

Date: 21.01.2020

AUTHORIZATION LETTER

To whom it may concern

We, Shenzhen Mindray Bio-Medical Electronics Co., Ltd. ("Mindray"), who is official manufacturer of in-vitro diagnostic products ("Products"), located in Mindray Building, Keji 12th Road South, High-Tech Industrial Park, Nanshan, 518057 Shenzhen, PEOPLE'S REPUBLIC OF CHINA, do hereby authorize:

"Echipamed-Plus" SRL str. Valea Trandafirilor, 24B, of. 80 MD-2001, Chisinau Republic of Moldova

as the exclusive one company to submit a bid, subsequently negotiate and sign the Contract against Tender nr. ocds-b3wdp1-MD-1579165718042 (21018001) dated 20.02.2020 organized by IMSP Institutul de Medicina Urgenta for the Mindray Products in Moldova.

As the manufacturer, Mindray guarantees the Products against defects in materials and workmanship, and provide services based on the standard terms and conditions of Mindray's warranty policy.

The present authorization is valid from the date of issuance up to December 31st, 2020.

Yours faithfully,

Peno

Sales Manager, of International Sales and Marketing Division, CIS

Shenzhen Mindray Bio-Medical Electronics Co., Ltd.

SHENZHEN MINDRAY BIO-MEDICAL ELECTRONICS CO., LTD.

Mindray Building, Keji 12th Road South, High-tech Industrial Park, Nanshan, Shenzhen 518057, P.R. China Tel: +86 755 81888998

Fax: +86 755 26582680 Website: www.mindray.com





CERTIFICATE

No. QS5 17 07 44751 097

Certificate Holder:

Shenzhen Mindray Bio-Medical

Electronics Co., Ltd. Mindray Building Keji 12th Road South High-Tech Industrial Park

Nanshan

518057 Shenzhen

PEOPLE'S REPUBLIC OF CHINA

Certification Mark:





Scope of Certificate:

Design and Development, Production and Distribution of Medical Electronic Equipment (Including Patient Monitor and Accessories, Vital Signs Monitor, Center Monitoring System, Telemetry Monitoring System, Pulse Oximeter, Temperature Probe, Flow Sensor, Ambulatory Blood Pressure Monitor, Defibrillator/Monitor and Accessories. Electrocardiograph, Wearable ECG Recorder, Anesthesia Machine and Accessories, Ventilator, Infusion Pump, Syringe Pump, Enteral Feeding Pump, Infusion Supervision System Ultrasonic Diagnostic Equipment and Accessories, Digital Radiography System, Radiography System, Magnetic Resonance Imaging System, Hematology Analyzer, Clinical Chemistry Analyzer, Urine Analyzer, Microplate Reader, Microplate Washer for Invitro Diagnostic Use, Chemiluminescence Immunossay Analyzer, Flow Cytometer, (Auto) Sample Processing System, Auto Slide Maker & Stainer, Glycohemoglobin Analyzer, Specific Protein Analyzer), Reagents for Hematology Analyzer, Reagents for Clinical Chemistry Analyzer, Chemiluminescence Immunoassay Reagents, Chemiluminescence Immunoassav Calibrators and Controls, Reagents for Flow Cytometer, Reagents for Glycohemoglobin Analyzer, Calibrators and Controls for Glycohemoglobin Analyzer, Disposable Anesthesia Mask, Reusable Anesthesia Mask, Respiratory Mask, Disposable Breathing Circuit, Reusable Breathing Circuit, Heat and Moisture Exchanger,

Standard(s):

ISO 9001:2015

The Certification Body of TÜV SÜD America Inc. certifies that the company mentioned above has established and is maintaining a quality management system that meets the requirements of the listed standards.

Report No.:

M2606

Effective Date:

2017-07-01

Expiry Date:

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USA

2020-06-30

Earl Buckmiller

Director, Quality Systems & MS Cert. Body

fail Buermiller

TÜV SÜD America Inc. 10 Centennial Drive Peabody, MA 01960





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DAKKS CRT2 / A4 07.17



CERTIFICATE

No. Q5 18 03 44751 111

Holder of Certificate: Shenzhen Mindray Bio-Medical

Electronics Co., Ltd.

Mindray Building Keji 12th Road South High-Tech Industrial Park

Nanshan

518057 Shenzhen

PEOPLE'S REPUBLIC OF CHINA Certification Mark:



Scope of Certificate: Design and development,

production and distribution of

Active medical devices (intended) for monitoring, diagnosis, anesthesia, breathing and intensive care;

In-vitro diagnostic instruments;

Non-active accessories

for breathing therapy and anesthesia;

In-vitro diagnostic reagents and kits (intended)

for hematology, clinical chemistry, immunology and cell analysis

(For detail information see following pages)

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.:

SH1805531

Valid from:

2018-07-09

Valid until:

2020-08-31

Date, 2018-07-09

Stefan Preiß

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CE₀₁₂₃ DECLARATION OF CONFORMITY

To whom it may concern,

We, Shenzhen Mindray Bio-Medical Electronics Co., Ltd., located at Mindray Building, Keji 12th Road South, Hi-tech Industrial Park, Nan-shan, 518057, Shenzhen, P. R. China, hereby confirm that:

We herewith declare that the products listed in Attachment I and Attachment II meet the provisions of the Directive 98/79/EC on In Vitro Diagnostic Medical Devices.

Product Category I: Reagents for Chemiluminescence Immunoassay Analyzer

Products:

Attachment I

Classification:

List B in IVDD annex II

Conformity Assessment Route: Annex IV.3

Notified Body:

TÜV SÜD Product Service GmbH,

Ridlerstraße 65, 80339 München, Germany.

Notified Body NO: 0123

Product Category II: Reagents for Chemiluminescence Immunoassay Analyzer

Products:

Attachment II

Classification: List A in IVDD annex II

Conformity Assessment Route: Annex IV.3 and IV.4

Notified Body:

TÜV SÜD Product Service GmbH,

Ridlerstraße 65, 80339 München, Germany.

Notified Body NO: 0123

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80, 20537 Hamburg, Germany

We hereby certify that the forgoing is a true and accurate statement.

Place, Date Of Issue: Shenzhen, 2017-11-1

Signatory name: Xinbing Wang

Signatory title: Technical Regulation Manager

Shenzhen Mindray Bio-Medical Electronics Co., Ltd.

ATTACHMENT I

Mindray Product list

Total Prostate Specific Antigen (CLIA)

Total PSA Calibrators

Free Prostate Specific Antigen (CLIA)

Free PSA Calibrators

Tumor Marker Multi Control

ATTACHMENT II

Mindray Product list

Hepatitis B Surface Antigen (CLIA)

Antibody to Hepatitis B Surface Antigen (CLIA)

Hepatitis B e Antigen (CLIA)

Antibody to Hepatitis B e Antigen (CLIA)

Antibody to Hepatitis B Core Antigen (CLIA)

Antigen and Antibodies to Human Immunodeficiency Virus (CLIA)

HBsAg Calibrators

Anti-HBs Calibrators

HBeAg Calibrators

Anti-HBe Calibrators

Anti-HBc Calibrators

HIV Calibrators

HBsAg Positive Control

HBsAg Negative Control	
Anti-HBs Positive Control	
Anti-HBs Negative Control	
HBeAg Positive Control	
HBeAg Negative Control	
Anti-HBe Positive Control	
Anti-HBe Negative Control	
Anti-HBc Positive Control	
Anti-HBc Negative Control	
HIV Ag/Ab Positive Control	
HV Ag/Ab Negative Control	



Keji 12th Road South, Hi-tech Industrial Park, Shenzhen

518057, P. R. China Tel: +86 755 26582888

Fax: +86 755 26582500



DECLARATION OF CONFORMITY

To whom it may concern,

We, Shenzhen Mindray Bio-Medical Electronics Co., Ltd., located at Mindray Building, Keji 12th Road South, Hi-tech Industrial Park, Nan-shan, 518057, Shenzhen, P. R. China, hereby confirm that:

We herewith declare that the products listed in Attachment I meet the provisions of the Directive 98/79/EC on In Vitro Diagnostic Medical Devices.

Product Category: Auto Hematology Analyzer and the reagents for Auto Hematology Analyzer, Semi-auto Chemistry Analyzer, Microplate reader, Microplate washer, Chemiluminescence Immunoassay Analyzer and the reagents for Chemiluminescence Immunoassay Analyzer

Products:

Attachment I

Classification: The device not in IVDD annex II and not for self testing/performance evaluation Conformity Assessment Route: Annex III (not includes Section 6)

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80, 20537 Hamburg, Germany

We hereby certify that the forgoing is a true and accurate statement.

Place, Date Of Issue: Shenzhen, 2017-11-1

Signatory name: Xinbing Wang

Signatory title: Technical Regulation Manager

Shenzhen Mindray Bio-Medical Electronics Co., Ltd.



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Fax: +86 755 26582500

ATTACHMENT I

Equipments

Product Name	Model
Auto Hematology Analyzer	BC-2800
Auto Hematology Analyzer	BC-2800Vet
Auto Hematology Analyzer	BC-3000Plus
Auto Hematology Analyzer	BC-3200
Auto Hematology Analyzer	BC-3600
Auto Hematology Analyzer	BC-20s
Auto Hematology Analyzer	BC-30s
Semi-auto Chemistry Analyzer	BA-88A
Microplate reader	MR-96A
Microplate washer	MW-12A
Chemiluminescence Immunoassay Analyzer	CL-1000i

Reagents, Calibrators, Controls and Consumables for Auto Hematology Analyzer

Product Name	Model	
M-30D DILUENT	M-30D	
M-30CFL LYSE	M-30CFL	
M-30R RINSE	M-30R	
M-30E E-Z CLEANSER	M-30E	
M-30P PROBE CLEANSER	M-30P	
PROBE CLEANSER	\	
Hematology Control	B30	
Calibrator	S30	
Hematology Control	BC-3D	
Hematology Calibrator	SC-CAL Plus	

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Reagents for Chemiluminescence Immunoassay Analyzer

Free Thyroxine (CLIA)

Free Triiodothyronine (CLIA)

Total Triiodothyronine (CLIA)



Shenzhen Mindray Bio-Medical Electronics Co., Ltd. Keji 12th Road South, Hi-tech Industrial Park, Shenzhen 518057, P. R. China

Tel: +86 755 26582888 Fax: +86 755 26582500

Fax: +86 755 2658250	00
Total Thyroxine (CLIA)	
Thyroid-Stimulating Hormone (CLIA)	
Thyroglobulin (CLIA)	
Antibody to thyroglobulin (CLIA)	
Antibody to thyroid peroxidase (CLIA)	
Cancer Antigen 125 (CLIA)	
Carbohydrate Antigen 19-9 (CLIA)	
Carcinoembryonic Antigen (CLIA)	
Alpha-fetoprotein (CLIA)	
Cancer Antigen 15-3 (CLIA)	
Cancer Antigen 72-4 (CLIA)	
CYFRA 21-1 (CLIA)	
Neuron-specific enolase (CLIA)	
Total β Human Chorionic Gonadotropin (CLIA)	
Luteinizing Hormone (CLIA)	
Follicle Stimulating Hormone (CLIA)	
Prolactin (CLIA)	
Estriol (CLIA)	
Progesterone (CLIA)	
Testosterone (CLIA)	SINAU. SOCIE
Estradiol (CLIA)	July Settle amed The
Insulin (CLIA)	

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Tel: +86 755 26582888

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C-peptide (CLIA)
Dehyroepiandrosterone sulfate (CLIA)
Cortisol (CLIA)
Adrenocorticotropic hormone (CLIA)
Troponin I (CLIA)
Myoglobin (CLIA)
Creatine kinase MB (CLIA)
B-type natriuretic peptide (CLIA)
Parathyroid hormone (CLIA)
Calcitonin (CLIA)
25-OH-Vitamin D Total (CLIA)
Ferritin (CLIA)
Vitamin B12 (CLIA)
Folate (CLIA)
Red Blood Cell Folate Releasing Reagent
Antibody to Treponema pallidum (CLIA)

Calibrators for Chemiluminescence Immunoassay Analyzer

Free T3 Calibrators Free T4 Calibrators Total T3 Calibrators Total T4 Calibrators



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TSH Calibrators	
Thyroglobulin Calibrators	
Anti-Tg Calibrators	
Anti-TPO Calibrators	
CA125 Calibrators	
CA19-9 Calibrators	
CEA Calibrators	
AFP Calibrators	
CA15-3 Calibrators	
CA72-4 Calibrators	
CYFRA21-1 Calibrators	
NSE Calibrators	
Total β HCG Calibrators	
LH Calibrators	
FSH Calibrators	
Prolactin Calibrators	
Estriol Calibrators	
Progesterone Calibrators	
Testosterone Calibrators	
Estradiol Calibrators	
Insulin Calibrators	
C-peptide Calibrators	
DHEA-S Calibrators	
Cortisol Calibrators	
ACTH Calibrators	
Troponin I Calibrators	
MYO Calibrators	
CK-MB Calibrators	
BNP Calibrators	
PTH Calibrators	
Calcitonin Calibrators	
25-OH-Vitamin D Total Calibrators	
Ferritin Calibrators	MAU. SO
Vitamin B12 Calibrators	S S S A P A M C

Folate Calibrators Anti-TP Calibrators

Keji 12th Road South, Hi-tech Industrial Park, Shenzhen

518057, P. R. China

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Controls for Chemiluminescence Immunoassay Analyzer

Thyroid Function Multi Control
Reproductive Multi Control
Anti-thyroid Antibodies Control
Cardiac Marker Multi Control
Immunoassay Multi Control
ACTH Control
Metabolic Multi Control
NSE control
Anti-TP Control

Consumables for Chemiluminescence Immunoassay Analyzer

Shenzhen Mindray Bio-Medical Electronics Co., Ltd. Keji 12th Road South, Hi-tech Industrial Park, Shenzhen 518057, P. R. China

Tel: +86 755 26582888

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Substrate Solution
Wash Buffer
Detergent CD80
Sample Diluent
Reaction Cuvette



Declaration of Conformity



Manufacturer:

Shenzhen Mindray Bio-Medical Electronics Co., Ltd.

Mindray Building, Keji 12th Road South, Hi-tech Industrial

Park, Nanshan, Shenzhen, 518057, P. R. China

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80

20537 Hamburg, Germany

Product:

DETERGENT C

Classification:

The device not in IVDD annex II and not for self

testing/performance evaluation

Conformity Assessment Route: IVDD Annex III(not includes Section 6)

We herewith declare that the above mentioned products meet the provisions of the Directive 98/79/EC on In Vitro Diagnostic Medical Devices. All supporting documentations are retained under the premises of the manufacturer.

Standards Applied:

List of (harmonized) standards for which documented evidence for compliance can be provided as attachment.

Start of CE-Marking:

2018-4-17

Place, Date of Issue:

Shenzhen, 2018-4-17

Signature:

Mr.WangXinBing

Name of Authorized Signatory:

Position Held in Company:

Manager of Technical Regulation

