

Anexa nr. 7  
la Documentația standard nr. \_\_\_\_\_  
din “ \_\_\_\_\_ ” \_\_\_\_\_ 20 \_\_\_\_\_

## CERERE DE PARTICIPARE

Către: **IMSP Spitalul Clinic Republican „Timofei Moșneaga”,**  
**(mun.Chișinău, str.N.Testemițanu 29)**

**Stimați domni,**

Ca urmare a anunțului de participare apărut în Buletinul achizițiilor publice nr. 53 din 07.07.2023, privind aplicarea procedurii pentru atribuirea contractului de achiziționare a **Necesarului de piese de schimb pentru dispozitive medicale pentru anul 2023 etapa II**, noi Medist Grup SRL, am luat cunoștință de condițiile și de cerințele expuse în documentația de atribuire și exprimăm prin prezenta interesul de a participa, în calitate de ofertant/candidat, neavând obiecții la documentația de atribuire.

Data completării 25.07.2023

Cu stimă,  
Ofertant/candidat  
**Gabriela-Cristina Anghel**

Tip.doc. 1

PLATITI:2302-00

LEI: Doua Mii Trei Sute Doi, 00

CODUL IBAN:MD57VI022242600000269MDL:

CODUL FISCAL:1018600004516

B.C. VictoriaBank S.A. s.26 Chisinau

BENEFICIAR:(R) IMSP Spitalul Clinic Repub CODUL IBAN:MD57M02251ASV96476607100:  
 lican Timofei Mosneaga CODUL FISCAL:1003600150783 :

OTP Bank

DESTINATIA PLATII: Garantia pentru oferta la LP nr :  
 . 21084566 din 12.07.2023. :

NORMAL/URGENT:NO

L.S.

CODUL TRANZACTIEI:001

DATA PRIMIRII:

DATA EXECUTARII:

SEMNATURILE

EMITENTULUI

SEMNATURA PRESTATORULUI

### MOTIVUL REFUZULUI

10:56:02 25 JUL 2023

Semnatura electronica:

ZbrBv2bGnqjQcjgJdE1zCwD5zrv7JK4VUgCF23wVI6oPiicZ4T2k6PKtBP/QJXq+FShu845tUNnmJmW2kaV/EdAiRzv+2N4XgaxpHE+ED+FHwmpwWRmgob0rzlyZbGVX08zavNVvCdqhRguG7XhYX7jeeyFOPI50LxVEA7p+k+jMK8o+Ui1rHCEQ7BBnk8AeISi+fLx10tWrH267qea4Gg90C35v0gZdMAq+WuH0p2vyDEkGYP98j5L+2e6pL/dz0ErQVez9v090c5VU9aidkrSWts4Yc2hvesjBYftq1V/DFGTAY3Y2z0Kjv411UsGUsPlug0p1y/HjNMl2jPvtw==



**I.P. "AGENȚIA SERVICII PUBLICE"**

Departamentul înregistrare și licențiere a unităților de drept

**Extras  
din Registrul de stat al persoanelor juridice  
nr. 108035 din 09.02.2023**



Denumirea completă: **Societatea cu Răspundere Limitată "MEDIST GRUP"**

Denumirea prescurtată: **"MEDIST GRUP" S.R.L.**

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

Numărul de identificare de stat și codul fiscal: **1018600004516**

Data înregistrării de stat: **02.02.2018**

Sediu: **MD-2012, strada Mitropolit Gavriil Bănulescu-Bodoni 25, ap. 33, mun. CHIȘINĂU**

**Republica Moldova**

Genurile de activitate:

- 1. Comerț cu ridicata al produselor farmaceutice;**
- 2. Comerț cu ridicata nespecializat;**
- 3. Repararea echipamentelor electronice și optice;**
- 4. Activități de testare și analize tehnice;**
- 5. Comerț cu amănuntul al articolelor medicale și ortopedice, în magazine specializate;**

Capitalul social: **373026 Lei**

Administrator: **ANGHEL GABRIELA-CRISTINA**

Asociați:

- 1. MEDIST IMAGING & P.O.C. S.R.L., partea socială 6244 Euro, ce constituie 33.0004%**
- 2. MEDIST LIFE SCIENCE S.R.L., partea socială 6244 Euro, ce constituie 33.0004%**
- 3. MEDIST S.R.L., partea socială 6433 Euro, ce constituie 33.9993%**

Beneficiari efectivi: **MANOLE IONEL, KLUMPNER CATALINA ANA, VLĂDESCU CARMEN, VLĂDESCU SEBASTIAN-ALEXANDRU**

Prezentul extras este eliberat în temeiul art. 34 al Legii nr.220/2007 privind înregistrarea de stat a persoanelor juridice și a întreprinzătorilor individuali și confirmă datele din Registrul de stat la data de 09.02.2023

Specialist coordonator

**Victoria Burcovschi**

tel. 022-207862

**DECLARAȚIE  
privind valabilitatea ofertei**

**Către IMSP Spitalul Clinic Republican „Timofei Moșneaga”,  
(mun.Chișinău, str.N.Testemițanu 29)**

**Stimați domni,**

Ne angajăm să menținem oferta valabilă, **privind achiziționarea Necesarului de piese de schimb pentru dispozitivele medicale pentru anul 2023 etapa II**, prin procedura de achiziție – **Licitație deschisă**, pentru o durată de **90 zile** (nouăzeci zile), respectiv până la data de 30.11.2023 și ea va rămâne obligatorie pentru noi și poate fi acceptată oricând înainte de expirarea perioadei de valabilitate.

Data completării 25.07.2023

Cu stimă,  
Ofertant/candidat  
Gabriela-Cristina Anghel

## DECLARAȚIE

Subsemnata Gabriela Anghel, reprezentant împuternicit al MEDIST GRUP S.R.L, cu sediul în mun. Chișinău, str. M.G. Bănulescu-Bodoni 25, Oficiul 33, declar pe propria răspundere că:

- livrarea produselor la destinatar va avea loc cu respectarea condițiilor de păstrare și transportare.
- Termenul de garanție va fi conform specificației tehnice,
- Seturile vor fi livrate în ambalaj original, securizat, marcat și etichetat de producător, fără preambalare; Date de identitate (denumirea, numărul lotului, seria, termenii de valabilitate, condițiile de păstrare) ale produsului indicate pe ambalaj vor coincide în mod obligatoriu cu cele de pe etichetele componentelor incluse în set. Instrucțiunile privind modul de utilizare sunt prezentate în limba de stat sau limba rusă.
- În caz de solicitare, mostrele vor fi prezentate în 5 zile.

Data: 25.07.2023

**MEDIST GRUP S.R.L.**  
**DIRECTOR**  
**GABRIELA ANGHEL**

## EU DECLARATION OF CONFORMITY

We, **Leica Biosystems Nussloch GmbH, Heidelberger Str. 17-19, 69226 Nussloch, Germany**

declare on our own responsibility, that the device

### **Leica IP C**

complies with

- Directive 2014/35/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to the making available on the market of electrical equipment designed for use within certain voltage limits (OJ L 96, 29.3.2014, p. 357–374)

EN 61010-1:2010

- Directive 2014/30/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to electromagnetic compatibility (OJ L 96, 29.3.2014, p. 79–106)

EN 61326-1:2013

- Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (OJ L 174, 1.7.2011, p. 88–110)

EN 50581:2012

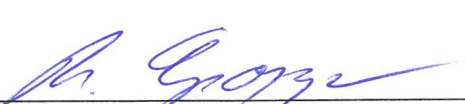
Quality Management System: Certified according to ISO 9001:2015

Manufacturing site: Leica Biosystems Nussloch GmbH, Heidelberger Str. 17-19,  
69226 Nussloch, Germany

Nussloch, 22.06.2020



Andreas Hahn  
Senior Director Core Histology



Robert Gropp  
RA/QA Manager

Rev. B

## EU DECLARATION OF CONFORMITY

We, **Leica Biosystems Nussloch GmbH, Heidelberger Str. 17-19, 69226 Nussloch, Germany**

declare on our own responsibility, that the device

### **Leica IP S**

complies with

- Directive 2014/35/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to the making available on the market of electrical equipment designed for use within certain voltage limits (OJ L 96, 29.3.2014, p. 357–374)

EN 61010-1:2010

- Directive 2014/30/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to electromagnetic compatibility (OJ L 96, 29.3.2014, p. 79–106)

EN 61326-1:2013

- Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (OJ L 174, 1.7.2011, p. 88–110)

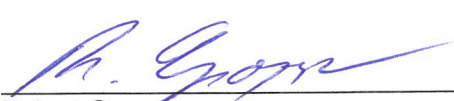
EN 50581:2012

Quality Management System: Certified according to ISO 9001:2015

Manufacturing site: Leica Biosystems Nussloch GmbH, Heidelberger Str. 17-19,  
69226 Nussloch, Germany

Nussloch, 22.06.2020

  
\_\_\_\_\_  
Andreas Hahn  
Senior Director Core Histology

  
\_\_\_\_\_  
Robert Gropp  
RA/QA Manager

Rev. B

certificate  
no. IFA 2205007  
dated 2022-05-20



**IFA**

Institut für Arbeitsschutz der  
Deutschen Gesetzlichen Unfallversicherung  
Prüf- und Zertifizierungsstelle im DGUV Test

Translation In any case, the German original shall prevail.

## DGUV Test Certificate

Name and address of the holder of the certificate (customer): TEKA Absaug- und Entsorgungstechnologie GmbH  
Millenkamp 9  
48653 Coesfeld  
GERMANY

Product designation: **Mobile welding fume separation equipment**

Type: filtoo

Testing based on: DIN EN ISO 21904-1: (06/2022);  
DIN EN ISO 21904-2: (06/2022)

Test report: 202221229/1140 of 2022-05-20, IFA - Sankt Augustin

Further details: The model named above complies with the standard and meets the requirements of welding fume separation class "W3".  
The device is intended to be used for separation of fume, generated by welding, cutting and allied processes of non-ferrous metal, mild and high-alloy steel including chromium nickel steel.  
For recirculation of the exhaust air of the device, national regulations for health and safety at work have to be observed.  
In Germany exhaust air may be recirculated under consideration of the national technical rule TRGS 528.  
Capture of welding fume and separation of gas emissions were not within the scope of this test.

This DGUV Test-Certificate replaces the GS-Certificate with the certificate no: 1102002, dated 2017-01-31.

The type tested complies with the test basis specified above. The holder of the certificate is entitled to affix the DGUV Test mark shown overleaf to the products complying with the type tested, including the specification given under the heading further details.

The present certificate including the right to affix the DGUV Test mark is valid until: **2027-05-19**

Further provisions concerning the validity, the extension of the validity and other conditions are laid down in the Rules of Procedure for Testing and Certification.

  
Dr. rer. nat. Peter Paszkiewicz  
Head of testing and certification body

  
Dipl.-Chem. Thomas von der Heyden  
Certification officer

# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Leica Biosystems Nussloch GmbH  
Heidelberger Strasse 17-19  
69226 Nussloch  
Germany

Holds Certificate Number:

**MD 542493**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

The design & development, manufacturing and service of IVD instruments and accessories including consumables for specimen preparation for histological analysis.



For and on behalf of BSI:

Gary E Slack, Senior Vice President - Medical Devices

Original Registration Date: 2008-12-24

Latest Revision Date: 2020-12-07

Effective Date: 2020-12-24

Expiry Date: 2023-12-23

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# Certificate of Registration

## QUALITY MANAGEMENT SYSTEM - ISO 9001:2015

This is to certify that:

Leica Biosystems Nussloch GmbH  
Heidelberger Strasse 17-19  
Nussloch  
69226  
Germany

Holds Certificate No:

**FM 543970**

and operates a Quality Management System which complies with the requirements of ISO 9001:2015 for the following scope:

The design & development, manufacturing and service for instruments and accessories for research and industrial specimen preparation.

Previous certificate expired on 23/12/2020.  
Recertification audit ended on 06/11/2020.

For and on behalf of BSI:



Andrew Launn, EMEA Systems Certification Director

Original Registration Date: 2008-12-24

Latest Revision Date: 2021-01-08

Effective Date: 2021-01-08

Expiry Date: 2023-12-23



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Cartridge Kit, 280 ml

Cartridge Kit, 280 ml



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Lamp Flash bulb UV-flash dry

Lamp Flash bulb UV-flash dry



Leica Biosystems  
21440 W. Lake Cook Road  
Floor 5  
Deer Park, IL 60010 United States

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SKU: 14049351403

Basket assembly

Basket assembly



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## 1. List of spare parts

| Spare filters                                                      | Art. No. |
|--------------------------------------------------------------------|----------|
| Gross filter, 10 pieces                                            | 978003   |
| Prefilter, 1 piece                                                 | 978004   |
| Main filter, 1 piece                                               | 978005   |
| Activated carbon filter, 1 piece                                   | 978006   |
| Other spare parts                                                  | Art. No. |
| Suction arm type filtoo complete, diam. 150                        | 978009   |
| Spare hose for suction arm type filtoo                             | 978007   |
| Spare hood for suction arm type filtoo                             | 978008   |
| Spare frame for suction arm type filtoo                            | 978010   |
| Set suction hose: 3 m hose diam. 150 with suction nozzles and hood | 978011   |
| PE bags for the disposal of the filters, 3 pieces                  | 10030257 |

Product number: 978003

Gross filter set for filtoo,  
ViroLine and AirToo



| Description  | Properties | Reviews |
|--------------|------------|---------|
| Width [mm]:  | 490        |         |
| Depth [mm]:  | 490        |         |
| Length [mm]: | 490        |         |

[Products](#) > [Spare filters](#) > [Pre-filter](#)

Product number: 978004

## Pre filter for filtoo, ViroLine and AirToo



Description

**Properties**

Reviews

|                                   |     |
|-----------------------------------|-----|
| Filter surface [m <sup>2</sup> ]: | 4   |
| Weight [kg]:                      | 1   |
| Width [mm]:                       | 484 |
| Depth [mm]:                       | 484 |
| Length [mm]:                      | 48  |

[Products](#) > [Spare filters](#) > [Particle filter](#)

Product number: 978005

## Main filter for filtoo and AirToo



Description

**Properties**

Reviews

Filter surface [m<sup>2</sup>]: 11.5

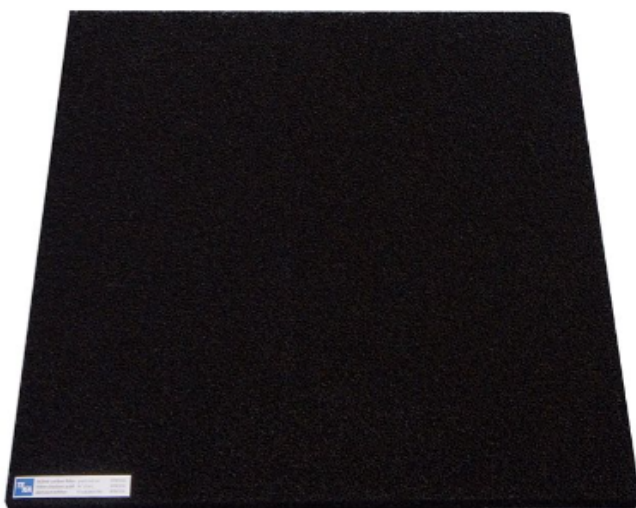
Width [mm]: 520

Depth [mm]: 520

[Products](#) > [Spare filters](#) > [Activated carbon filter](#)

Product number: 978006

## Activated carbon filter



Description

**Properties**

Reviews

Width [mm]: 484

Depth [mm]: 484

Length [mm]: 20