



CERTIFICATE

Certificate No: 1421C01230503
Effective Date: 21/05/2023
Expiry Date: 20/05/2026
Initial Certification Date: 21/05/2020
Last revision date: References: W001142104

This is to certify that the quality management system of Ultra For medical products (Ultramed) Co. (U.M.I.C.) S.A.E.

Part No. (304:310) & part no. (3

nanagement system of Part No. (304:310) & part no. (312) -Industrial Area,

Arab El Awamer - Abnoub - Assiut - Egypt

has been assessed and found compliant to ISO 9001:2015 EN ISO 9001:2015

EA Code(s) 23

Scope of certificate: Manufacturing of General Non-Active, Non-Implantable, Sterile & Non-Sterile

Medical Devices, as detailed on the next page (attachment)

For and on behalf of HTCert

GEORGE PAPPOUS Managing Director FILIPPOS KOTTIS
Certification Director





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ATTACHMENT TO CERTIFICATE

Certificate No: **1421C01230503** Effective Date: **21/05/2023**

Products:

- Sterile Intravenous Infusion Sets with and without needle
- Sterile Blood Transfusion Sets
- Sterile Umbilical Cord clamps
- Sterile Urine Bags
- Sterile I.V. Cannulas
- Sterile Surgical Glove
- Sterile Guedel Airway
- Sterile Mucus Extractors
- Sterile Burette Sets
- Sterile Nelaton Catheters
- Sterile Suction Catheters
- Sterile Suction Units
- Sterile Ryle's Tubes
- Sterile Infant Feeding Tubes
- Sterile 3 ways Stop Cocks
- Sterile Nasal Oxygen Cannulas
- Sterile Silicon Foley Catheters
- Sterile Latex Foley Catheters
- Sterile Fistula NeedlesSterile Oxygen Masks
- Sterile Flow Regulators
- Sterile I.V. Tubing
- Sterile Endotracheal Tubes with and without cuff
- Sterile Extension Tubes with and without stop cock
- Non-sterile Examination Gloves
- Non-sterile 3-ply face mask
- Non-sterile Nasal Oxygen Cannulas (Twin Bore Nasal Oxygen Cannula)
- Non-sterile Oxygen Face Masks
- Non-sterile Urine Collection Bags

For and on behalf of HTCert

GEORGE PAPPOUS Managing Director FILIPPOS KOTTIS
Certification Director





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CERTIFICATE

Certificate No: 1421C02230503

Effective Date: 21/05/2023

Expiry Date: 20/05/2026

Initial Certification Date: 21/05/2020

Last revision date: References: W001 1421 04

This is to certify that the quality management system of Ultra For medical products (Ultramed) Co. (U.M.I.C.) S.A.E.

Part No. (304:310) & part no. (3

nanagement system of Part No. (304:310) & part no. (312) -Industrial Area, Arab El Awamer - Abnoub - Assiut - Egypt

has been assessed and found compliant to ISO 13485:2016 EN ISO 13485:2016

Technical Area(s): - General non-active, non-implantable medical devices

- Ethylene Oxide Gas Sterilization

Scope of certificate: Manufacturing of General Non-Active, Non-Implantable, Sterile & Non-

Sterile Medical Devices, as detailed on the next page

For and on behalf of HTCert

GEORGE PAPPOUS Managing Director FILIPPOS KOTTIS
Certification Director





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ATTACHMENT TO CERTIFICATE

Certificate No: **1421C02230503** Effective Date: **21/05/2023**

Products:

- Sterile Intravenous Infusion Sets with and without needle
- Sterile Blood Transfusion Sets
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- Sterile Urine Bags
- Sterile I.V. Cannulas
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- Sterile Guedel Airway
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- Sterile Burette Sets
- Sterile Nelaton Catheters
- Sterile Suction Catheters
- · Sterile Suction Units
- Sterile Ryle's Tubes
- Sterile Infant Feeding Tubes
- Sterile 3 ways Stop Cocks
- Sterile Nasal Oxygen Cannulas
- Sterile Silicon Foley Catheters
- Sterile Latex Foley Catheters
- Sterile Fistula NeedlesSterile Oxygen Masks
- Sterile Flow Regulators
- Sterile I.V. Tubing
- Sterile Endotracheal Tubes with and without cuff
- Sterile Extension Tubes with and without stop cock
- Non-sterile Examination Gloves
- Non-sterile 3-ply face mask
- Non-sterile Nasal Oxygen Cannulas (Twin Bore Nasal Oxygen Cannula)
- Non-sterile Oxygen Face Masks
- Non-sterile Urine Collection Bags

For and on behalf of HTCert

GEORGE PAPPOUS Managing Director FILIPPOS KOTTIS
Certification Director





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EC-CERTIFICATE

PRODUCTION QUALITY ASSURANCE

This is to certify that the quality management system of

Ultra for medical products (Ultramed) Co (U.M.I.C) S.A.E.

Part No. (304:310) & part no. (312) - Industrial Area, Arab El Awamer - Abnoub - Assiut - Egypt

for manufacturing and final testing of

Sterile Disposable Medical Devices *Further details are given overleaf*

Certificate No: 1421C05201201
Issue Date: 14/12/2020
Original Approval: 14/12/2020
Valid until: 26/05/2024

References: W001 1421 01

HTCert is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification number 2803

fulfills the requirements of Annex V of Council Directive 93/42/EEC.

The use of CE Marking followed by the HTCert Notified Body identification number 2803 for the devices listed on the certificate is hereby authorised. The certificate remains valid subject to satisfactory surveillance audits, periodic or unexpected. Any significant changes in design or manufacture may render this certificate invalid. For class I sterile devices the certificate covers only the aspects of manufacture concerned with securing and maintaining sterile conditions. For class I devices with a measuring function the certificate covers only the aspects of manufacture concerned with the conformity of the products with metrological requirements.

For and on behalf of HTCert

GEORGE PAPPOUS

Managing Director

FILIPPOS KOTTIS

Certification Director

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Attachment to Certificate

No: 1421C05201201

Issued: 14/12/2020

Devices / device categories included in the certificate

Class I sterile devices:

- 3 ways Stop Cocks.
- Extension Tubes with and without stop cock.
- Umbilical Cord clamps.
- Urine Collection Bags.

Class IIa devices:

- Intravenous Infusion Sets with and without needle.
- Blood Transfusion Sets.
- I.V. Cannulas.
- Burette Sets.
- Endotracheal Tubes with and without cuff.
- Flow Regulators.
- Guedel Airway.
- Mucus Extractors.
- Nelaton Catheters.
- Suction Catheters.
- Suction Units.
- Ryle's Tubes.
- Infant Feeding Tubes.
- Surgical Gloves.
- Nasal Oxygen Cannulas.
- Silicon Foley Catheters.
- Latex Foley Catheters.
- Fistula Needles.
- Oxygen Masks.

For and on behalf of HTCert

GEORGE PAPPOUS

Managing Director

FILIPPOS KOTTIS

Certification Director

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ULTRA FOR MEDICAL PRODUCTS CO.

Declaration of Conformity

Manufacturer's Name : Ultra for medical products (Ultramed) Co (U.M.I.C) S.A.E.

Manufacturer's Address: Part No. (304:310) & part no. (312) -

Industrial Area, Arab El Awamer - Abnoub -

Assiut - Egypt

Internet : www.ultramedumic.com

Authorized Representative (AR)

Name : Obelis SA

Address : Bd. Général Wahis 53

1030 Brussels - Belgium

 Phone
 : +32 (0)2 732 59 54

 Fax
 : +32 (0)2 732 60 03

 E-mail
 : mail@obelis.net

 Internet
 : www.obelis.com

We hereby declare that the distributed CE marked products, specified in the attached product list, conform to the type covered by the provisions of the Council Directive 93/42/EEC, June 1993 concerning medical devices amended by Directive 2007/47/EC, September 2007 under the supervision of the Notified Body the Notified Body "HT CERT" (CE 2803).

Based on

Certificate No. : 1421C05201201 Issue Date : 14/12/2020 Expiry Date : 26/05/2024

- G.F.I. Health Technology Certification Ltd

Address : Jakovides Tower 81 - 83 Grivas Digenis

Av., CY 1090, Nicosia Cyprus

Phone : +357 (22) 503000 Fax : +357 (22) 503001 Internet : www.htcert.com

In addition, we ensure and declare that the distributed CE marked products, as mentioned, meet the provisions of the EC-Directive that apply to them.

This declaration is based on MDD 93/42/EEC (Annex-VII) & is subject to the procedure set out in Annex-V of directive 93/42/EEC, June 1993 amended by Directive 2007/47/EC, September 2007.

We also declare that the finished product shall conform to the specifications and functional requirements for 5 years from the date of sterilization.

Declaration of Conformity

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Storage Conditions: Not more than 5 cartoons on each other, Nice Ventilated place, Out of Sun light and a suitable temperature (18 - 32 °C). **Attached List:**

#	Product Name	Classification
1.	Intravenous Infusion Set (I.V. Infusion Set)	IIa
2.	Intravenous Cannula (I.V. Cannula)	IIa
3.	Blood Administration Set (Blood Transfusion Set)	IIa
4.	Extension Tube with/without Three Way Stop Cock	I
5.	Three Way Stop Cock	I
6.	Arterial Venous Fistula Needle (A.V. Fistula Needle)	IIa
7.	Infant Feeding Tube	IIa
8.	Ryle's Tube	IIa
9.	Suction Catheter	IIa
10.	Nelaton Catheter	IIa
11.	Mucus Extractor	IIa
12.	Umbilical Cord Clamp	I
13.	Close Wound Suction Unit (Hemosuc)	IIa
14.	Urine Collection Bag (Adult - Paediatric)	I
15.	Guedel Airway	IIa
16.	Burette Set	IIa
17.	Endotracheal Tube (Cuffed - Uncuffed)	IIa
18.	Latex Foley Catheter	IIa
19.	Silicon Foley Catheter	IIa
20.	Nasal Oxygen Cannula (Adult - Paediatric - Neonatal)	IIa
21.	Surgical Gloves Ultra Easy	IIa
22.	Surgical Gloves Ultra Touch	IIa
23.	Oxygen Face Mask (Adult - Paediatric)	IIa
24.	Flow Regulator	IIa
25.	Suction Tube (Yankaur & Handle)	IIa



Dated on: - 20/12/2020

Assiut Factory: Part No. 304, 305, 306, 307, 308, 309, 310, 312, Arab El Awamer- industrial zone, Abnoub, Assiut, Egypt.. **Tel**: 002-088-4964333 (500), 002-088-4964666 (600), 002-088-4964777 (700), 002-088-4964888 (800), 002-088-4964999 (900)

& Fax:002-088-4964222**& Mob:** 002-01001558853, 002-01068832355

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Assuit Office: 23, July Str. Tel: 088/2364111 – 2364222 **& Fax:** 088/2334964 & Mob. : 01223988202

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Declaration of Conformity

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