEFICIAR CODUL BANCII:
BC"Moldindconbank"S.A. :MOLDMD2X :

DESTINATIA PLATII:Pentru garantia pentru: TIPUL TRANSFERULUI :
oferta la procedura de achizitie public: NORMAL/URGENT : N:
a nr.oocds-b3wdp1-MD-1572532179763

L.S.

CODUL TRANZACTIEI:001:
DATA PRIMIRII:03/12/2019 : _____
DATA EXECUTARII: : _____
: SEMNATURILE
: EMITENTULUI
_____ :
SEMNATURA PRESTATORULUI :

MOTIVUL REFUZULUI : L.S.

Rechizite bancare ERICON SRL

INDNO: 1003600000316
IBAN: MD60VI222400008100401MDL
BCA"Victoriabank"filiala 8 VICBMD2X802
TVA: 0302060

Str. V.Lupu 6, Durlesti
Chişinău,MOLDOVA
www.ericon.md
Fax: +373 22 52 01 08
Phone: +373 79 41 00 42
e-mail: bunicgh@gmail.com

MED CERT®

Certification. Medical Only.

Certificate

The Certification Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith confirms that the company

Gynemed GmbH & Co. KG
Lübecker Str. 9
23738 Lensahn
Germany

has introduced, applies and maintains a Quality Management System in the area of:

Development, manufacture final inspection and distribution of Products for reproductive medicine

Distribution of Products for surgical and conservative gynaecology

The compliance of the Quality Management System with the requirements of the below mentioned standard was verified by an audit:

EN ISO 13485:2016

The license of certification is subject to surveillance by MEDCERT.

This certificate is valid from 28 July 2019 until 27 July 2022

Report No.:6587FS11F
Process No.:QS – 6587
Certificate No.: 6587GB445190717

Hamburg, 17. Juli 2019

MEDCERT Certification Body
(Dr. Andreas Schich)

DAkkS
Deutsche Akkreditierungsstelle
D-ZM-19630-04-00

EC CERTIFICATE

Number: 2154875CE01

Full Quality Assurance System

Directive 93/42/EEC on Medical devices, Annex II excluding (4)
(Devices in Class IIa, IIb or III)

Manufacturer:

Gynemed GmbH & Co. KG

Lubecker Straße 9
23738 Lensahn
Germany

For the product category(ies)

Media for the washing, cryopreservation and in vitro culture of human embryos, ova and spermatozoa for use in In Vitro Fertilization (IVF) / IntraCytoplasmatic Sperm Injection (ICSI), Assisted Reproductive Techniques (ART) and Embryo Transfer (ET).

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EC-Directive which apply to them:

0344

Documents, that form the basis of this certificate:

Certification Notice 2154875CN, initially dated 27 November 2012
Addendum, initially dated 27 November 2012

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Council Directive 93/42/EEC of June 14, 1993 concerning Medical devices, including all subsequent amendments. The manufacturer has implemented a quality assurance system for design, manufacture and final inspection for the above mentioned product category in accordance to the provisions of Annex II of Council Directive 93/42/EEC of June 14, 1993 and is subject to periodical surveillance. For placing on the market of Class III devices an additional EC design examination certificate according to Annex II (4) is mandatory. The necessary information related to the quality management system of the manufacturer, including facilities and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: 1 October 2023
Issued for the first time: 27 November 2012
Reissued: 12 November 2018

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 F +31 88 96 83100 www.dekra-certification.com Company registration 09085396

ADDENDUM

Belonging to certificate: 2154875CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Media for the washing, cryopreservation and in vitro culture of human embryos, ova and spermatozoa for use in In Vitro Fertilization (IVF) / IntraCytoplasmatic Sperm Injection (ICSI), Assisted Reproductive Techniques (ART) and Embryo Transfer (ET).

Issued to:

Gynemed GmbH & Co. KG
Lubecker Stra?e 9
23738 Lensahn
Germany

This certificate covers the following product(s):

Media with gentamicin (Class III)
Media with human albumin (Class III)
Media with human albumin and gentamicin (Class III)
Media with specific medium supplement (Class III)

Initial date: 27 November 2012
Revision date: 13 October 2016

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
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Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 106138

Manufacturer:

Marflow AG
Soodstrasse 57
8134 Adliswil, Zurich
SWITZERLAND

Product Category(ies):

Class IIb
Double J stent & set

Class IIa
PCN catheter & set
Ureteral catheter
Malecot catheter
Re-entry malecot catheter
Suprapubic catheter
Braided shaft catheter
Dual lumen catheter
Facial dilator
Amplatz dilator & set
Ureteral dilator & set
Ureteral balloon dilator
Double J stent & set
Mono J stent
Endopyelotomy stent
Guidewire
IP Needle
Chiba needle
Stone basket
Perk basket

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: IND20190101

Valid from: 2019-10-22
Valid until: 2024-10-21

Date, [#ISU_DT#]

Stefan Preiß
Head of Certification/Notified Body

Draft From CBW 2.0 - Production System (Build - 20190829.1)

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 106138

Facility(ies):

Marflow AG
Soodstrasse 57, 8134 Adliswil, Zurich, SWITZERLAND

-/-

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bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in class I in sterile conditions, sterilized systems or procedure packs)

No. G1S 106138

Manufacturer:

Marflow AG
Soodstrasse 57
8134 Adliswil, Zurich
SWITZERLAND

Product Category(ies):

Class Is
Urine bag connector
Penile clamp
Evacuator
IUI catheter without syringe

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex II. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: IND20190101

Valid from: 2019-10-22
Valid until: 2024-10-21

Date, [#ISU_DT#]

Stefan Preiß
Head of Certification/Notified Body

Draft From CBW 2.0 - Production System (Build - 20190829.1)

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in class I in sterile conditions, sterilized systems or procedure packs)

No. G1S 106138

Facility(ies):

Marflow AG
Soodstrasse 57, 8134 Adliswil, Zurich, SWITZERLAND

-/-

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TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-11321-01-00

Product Service

Certificate

No. Q5 106138

Holder of Certificate: **Marflow AG**
Soodstrasse 57
8134 Adliswil, Zurich
SWITZERLAND

Facility(ies): Marflow AG
Soodstrasse 57, 8134 Adliswil, Zurich, SWITZERLAND

Certification Mark:

Scope of Certificate: **The Design, Manufacture and Supply of Medical Disposables, Surgical Tools, Equipment & Accessories in the field of Urology, Gastroenterology, Radiology, Gynaecology & Cardiology.**

Applied Standard(s): EN ISO 13485:2016
Medical devices - Quality management systems - Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: IND20190101

Valid from: 2019-10-22
Valid until: 2022-10-21

Date, [#ISU_DT#]

Stefan Preiß
Head of Certification/Notified Body

Draft From CBW 2.0 - Production System (Build - 20190829.1)

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