ZERTIFIKAT ◆ CERTIFICATE ◆ 证书 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 12 66097 065

Manufacturer: **B. Braun Avitum AG**
Schwarzenberger Weg 73 - 79
34212 Melsungen
GERMANY

Facility(ies):
B. Braun Avitum AG
Schwarzenberger Weg 73 - 79, 34212 Melsungen, GERMANY

B. Braun Avitum Italy S.p.A.
Via XXV Luglio, 11, 41037 Mirandola (MO), ITALY

B. Braun Medical Kft Production Division
Déli-Külhatár út 2-4, 3200 Gyöngyös, HUNGARY

B. BRAUN Vietnam Co., Ltd.
Thanh Oai Industrial Complex, Thanh Oai District, 156800 Hanoi,
VIETNAM

Product Category(ies): **Kit for Dialysis and Haemo(dia)filtration (extracorporeal circuit and dialyser)**
Lines for Dialysis and Hemo(dia)filtration

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.: 713053685

Valid from: 2015-06-23
Valid until: 2020-06-22

Date, 2014-12-18

Hans-Heiner Junker

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

TÜV®

A1 / 04.11

ZERTIFIKAT ◆ CERTIFICATE ◆ 证书 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Traducere din limba engleză

CERTIFICAT CE
SISTEMUL COMPLET DE ASIGURARE A CALITATII

(Directiva 93/42/CEE privind Dispozitivele medicale), Anexa II excluzând (4)
(Dispozitive in clasa IIa, IIb sau III)
Număr G1 14 12 66097 065

Sigla TUV

Producător: **B. Braun Avitum AG**
Schwarzenberger Weg 73 – 79
34212 Melsungen
GERMANIA

Unități:
B. Braun Avitum AG
Schwarzenberger Weg 73 - 79, 34212 Melsungen, GERMANIA

B. Braun Avitum Italy S.p.A.
Via XXV Luglio, 11, 41037 Mirandola (MO), ITALIA

B. Braun Medical Kft Production Division
Deli-Kulhater ut2-4, 3200 Gyongyos, UNGARIA

B. BRAUN Vietnam Co, Ltd.
Complexul industrial Thanh Oai, districtul Thanh Oai, 156800 Hanoi,
VIETNAM

Categorie (ii) Produse: Kit pentru dializă și hemo(dia)filtrare (circuit extracorporal și dializor)
Linii pentru dializă și hemo(dia)filtrare

Organismul de certificare al TUV Product Service GmbH declara ca producătorul mai sus menționat a implementat un sistem de asigurare a calității pentru producție în conformitate cu Anexa II din Directiva privind Dispozitivele medicale. Acest sistem de asigurare a calității este in conformitate cu cerințele prezentei directive si este supus unei supravegheri periodice. Pentru punerea pe piață a dispozitivelor din clasa III, este obligatoriu un certificat suplimentar Anexa II (4). A se vedea, de asemenea, notele de pe verso.

Raport nr. 713053585
Valabil de la: 23.06.2015
Valabil până la: 22.06.2020

Data, 18.12.2014
/semnătură indescifrabilă/
Hans-Heiner Junker

Sigla TUV

TUV Product Service GmbH este un Organism Certificat cu număr de identificare 0123.

Subsemnata, **VALERICA PĂTRU**, interpret si traducător autorizat pentru limbile ENGLEZĂ, FRANCEZĂ şi ITALIANĂ în temeiul autorizaţiei nr. 17602, eliberată de Ministerul Justiţiei din România, certific exactitatea traducerii efectuate din limba *ENGLEZĂ* în limba *ROMÂNĂ*, că textul prezentat spre traducere a fost tradus în întregime, fără omisiuni, şi că, prin traducere, înscrisului nu i-au fost denaturate conţinutul şi sensul.

Traducător şi interpret autorizat,
Valerica Pătru (17602)

ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 17 08 66097 082

Manufacturer: **B. Braun Avitum AG**
Schwarzenberger Weg 73-79
34212 Melsungen
GERMANY

Facility(ies): B. Braun Avitum Italy S.p.A.
Via XXV Luglio, 11, 41037 Mirandola (MO), ITALY

Product Category(ies): **Accessories for dialysis, infusion and apheresis (class I sterile)**

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 V. 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.: 713113517

Valid from: 2018-01-10
Valid until: 2023-01-09

Date, 2017-10-11

Stefan Preiß

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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A1 / 04.11

ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

DAkkS CRT2 / 10.13

Product Service

CERTIFICATE

No. Q1N 16 06 66097 071

Holder of Certificate: **B. Braun Avitum AG**
Schwarzenberger Weg 73-79
34212 Melsungen
GERMANY

Facility(ies):
B. Braun Avitum AG
Schwarzenberger Weg 73-79, 34212 Melsungen, GERMANY

B. Braun Avitum AG
Am Buschberg 1, 34212 Melsungen, GERMANY

B. Braun Avitum AG, Werk Glandorf
Kattenvenner Straße 32, 49219 Glandorf, GERMANY

Certification Mark:

Scope of Certificate: **Design and Development, Production, Distribution and Servicing of Active and Non-Active Medical Devices for Extracorporeal Blood Treatment (Hemodialysis, Acute Dialysis, Apheresis);**
Design and Development, Production and Distribution of Non-Active Medical Devices in various Therapies;
Distribution and Servicing of Active Medical Devices for Reverse Osmosis Systems, Central Concentrate Supply, Ring Piping and Hot Rinse Disinfection Systems

Applied Standard(s): EN ISO 13485:2012 + AC:2012
Medical devices - Quality management systems - Requirements for regulatory purposes
(ISO 13485:2003 + Cor. 1:2009)
DIN EN ISO 13485:2012

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.: 713078602
Valid from: 2016-08-01
Valid until: 2019-07-31

Date, 2016-07-27

Stefan Preiß

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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ZERTIFIKAT ◆ CERTIFICATE ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 17 11 66097 086

Manufacturer: **B. Braun Avitum AG**
Schwarzenberger Weg 73-79
34212 Melsungen
GERMANY

Facility(ies): B. Braun Avitum AG, Werk Glandorf
Kattenvenner Straße 32, 49219 Glandorf, GERMANY

B. Braun Avitum Italy S.p.A.
Via XXV Luglio, 11, 41037 Mirandola (MO), ITALY

Product Category(ies): **Sterile and non-sterile hemodialysis concentrates (class IIb), solid dosage form for hemodialysis (class IIb), and irrigation solutions (class IIa)**

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.: 713120020

Valid from: 2018-02-17
Valid until: 2023-02-16

Date, 2017-11-29

Stefan Preiß

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

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ZERTIFIKAT ◆ CERTIFICATE ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Traducere din limba engleză

CERTIFICAT CE
SISTEMUL COMPLET DE ASIGURARE A CALITATII

(Directiva 93/42/CEE privind Dispozitivele medicale), Anexa II excluzând (4)
(Dispozitive în clasa IIa, IIb sau III)
Număr G1 17 11 66097 086

Sigla TUV

Producător: **B. Braun Avitum AG**
Schwarzenberger Weg 73 – 79
34212 Melsungen
GERMANIA

Unități: **B. Braun Avitum AG, Werk Glandorf Kattenvenner Strasse 32, 49219 Glandorf, GERMANIA**

B. Braun Avitum Italia S.p.A.
Via XXV Luglio, 11, 41037 Mirandola (MO), ITALIA

Categorie (ii) Produse: Concentrate de hemodializă sterile și nesterile (clasa IIb), formă solidă de dozare pentru hemodializă (clasa IIb) și soluții de irigare (clasa IIa)

Organismul de certificare al TUV Product Service GmbH declară ca producătorul mai sus menționat a implementat un sistem de asigurare a calității pentru producție în conformitate cu Anexa II din Directiva privind Dispozitivele medicale. Acest sistem de asigurare a calității este în conformitate cu cerințele prezentei directive si este supus unei supravegheri periodice. Pentru punerea pe piață a dispozitivelor din clasa III, este obligatoriu un certificat suplimentar Anexa II (4). A se vedea, de asemenea, notele de pe verso.

Raport nr. 713120020
Valabil de la: 17.02.2018
Valabil până la: 16.02.2023

Data, 29.11.2017
/semnătură indescifrabilă/
Stefan Preiss

Sigla TUV

TUV Product Service GmbH este un Organism Certificat cu număr de identificare 0123.

Subsemnata, **VALERICA PĂTRU**, interpret si traducător autorizat pentru limbile ENGLEZĂ, FRANCEZĂ și ITALIANĂ în temeiul autorizației nr. 17602, eliberată de Ministerul Justiției din România, certific exactitatea traducerii efectuate din limba *ENGLEZĂ* în limba *ROMÂNĂ*, că textul prezentat spre traducere a fost tradus în întregime, fără omisiuni, și că, prin traducere, înscrisului nu i-au fost denaturate conținutul și sensul.

Traducător și interpret autorizat,
Valerica Pătru (17602)

Ministerul Justiției
VALERICA PĂTRU
Traducător autorizat
Engleză, Franceză, Italiană
Autorizația Nr. 17602

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JOLINE GmbH & Co. KG • Neue Rottenburger Str. 50 • D-72379 Hechingen

KONFORMITÄTSERKLÄRUNG / DECLARATION OF CONFORMITY

Name und Adresse der Firma / Name and address of the firm

Joline GmbH & Co. KG
Neue Rottenburger Str. 50
72379 Hechingen
Germany

Wir erklären in alleiniger Verantwortung, dass das Medizinprodukt /
We declare under our sole responsibility that the medical device

Dialyse Katheter ST gemäß Anhang /
Dialysis Catheter ST according to the annex

der Klasse IIa / of class IIa
nach Anhang IX der Richtlinie 93/42/EWG / according to Annex IX of directive 93/42/EEC

allen Anforderungen der Richtlinie 93/42/EWG entspricht, die anwendbar sind /
meets all the provisions of the directive 93/42/EEC which apply to it.

Konformitätsbewertungsverfahren / Conformity assessment procedure

gemäß Richtlinie 93/42/EWG, Anhang II ohne Abschnitt (4) /
according to directive 93/42/EEC Annex II without section (4)

Konformitätsbewertungsstelle / Notified Body

DEKRA Certification GmbH
Handwerkstr. 15
70565 Stuttgart
Germany
ID: 0124

Diese Erklärung ist gültig bis zum 29.11.2023 bzw. bis zur Ausstellung einer revidierten Erklärung. /
This declaration is valid until 2023-11-29 or until a revised declaration comes into effect.

Hechingen, 2018-11-30

Michael Eisenlohr
Site Manager

Dr. Marian Wenzel
Director QA/RA

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ANHANG – PRODUKTLISTE / ANNEX – PRODUCT LIST

Single Lumen Short Term

DCPT 8/10	DCPT 8/17,5 PH	PKSL08P150C	PKSL08P250
DCPT 8/15	DCPT 8/20 H-PH	PKSL08P175	PKSL08P250C
DCPT 8/15 H-PH	DCPT 8/20 PH	PKSL08P175C	
DCPT 8/15 PH	DCPT 8/25 PH	PKSL08P200	
DCPT 8/17,5 H-PH	PKSL08P150	PKSL08P200C	

Single Lumen ST (Händler / Distributor)

-PKSL08P150	-PKSL08P175	-PKSL08P200	-PKSL08P250

Double Lumen Short Term Extra Flow Kits

HF-DLS 11/12,5	KEF11P100	KEF11P250R	PKHF11P175
HF-DLS 11/12,5 C	KEF11P125	KEF11P300	PKHF11P200
HF-DLS 11/15	KEF11P125C	KEF11P300C	PKHF11P250
HF-DLS 11/15 C	KEF11P125R	KEF11P300R	PKHF11P150R
HF-DLS 11/15 C 14	KEF11P150	KEF11PH125R	PKHF11P175R
HF-DLS 11/17,5	KEF11P150C	KEF11PH150C	PKHF11P125
HF-DLS 11/17,5 C	KEF11P150R	KEF11PH150R	PKHF11P225
HF-DLS 11/20	KEF11P175	KEF11PH175	PKHF11P125R
HF-DLS 11/20 C	KEF11P175C	KEF11PH175R	PKHF11P200R
HF-DLS 11/20 C 14	KEF11P175R	KEF11PH200	PKHF11P225R
HF-DLS 11/22,5	KEF11P200	KEF11PH200C	PKHF11P250R
HF-DLS 11/25	KEF11P200C	KEF11PH200R	PKHF11P150R 21
HF-DLS 11/25 C	KEF11P200R	KEF11PH250	PKHF11P150C 21
HF-DLS 11/25 C 14	KEF11P225R	KEF11PH300	PKHF11P175C 21
HF-DLS 11/30	KEF11P250	PKHF11P150	

Double Lumen ST Extra Flow (Händler / Distributor)

-PKHF11P150	-PKHF11P200	-PKHF11P150R	-PKHF11P200R
-PKHF11P175	-PKHF11P250	-PKHF11P175R	-PKHF11P250R

Double Lumen Short Term Standard - Einzelkatheter und Kits

DLS 11/20 C	KDL11P150R	KDL11PH125C	PKDL11P150 21
DLS 11/12,5	KDL11P175	KDL11PH125R	PKDL11P150C 21
DLS 11/15	KDL11P175C	KDL11PH150	PKDL11P150R
DLS 11/15 C	KDL11P175R	KDL11PH150C	PKDL11P175
DLS 11/17,5	KDL11P200	KDL11PH150R	PKDL11P175 21
DLS 11/17,5 C	KDL11P200C	KDL11PH175	PKDL11P175C 21
DLS 11/20	KDL11P200R	KDL11PH175C	PKDL11P175R
DLS 11/25	KDL11P225	KDL11PH175R	PKDL11P200
KDL11P100	KDL11P225C	KDL11PH200	PKDL11P200R
KDL11P100C	KDL11P225R	KDL11PH200C	PKDL11P250
KDL11P125	KDL11P250	KDL11PH200R	PKDL11P250R
KDL11P125C	KDL11P250C	KDL11PH250	PKDL11PH200
KDL11P125R	KDL11P250R	PKDL11P125	
KDL11P150	KDL11PH100R	PKDL11P125R	
KDL11P150C	KDL11PH125	PKDL11P150	

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Double Lumen ST (Händler / Distributor)

-PKDL11P125	-PKDL11P200	-PKDL11P150R	-PKDL11P250R
-PKDL11P150	-PKDL11P250	-PKDL11P175R	
-PKDL11P175	-PKDL11P125R	-PKDL11P200R	

Double Lumen Short Term High Flow Kits

HF-DLS 13/15	HF-DLS 13/20 C	PKHF13PH150C 21	PKHF13PH200R
HF-DLS 13/15 C	HF-DLS 13/25	PKHF13PH150R	PKHF13PH250
HF-DLS 13/17,5	HF-DLS 13/22,5	PKHF13PH175	PKHF13PH250R
HF-DLS 13/17,5 C	PKHF13PH150	PKHF13PH175R	PKHF13PH300
HF-DLS 13/20	PKHF13PH150 21	PKHF13PH200	

High Flow Double Lumen ST (Händler / Distributor)

-PKHF13PH150	-PKHF13PH200	-PKHF13PH150R	-PKHF13PH200R
-PKHF13PH175	-PKHF13PH250	-PKHF13PH175R	-PKHF13PH250R

Double Lumen Extra Flow - Pädiatrisch

KDL06P075	KDL08P075C	P-DLS 6,5/7,5	PKDL06P125R
KDL06P075C	KDL08P075R	P-DLS 6,5/7,5 C	PKDL08P075 21
KDL06P075R	KDL08P100	P-DLS 8/15	PKDL08P100
KDL06P100	KDL08P100C	P-DLS 8/10 C	PKDL08P100R
KDL06P100C	KDL08P100R	P-DLS 8/12,5	PKDL08P125
KDL06P100R	KDL08P125	P-DLS 8/12,5 C	PKDL08P125R
KDL06P125	KDL08P125C	P-DLS 8/7,5	PKDL08P150
KDL06P125C	KDL08P125R	P-DLS 8/10	PKDL08P150 21
KDL06P125R	KDL08P150	PKDL06P075	PKDL08P150R
KDL06P150	KDL08P150C	PKDL06P075R	
KDL06P150C	KDL08P150R	PKDL06P100	
KDL06P150R	P-DLS 6,5/10	PKDL06P100R	
KDL08P075	P-DLS 6,5/10 C	PKDL06P125	

Extra Flow Double Lumen (Händler / Distributor)

-PKDL08P100	-PKDL08P100R	-PKDL06P075	-PKDL06P075R
-PKDL08P125	-PKDL08P125R	-PKDL06P100	-PKDL06P100R
-PKDL08P150	-PKDL08P150R	-PKDL06P125	-PKDL06P125R

Triple Lumen Short Term

KTL12P125	KTL12P175R	PKTL12P150 C 21	PKTL12P200C 21
KTL12P125R	KTL12P200	PKTL12P150R	PKTL12P200 21
KTL12P150	KTL12P200C	PKTL12P175	PKTL12P200R
KTL12P150C	KTL12P200R	PKTL12P175 21	PKTL12P250R
KTL12P150R	KTL12P225	PKTL12P175C 21	PKTL12P250
KTL12P175	KTL12P250	PKTL12P175R	
KTL12P175C	PKTL12P150	PKTL12P200	

Triple Lumen ST Händler

-PKTL12P150	-PKTL12P200	-PKTL12P150R	-PKTL12P200R
-PKTL12P175	-PKTL12P250	-PKTL12P175R	-PKTL12P250R

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Triple Lumen Short Term High Flow

HF-TLK 13/15	HF-TLK 13/25	PKTHF13P150R	PKTHF13P200R
HF-TLK 13/15 C	HF-TLK 13/30	PKTHF13P175	PKTHF13P250
HF-TLK 13/17,5	HF-TLK 13/30 C	PKTHF13P175R	PKTHF13P250R
HF-TLK 13/17,5 C	PKTHF13P150	PKTHF13P175C 21	
HF-TLK 13/20	PKTHF13P150 21	PKTHF13P200	
HF-TLK 13/20 C	PKTHF13P150 C 21	PKTHF13P200 21	

Triple Lumen ST Händler

-PKTHF13P150	-PKTHF13P200	-PKTHF13P150R	-PKTHF13P200R
-PKTHF13P175	-PKTHF13P250	-PKTHF13P175R	-PKTHF13P250R

R14

EC CERTIFICATE

Number: 3804606CE01

Full Quality Assurance System

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

Manufacturer:
Cytosorbents, Inc.
7 Deer Park Dr., Suite K
Monmouth Jct. NJ 08852
United States Of America

For the product category(ies)

Polymer Based Adsorption Systems

DEKRA grants the right to use the EC Notified Body identification Number mentioned 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 3804606CN, initially dated 20 September 2010
Addendum, initially dated 25 March 2011

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 September 2019
Issued for the first time: 25 March 2011
Reissued: 1 September 2016

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

ing. A.A.M. Laan
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: 3804606CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Polymer Based Adsorption Systems

Issued to:

Cytosorbents, Inc.
7 Deer Park Dr., Suite K
Monmouth Jct. NJ 08852
United States Of America

This certificate covers the following product(s):

Cytokine, Bilirubin, and Myoglobin, Adsorption (Class IIb)

Initial date: 25 March 2011
Revision date: 9 June 2016

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

ing. A.A.M. Laan
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

CERTIFICAT CE

Numărul: 3804606CE01
Sistemul Complet pentru Asigurarea Calității
Directiva 93/42/CEE privind Dispozitivele Medicale, Anexa II excluzând (4)
(Dispozitivele din Clasele IIa, IIb sau III)

Producător:
Cytosorbents, Inc.
7 Deer Park Dr., Suite K
Monmouth Jct. NJ 08852
Statele Unite ale Americii

Pentru categoria(categoriile) de produse

Sisteme de Absorbție pe Bază de Polimeri

DEKRA oferă dreptul de a utiliza Numărul de Identificare al Organismului Notificat CE de mai jos pentru a însoți Marca de Conformitate CE a produselor implicate care sunt conforme cu Documentația Tehnică necesară și îndeplinesc prevederile Directivei CE care se aplică acestora:

0344

Documentele care stau la baza acestui certificat:

Notificare de Certificare 3804606CN, inițială din data de 20 septembrie 2010
Act Adițional, inițial din data de 25 martie 2011

DEKRA declară că producătorul menționat mai sus îndeplinește prevederile aplicabile ale „Besluit Medische Hulpmiddelen", transpunerea olandeză a Directivei 93/42/CEE din 14 iunie 1993 privind dispozitivele medicale, inclusiv toate modificările ulterioare, și care pentru categoria de produse menționată mai sus, Procedura de Evaluare a Conformității Anexa II pentru produsele din clasa IIa, este realizată de Producător în conformitate cu prevederile Directivei Consiliului 93/42/CEE din 14 iunie 1993. Pentru plasarea pe piață a dispozitivelor din Clasa III, este obligatoriu un certificat suplimentar de examinare a design-ului CE conform Anexei II, secțiunea 4. Informațiile necesare și referirea la documentația aplicabilă privind produsele implicate și evaluările realizate, sunt menționate în Notificarea de Certificare care constituie o parte integrantă din acest certificat.

Acest certificat este valabil până în data de: 1 septembrie 2019
Data eliberării inițiale: 25 martie 2011
Reeliberat: 1 septembrie 2016

DEKRA Certification B.V.

drs. G.J. Zoetbrood	ing. A.A.M. Laan
Director Executiv	Director de Certificare
Semnătură indescifrabilă	*Semnătură indescifrabilă*

ANEXĂ

Atașată la certificatul cu nr. 3804606CE01

DECLARAȚIE DE CONFORMITATE CE

DISPOZITIVE MEDICALE

Sisteme de absorbție pe bază de polimer

Emis către:

Cytosorbents Inc.
7 Deer Park Dr., Suite K
Monmouth Jct. NJ 08852
Statele Unite ale Americii

Acest certificat acoperă următoarele produse:

Citokină, bilirubină, mioglobină, absorbție (clasa IIb).

Data inițială de emitere: 25 martie 2011

Data revizuirii: 9 iunie 2018

DEKRA Certification B.V.

semnătură indescifrabilă	*semnătură indescifrabilă*
Dr. G. J. Zoetbrood	Ing. A. A. M. Laan
Director general	Manager de certificare

DEKRA Certification B.V. este un organism notificat cu nr. 0344

DEKRA Certification B.V. Meander 1051, 8825 MJ Arnhem Cod poștal 5185, 6802 ED Arnhem, Olanda

T +31 88 96 83000 F +88 96 83100 www.dekra-certification.com Nr. de înregistrare 09085396

EC CERTIFICATE

for the Quality Assurance System

according the Directive 93/42/EEC, Annex II excluding section (4)

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company

Joline GmbH & Co. KG

Neue Rottenburger Straße 50, 72379 Hechingen, Germany

applies a quality assurance system according to the Directive 93/42/EEC Annex II for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 50565-Z5-00, the decision dated 2018-10-04 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2018-11-30 to 2023-11-29

Registration No.: 50565-16-06

Benannt durch/Designated by Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten www.zlg.de
ZLG-BS-295,10.02

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart; 2018-10-04
Notified Body ID-number: 0124

Annex to the EC Certificate No. 50565-16-06

Valid from 2018-11-30 to 2023-11-29

Revision status of the annex: 0 dated 2018-11-30

Devices/device categories included in the certificate:

Class II a:

MD 0102

- Dialysis Catheter ST
  - Kits
  - Catheter

MD 0106

- Kyphoplasty Systems ALLEVO
  - Kits
  - Individual Instruments
- Dialysis Accessories
  - Introducer Needle
  - Guide Wire
  - Dilator
  - Trocar
  - Connector LT
- Stone Extraction Catheter

Class III:

MD 0203

- Dialysis Catheter PU-LT
  - Kits
  - Catheter
- Dialysis Catheter Silicone LT
  - Kits
  - Catheter

MD 0106

- Biopsy Forceps KNIPSA

For the placing on the market of class III devices covered by this certificate an EC design-examination certificate according to directive 93/42/EEC annex II (4) is required.

Ruth Delbeck-Bayer
DEKRA Certification GmbH, Stuttgart, 2018-10-04
Notified Body ID-number: 0124

EC CERTIFICATE

for the Quality Assurance System

according the Directive 93/42/EEC, Annex V

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company

Joline GmbH & Co. KG

Neue Rottenburger Straße 50, 72379 Hechingen, Germany

applies a quality assurance system according to the Directive 93/42/EEC Annex V for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 50565-Z5-00, the decision dated 2018-10-04 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2018-11-30 to 2023-11-29

Registration No.: 50565-17-05

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-295.10.02

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart: 2018-10-04
Notified Body ID-number: 0124

Annex to the EC Certificate No. 50565-17-05

Valid from 2018-11-30 to 2023-11-29

Revision status of the annex: 0 dated 2018-11-30

Devices/device categories included in the certificate:

Class I s:

For the products listed below, review of the Quality Assurance System refers exclusively to aspects of manufacture concerned with securing and maintaining sterile conditions.

MD 0101

- Miniclamp

MD 0106

- Mixer

Ruth Delbeck-Bayer
DEKRA Certification GmbH, Stuttgart, 2018-10-04
Notified Body ID-number: 0124