Declaration Ref No: DC11-0051

Atlas Medical

# CE Declaration of Conformity

## According to Annex III of the IVD Directive 98/79/EC

We,

**Atlas Medical**


Head office: Ludwig-Erhard-Ring 3
D-15827 Blankenfelde-Mahlow
Tel: +49 - 33708 – 3550 30
Email: info@atlas-medical.com

Middle East Site: Sahab Free Zone Area, P. O. Box 212555, Amman, Jordan.
Tel.: +962 6 4026468
Fax: +962 6 4022588
Email: info@atlas-medical.com


Declare our responsibility that the following product:

**See Attached list**

- Comply with all essential requirements (AnnexI) of the IVD Directive 98/79/EC. This compliance has been properly documented and covers the items listed in Annex I of the IVD Directive.
- This product is produced under Atlas quality system (ISO13485:2016) issued by Lloyd's Register Quality Assurance.
- Comply with the essential requirements of following standards (EN 18113-1, -2,-4:2011, EN ISO 15223:2016 , EN ISO 23640:2015, EN ISO 14971:2012, ISO 2859/1:1999, EN ISO 13612:2002, EN ISO 13641:2002.

And

Intended for In-Vitro Professional use only.

"SANMEDICO"

**Manufacturer**
**Atlas Medical**
**Ludwig-Erhard-Ring 3**
**D-15827 Blankenfelde-Mahlow**

| Atlas Medical | Issue date | Date of review | Management approval | MRXDO10F.10 08.02.2011 |
|---|---|---|---|---|
| | December.2011 | 26.11.2019 | | |

Digitally signed by Monastîrschii Viorica
Date: 2020.11.04 16:52:00 EET
Reason: MoldSign Signature
Location: Moldova

# СЕРТИФІКАТ

CERTIFICATE * CERTIFICAT * ZERTIFIKAT * СЕРТИФИКАТ * CERTIFICADO

ОРГАН СЕРТИФІКАЦІЇ СИСТЕМ УПРАВЛІННЯ
ДП «УКРМЕТРТЕСТСТАНДАРТ»
ЗАСВІДЧУЄ, ЩО

## СИСТЕМА УПРАВЛІННЯ ЯКІСТЮ

## ТОВАРИСТВА З ОБМЕЖЕНОЮ ВІДПОВІДАЛЬНІСТЮ «ВІТРОТЕСТ БІОРЕАГЕНТ»

Юридична адреса: вул. Бойчука, 18-Б, кв. 56, м. Київ, 01103, Україна
Адреса виробництва: вул. Курортна, 11, м. Київ, 04075, Україна

код ЄДРПОУ 42149820

стосовно
розроблення та виробництва тест-систем імуноферментних

## ВІДПОВІДАЄ ВИМОГАМ ДСТУ EN ISO 13485:2018 (EN ISO 13485:2016, IDT; ISO 13485:2016, IDT)

Сертифікат № UA.C.378–19 в Реєстрі Органу сертифікації
зареєстрований " 25 " листопада 2019 року
чинний до " 24 " листопада 2022 року

**Заступник керівника Органу сертифікації** **В.Д. Ример**

ДЕРЖАВНЕ ПІДПРИЄМСТВО «ВСЕУКРАЇНСЬКИЙ ДЕРЖАВНИЙ НАУКОВО-ВИРОБНИЧИЙ ЦЕНТР СТАНДАРТИЗАЦІЇ, МЕТРОЛОГІЇ, СЕРТИФІКАЦІЇ ТА ЗАХИСТУ ПРАВ СПОЖИВАЧІВ»
(ДП «УКРМЕТРТЕСТСТАНДАРТ»)
вул. Метрологічна, 4, м. Київ, 03143, Україна, тел./факс +38 044 452-67-38
Атестат акредитації НААУ № 8О020

№ 8О020
ДСТУ EN ISO/IEC 17021-1

Чинність сертифікату можна перевірити на сайті www.certsystems.kiev.ua в розділі «Послуги / Сертифікація систем управління»

CE

# KONFORMITÄTSERKLÄRUNG

# DECLARATION OF CONFORMITY

Doc#022/06-2014

Wir / *We*

**TECO Medical Instruments Production and Trading GmbH**
Name des Herstellers / Manufacturer's *name*
**Dieselstrasse 1, D-84088 Neufahrn NB**
Anschrift / *Address*

erklären in alleiniger Verantwortung, dass unsere im beigefügten Anhang (2 Seiten) spezifizierten Produkte wie folgt gemäß der Richtlinie für In-vitro-Diagnostika Medizinprodukte 98/79/EC klassifiziert sind:
*declare under our own responsibility, that our products specified in the enclosed addendum (2 pages) classified as follows according to the directive on in vitro diagnostic medical devices 98/79/EC:*

## Übrige Produkte – Reagenzien für In-vitro-Diagnostika
## Other Products – Reagents for in vitro diagnostic

| | |
|---|---|
| Allen anwendbaren Anforderungen der folgenden Richtlinien entsprechen: | *Meet all applicable requirements of:* |
| Richtlinie 98/79/EG über In-vitro-Diagnostika klassifiziert gemäß Artikel 9 als "alle anderen Produkte" | *Directive 98/79/EC on in-vitro-diagnostic medical devices classified according to article 9 as „all other products"* |

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| | |
|---|---|
| Das QM-System des Herstellers ist zertifiziert nach: | *The QM-system of the manufacturer is certified for:* |
| **EN ISO 13485:2016** | ***EN ISO 13485:2016*** |
| Die vorstehende Konformitätserklärung ist gültig für alle Chargen dieser Produkte, die nach dem Datum der Unterzeichnung in Verkehr gebracht wurden. | *The above mentioned declaration of conformity is valid for all lots of this product, which are distributed after the date of signature.* |
| Konformitätsbewertungsverfahren: | *Conformity assessment procedure:* |
| Gemäß Anhang III der Richtlinie 98/79/EG | *According to Annex III of Directive 98/79/EC* |

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Ort und Datum der Unterzeichnung: Neufahrn, 26.03.2019
Place and date of issue: Neufahrn, March 26, 2019

Christian Hötzl
General Manager

Medical Instruments TECO Production+Trading GmbH

CE

# KONFORMITÄTSERKLÄRUNG

# DECLARATION OF CONFORMITY

Doc#001/35-2014

Wir / *We*

**TECO Medical Instruments Production and Trading GmbH**
Name des Herstellers / Manufacturer's *name*
**Dieselstrasse 1, 84088 Neufahrn, Germany**
Anschrift / *Address*

erklären in alleiniger Verantwortung, dass die Produkte – IVD-Blutgerinnungsmessgeräte,
*declare under our own responsibility, that the products – IVD Coagulation analyzers*

## Coatron X Eco, Pro, Top

Bezeichnung, Typ oder Modellname / *name, type or model*

| | |
|---|---|
| allen anwendbaren Anforderungen der folgenden Richtlinien entsprechen: | *meet all applicable requirements of:* |
| 1. Richtlinie 98/79/EG über In-vitro Diagnostika klassifiziert gemäß Artikel 9 als: "alle anderen Produkte" | *1. Directive 98/79/EC on In-vitro diagnostic medical devices classified according to article 9 as: "all other products"* |
| 2. Richtlinie 2014/30/EU über Elektromagnetische Verträglichkeit | *2. Directive 2014/30/EU on electromagnetic Compatibility* |
| 3. Richtlinie 2011/65/EU RoHS II | *3. Directive 2011/65/EU RoHS II* |
| Das QM-System des Herstellers ist zertifiziert nach: | *The QM-system of the manufacturer is certified for:* |
| **EN ISO 13485:2016** | ***EN ISO 13485:2016*** |
| Diese Erklärung bescheinigt die Übereinstimmung mit den genannten Harmonisierungsrechtsvorschriften, beinhaltet jedoch keine Zusicherung von Eigenschaften. | *This declaration attests the accordance with the mentioned harmonization rule but does not include a warranty of properties.* |
| Konformitätsbewertungsverfahren: | *Conformity assessment procedure:* |
| Gemäß Anhang III der Richtlinie 98/79/EG | *According to Annex III of Directive 98/79/EC* |

Ort und Datum der Unterzeichnung:
Place and date of issue:

Neufahrn, 27.03.2019
Neufahrn, March 27, 2019

Christian Hötzl
General Manager

Medical Instruments
TECO
Production+Trading GmbH

LumiQuick Diagnostics, Inc.
2946 Scott Blvd., Santa Clara, CA 95054, USA

Tel: 408-855-0061
Fax: 408-855-0063
E-mail: info@LumiQuick.com
Web: www.lumiquick.com

# Declaration of Conformity

| PRODUCT IDENTIFICATION | |
|---|---|
| **Product name** | **Model/number** |
| Chikungunya Test Devices<br>QuickProfile Chikungunya IgG/IgM Combo Test Card | 71031 |

| MANUFACTURER | | |
|---|---|---|
| **Name of company** | **Address** | **Representative** |
| LumiQuick Diagnostics, Inc. | 2946 Scott Blvd.<br>Santa Clara, CA<br>95054<br>USA | Jeff Wang |

| AUTHORIZED REPRESENTATIVE | | |
|---|---|---|
| **Name of company** | **Address** | **Telephone/email** |
| Emergo Europe | Prinsessegracht 20<br>2514 AP<br>The Hague, Netherlands | +31.70.345.8570 - phone<br>+31.70.346.7299 - fax<br>europe@emergogroup.com |

| CONFORMITY ASSESSMENT | | |
|---|---|---|
| **Device classification** | **Route to compliance** | **Standards applied** |
| Class: Self-Certify | Annex III of IVDD 98/79/EC Council Directive | ISO 13485:2003 |

LumiQuick Diagnostics, Inc. declares that the above mentioned products meet the provision of the Council Directive 98/79/EC for In Vitro Diagnostic Medical Devices and Directive 98/79/EC as transposed in the national laws of the Member States.

**COMPANY REPRESENTATIVE:** Jeff Wang

**TITLE:** Quality Systems Manager **SIGNATURE:**

**DATE:** 28/04/2017

EC_Declaration_Letter_Emergo_E2R0_NewAddress

# Declaration of Conformity

according to Directive 98/79/EC, on in vitro diagnostic medical devices

| | | | |
|---|---|---|---|
| **Maker** (Name, Address) | **Getein Biotech, Inc.** No. 9 Bofu Road, Luhe District, Nanjing, 211505, China | | |
| **Authorized Representative** (Name, Address) | **Lotus NL B.V.** Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands. | | |
| **Medical device** | Description | : | FIA8000 Quantitative Immunoassay Analyzer<br>FIA8600 Quantitative Immunoassay Analyzer<br>Cardiac Troponin I Fast Test Kit<br>One Step Test for cTnI (Colloidal Gold)<br>cTnI Rapid Test (Colloidal Gold Assay)<br>One Step Test for NT-proBNP (Colloidal Gold)<br>One Step Test for NT-proBNP/cTnI (Colloidal Gold)<br>One Step Test for CK-MB/cTnI/Myo (Colloidal Gold)<br>One Step Test for hs-CRP+CRP (Colloidal Gold)<br>One Step Test for D-Dimer (Colloidal Gold)<br>One Step Test for PCT (Colloidal Gold)<br>One Step Test for β2-MG (Colloidal Gold)<br>One Step Test for mAlb (Colloidal Gold)<br>One Step Test for NGAL (Colloidal Gold)<br>One Step Test for CysC (Colloidal Gold)<br>One Step Test for HCG+β (Colloidal Gold)<br>One Step Test for HbA1c (Colloidal Gold)<br>One Step Test for PCT/CRP (Colloidal Gold)<br>One Step Test for CK-MB/cTnI/H-FABP (Colloidal Gold)<br>One Step Test for H-FABP (Colloidal Gold)<br>One Step Test for CK-MB/cTnI (Colloidal Gold)<br>One Step Test for CK-MB (Colloidal Gold)<br>One Step Test for TSH (Colloidal Gold)<br>One Step Test for T4/T3 (Colloidal Gold)<br>One Step Test for T3 (Colloidal Gold)<br>One Step Test for T4 (Colloidal Gold)<br>One Step Test for 25-OH-VD (Colloidal Gold)<br>One Step Test for FOB (Colloidal Gold)<br>One Step Test for *H. pylori* (Colloidal Gold)<br>One Step Test for SAA (Colloidal Gold)<br>Getein1100 Immunofluorescence Quantitative Analyzer<br>Getein1600 Immunofluorescence Quantitative Analyzer<br>Getein1180 Immunofluorescence Quantitative Analyzer<br>Getein1200 Immunofluorescence Quantitative Analyzer<br>Cardiac Troponin I Fast Test Kit (Immunofluorescence Assay)<br>NT-proBNP Fast Test Kit (Immunofluorescence Assay)<br>hs-CRP+CRP Fast Test Kit (Immunofluorescence Assay)<br>NT-proBNP/cTnI Fast Test Kit (Immunofluorescence Assay)<br>CK-MB/cTnI/Myo Fast Test Kit (Immunofluorescence Assay)<br>D-Dimer Fast Test Kit (Immunofluorescence Assay) |

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-245.10.07

TÜV SÜD Product Service

# EC Certificate

Full Quality Assurance System
Directive 98/79/EC on In Vitro Diagnostic Medical Devices (IVDD), Annex IV excluding (4, 6)
(List A and B and devices for self-testing)

## No. V1 104507 0003 Rev. 01

**Manufacturer:** ACON Laboratories, Inc.
5850 Oberlin Drive, #340
San Diego CA 92121
USA

**Product Category(ies):** In Vitro diagnostics for the detection of human infections and tumor markers, blood glucose measuring self-testing systems, self-testing devices for clinical chemistry, hematology and pregnancy and ovulation

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 families in accordance with IVDD Annex IV. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of List A devices an additional Annex IV (4) certificate is mandatory. See also notes overleaf.

**Report no.:** SH1974310

**Valid from:** 2019-10-24
**Valid until:** 2022-09-12

**Date,** 2019-10-24

Stefan Preiß
Head of Certification/Notified Body

REPUBLIC MOLDOVA, mun. CHISINAU S.R.L. "SANMEDICO" IDNO 1003 SOCIETATEA CU RASPUNDERE LIMITATĂ


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

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

## DECLARATION OF CONFORMITY

1) Manufacturer *(Name, department)*: **HiMedia Laboratories Pvt. Ltd.**

**Address: 23 Vadhani Industrial Estate,LBS Marg, Mumbai - 86, MS, India**

and

2) European authorized representative: **CEpartner4U BV,**

**Address: ESDOORNLAAN 13, 3951DB MAARN, THE NETHERLANDS;**
(on product labels printed as:
CEpartner4U , ESDOORNLAAN 13, 3951DB MAARN, THE NETHERLANDS. www.cepartner4u.eu)

3) Product(s) *(groupnames /.)*:

| Group | Group name | NL registration no. | No. |
|---|---|---|---|
| DCM&S | Dehydrated Culture Media & Supplements | NL-CA002-2013-26442 | 1 |
| RPM | Ready Prepared Media<br>Subgroups: Ready Prepared Plates, Ready Prepared Liquid & Solid Medium, Ready Prepared Slants,Ready Prepared Dual Media, HiDip Slides, HiSafe Blood Culturing System, Transport Medium w/ swabs, Viral Transport Medium w/ swabs, L.J.Medium Slants & Kits, Biochemical Kits for Mycobacteria, UTI Diagnostic Kits, Biochemical Identification Kits | NL-CA002-2013-26448 | 2 |
| ESK | Epidemiological Screening Kit:<br>Subgroups: Hi Aureus Confirmation Kits | NL-CA002-2012-24117 | 3 |
| ASS | Antimicrobial Susceptibility Systems<br>Subgroups: Sensitivity Discs-Single & Multi Discs<br>MIC Strips: HiComb Strips & Ezy MIC Strips | NL-CA002-2013-26444 | 4 |
| BDA | Bacteriological Differentiation Aids<br>Subgroups: Readymade Stains, Indicators & Reagents in liquid, Differentiation Discs & Strips, HiDtect Rapid Identification Discs | NL-CA002-2013-26445 | 5 |
| CCM | Cell Culture Media<br>Subgroups: Karyotyping Media, Stem Cell Differentiation Media & Supplements, Stem Cell Freezing Medium, Stem Cell Differentiation Kits ,Viral Transport Medium, Balanced Salt Solutions, Antibiotic solutions, Animal Cell Culture Medium Liquid | NL-CA002-2013-26446 | 6 |
| MBP | Molecular Biology Products<br>Subgroups: DNA & RNA Isolation Kits, Latex Agglutination Kits, Haematology Kits, Density gradient Separation Medium, PCR Kits | NL-CA002-2013-26447 | 7 |

*type and model numbers: see appendix*

4) The product(s) described above is in conformity with:

| Title | Document No. |
|---|---|
| *In vitro* Diagnostic Medical Devices Directive | 98/79/EC |

5) Additional information *(Conformity procedure, Notified Body, CE certificate, Registration nr., etc.)*:

Conformity assessment procedure for CE marking: *In vitro* Diagnostic Medical Device Directive, Annex III

Mumbai, India; 2019-04-22

*(Place & date of issue (yyyy-mm-dd))*

Dr. G.M.Warke , Managing Director

*(name; function and signature of manufacturer)*

Lloyd's Register
LRQA

# EC CERTIFICATE – FULL QUALITY ASSURANCE SYSTEM

**In accordance with the requirements of the In Vitro Diagnostic Medical Devices Directive 98/79/EC and the Medical Devices Regulations 2002, UK Statutory Instrument 2002 No. 618**

This is to certify that the Quality Management System of:

**Mast Group Limited**
**Mast House, Derby Road,**
**Bootle, Merseyside**
**United Kingdom**

has been assessed against the requirements of Annex IV of the In Vitro Diagnostic Medical Devices Directive 98/79/EC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown on the attached certificate schedule.

Approval is subject to the maintenance of the quality system in accordance with the requirements of the above Directive and Regulations.

Authorisation is hereby given to use the LRQA Notified Body Registration Number in accordance with the requirements of the specified Directives/Regulations in relation to the products as identified above.

| | |
|---|---|
| Certificate No: | LRQ 0932114/C |
| Original Approval: | 25 May 2004 |
| Current Certificate: | 1 June 2018 |
| Certificate Expiry: | 31 May 2021 |

LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited

1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom.

CEpartner4U

Certificate number: 2017-IVD/193

# Certificate of CE-Notification

This is to certify that, in accordance with the *In Vitro* Diagnostic Medical Device Directive 98/79/EC, **CEpartner4U BV** agrees to perform all duties and responsibilities as the Authorized Representative for

## CJSC EKOlab

1 Budennogo Str., Elektrogorsk, Moscow region, 142530, Russia

as stipulated and demanded by the aforementioned Directive. The Dutch Competent Authorities have accepted the manufacturer's medical device registrations by CEpartner4U as listed on the product list attached to the manufacturer's Declaration of Conformity:

## Device group: Rabbit plasma

**IVD devices were registered under number:**

**Registration number Rabbit plasma: NL-CA002-2017-43242**

**with Dutch Competent Authorities as a consequently this IVD devices were entered in EUDAMED by Dutch Competent Authorities**

The manufacturer has provided CEpartner4U with all necessary documentation, together with an appropriate Declaration of Conformity that the IVD medical devices fulfil the essential requirements of Directive 98/79/EC.

2017-12-18

Olga Teirlinck
Consultant CEpartner4U BV

REPUBLICA MOLDOVA, mun. CHIŞINĂU S.R.L. "SANMEDICO" IDNO 1003602008154 SOCIETATEA CU RĂSPUNDERE LIMITATĂ

CEpartner4U
Esdoornlaan13
3951 DB Maarn NL
tel: +31 (0)343 442 524
www.cepartner4u.nl

SYNTESYS®

ICIM ISO9011 2008 Cert. N 6574/0 · CISQ · IQNet

SYNTESYS S.A.S. DI RINALDO R. & C.
VIA G. GALILEI, 10/3
35037 Z.I. SELVE DI TEOLO (PD)
TEL. +39 049 9903866 R.A. FAX +39 049 9903867
COD.FISCALE P.IVA N.REG.IMP. PADOVA 03573950288
E-MAIL INFO@SYNTESYS.IT - WEB WWW.SYNTESYS.IT

# DICHIARAZIONE DI CONFORMITA'
*Conformity declaration*

CE

Il sottoscritto, Rinaldo Ruggero legale rappresentante della ditta:
*The undersigned, Rinaldo Ruggero legal representative of the company:*

produttore/*manufacturer*

SYNTESYS S.a.s. di Rinaldo R. & C.

indirizzo/*address*

Via G. Galilei, 10/3 35037 Zona Industriale SELVE DI TEOLO (PADOVA) ITALY

O rappresentante il mandatario autorizzato entro la Unione Europea
*or representing the authorized mandatary within the European Community*

Mandatario autorizzato/*authorized mandatary*

indirizzo/*address*

Dichiara sotto la propria responsabilità che il prodotto/*declares under his own responsability that the product:*

| | |
|---|---|
| Denominazione/*Description* | Padella per ammalati, urinali uomo e donna, speculum vaginali, tamponcini cotonati, tamponi sterili in provetta, tamponi sterili con terreno Amies e Stuart in provetta/ *Bed pan, Urinal's man and woman, Vaginal speculum, Cotton swab, Sterile swab in test tube, Sterile swab with medium Amies or Stuart in test tube* |
| Materiale/*Material* | Polipropilene, Polietilene, Legno/ *Polypropylene, Polyethylene, Wood* |

È conforme alle disposizioni della direttiva 93/42/CE e smi, concernente i dispositivi medici ed al Decreto Legislativo di recepimento con D.lgs. del 24/02/1997 n° 46/97 e soddisfa a tutti i requisiti specificati.
Il dispositivo è stato classificato appartenente alla classe I° secondo i criteri stabiliti in base a quanto previsto dall'Art. 9 ed allegato IX della direttiva sopra citata /*It meets the EC Directive 93/42 about Medical Device, specifications established by the Italian law n 46/97, dated 24th February 1997. The device was classified as belonging to the 1st class, according to the specifications of the established by the art.9, IX enclosure of the above mentioned directive.*

Dichiara inoltre che la documentazione tecnica di supporto alla presente dichiarazione di conformità è conservata presso gli uffici dell'azienda e sarà posta alla disposizione di chi la richiede/ *declares that all technical documents attached to this conformity statment are filed in our company and can be consulted by any authorized body on demand.*

Data 07.01.2016
Issued on January 7th 2016

REPUBLICA MOLDOVA, mun. CHISINAU "SANMEDICO" S.R.L. IDNO 1007600208154 SOCIETATEA CU RASPUNDERE LIMITATA

SYNTESYS S.A.S.
Il legale rappresentante
Rinaldo Ruggero

SYNTESYS

ISO9001:2008
Cert. N.6574/0

SYNTESYS S.A.S. DI RINALDO R. & C.
VIA G. GALILEI, 10/3
35037 Z.I. SELVE DI TEOLO (PD)
TEL. +39 049 9903866 R.A. FAX +39 049 9903867
COD.FISCALE P.IVA N.REG.IMP. PADOVA 03573950288
E-MAIL INFO@SYNTESYS.IT - WEB WWW.SYNTESYS.IT

# DICHIARAZIONE DI CONFORMITA'
## *Conformity declaration*

CE

Il sottoscritto, Rinaldo Ruggero legale rappresentante della ditta:
*The undersigned, Rinaldo Ruggero legal representative of the company:*

produttore/*manufacturer*

SYNTESYS S.a.s. di Rinaldo Ruggero & C.

indirizzo/*address*

Via G. Galilei, 10/3 35037 Zona Industriale SELVE DI TEOLO (PADOVA) ITALY

*O rappresentante il mandatario autorizzato entro la Unione Europea* or representing the authorized mandatary within the European Community

Mandatario autorizzato/*authorized mandatary*

indirizzo/*address*

Dichiara sotto la propria responsabilità che il prodotto/*declares under his own responsability that the product:*

| | |
|---|---|
| Denominazione degli articoli prodotti/*Description of Manufacturer* | Contenitori per urina, contenitori per feci, contenitori universali, Pipette Pasteur, Piastre di Petri, Anse Sterili per batteriologia, Aste a "L", Puntali Eppendorf gialli e blue, cuvette per spettrofotometro, tazzine per campionamento siero, bacchette per distacco ed estrazione del coagulo, pinzette in polistirolo monouso, provette monouso in plastica, tappi alettati per provette diam. 12 mm e 16mm, provette con granuli ed acceleratore, provette sottovuoto per prelievo, Sistema SEDIPLAST, Microprovette, Portavetrini, Vetrini precolorati, Portaprovette, supporti per microprovette, bottiglie per raccolta urine.<br><br>*Urine container, faeces container, universal container, Pasteur pipette, Petri dishes, Sterile loops, Sterile loops open "L", Eppendorf tips yellow and blue, cuvettes for spectrophotometer, samples cups, Rod to detach clot, disposable forceps, Disposable plastic tubes, winged stoppers for tubes diam. 12mm & 16mm, Test tube with granules and clot activator, vacuum test tube, SEDIPLAST system, micro test tubes, Slides Mailer, "TESTSIMPLETS" slides rack for test tubes, rack for micro test tubes, Bottles for urine collection.* |

REPUBLICA MOLDOVA, mun. CHISINAU, SOCIETATEA CU RĂSPUNDERE LIMITATĂ "SANMEDICO" S.R.L. IDNO 1003602006154