Specification

Model	VS-10H/VS-10M/VS-10S
Dimensions	191(H) x 92 (W) x 112 (D) mm
Weight	About 0.25kg (including the battery)
Power Adapter	Input voltage: AC 100-240V 50/60Hz Input power: 25VA. DC output voltage: 5V 2A
Battery	Built-in battery: 3.6V 3400mAh Continuous operation duration of the lithium battery: 4 hours Lithium battery charge time: not longer than 4 hours
Rotation Angle of Display Screen	Maximum vertical rotation angle: 140°±10° Maximum horizontal rotation angle: 270°±10°
Screen	3.5 inches touchscreen with resolution of 640x960 pixels
Water Proof Level	IP66
Photo Record	Photo can be recorded in JPEG format, up to 10,000 photos can be recorded
Video Record	Video can be recorded in MPEG-4 format, up to 2 hours video can be recorded
Media Storage	Not less than 5GB
Data Port	USB Type C, HDMI*, and Wi-Fi** are available
Anti-Fog	Immediate anti-fog technology is supported with no need for heating of the anti-fog coating on the laryngoscope blade
Blades	Single use blade
Frequency for WiFi**	2.4GHz

Note: *For VS-10H & VS-10M, **For VS-10H only

Smaller Size, Bigger Vision

VS-10 Series Video Laryngoscope

MEDCAPTAIN

MEDCAPTAIN LIFE SCIENCE CO.,LTD.

Address: Room 103, 1st Floor, Room 601, 6th Floor, and Room 701, 7th Floor, Block C, Jinweiyuan Industrial Park, Julongshan Subdistrict, Pingshan District, 518118, Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

Tel: +86 0755-28380626
Fax: +86 0755-84517910
E-mail: disposable@medcaptain.com
Website: www.medcaptain.com

EN-Airway Management-10P-Version 1.0

Distributor:

US: Rx only

VS-10 Series Video Laryngoscope

Vision with Safety

- Smart Power Management
  Remaining Time Display
- IP66, Waterproof Design
- Instant anti-fog technology
  No heating time required

Vision with Ease

- 3.5"HD touchscreen: a step closer to reality
- Horizontal rotation angle: 0°~270°
- Vertical rotation angle: 0°~140°

HDMI：Bigger screen display

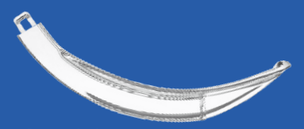
Wifi: Wireless data transportation

Vision For Everyone

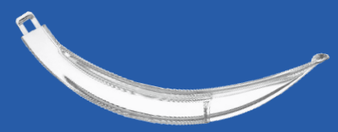
All 5 blades share a general handle
Disposable blades offer a better infection control

Size **M1** for infant

Size **M2** for child

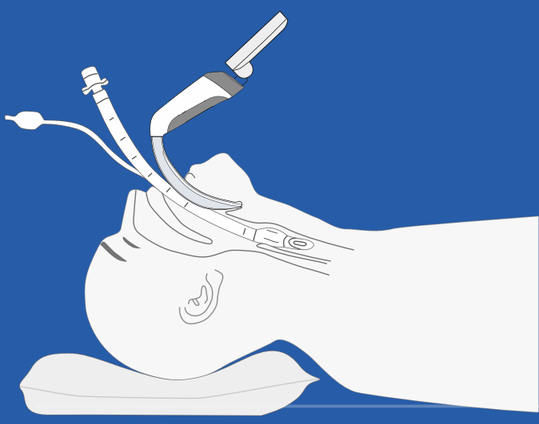


Size **M3** for adult



Size **M4** for obesity/big adult

Size **M3D** for difficult intubation



ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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

TÜV SÜD Product Service

Certificate

No. Q5 086916 0024 Rev. 05

Holder of Certificate:

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.
12th Floor, Baiwang Research Building
No.5158 Shahe West Road
Xili, Nanshan
518055 Shenzhen, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Certification Mark:

TÜV SÜD EN ISO 13485 tuvsud.com/ps-cert

Scope of Certificate:

Design and Development, Production and Distribution of Syringe Pumps, Infusion Pumps, Infusion Workstations, Infusion Central Monitoring Systems, DVT Preventive Pumps, Infusion Docking Stations, Thromboelastograph Instruments and Reagents, Vein Illuminators, Enteral Feeding Pumps, Video Laryngoscopes, Chemiluminescence Immunoassay Analyzers and Reagents, Hemostasis Reagents, Colloidal Gold Reagents, Blood Grouping Systems, Spirometers, Nucleic Acid Extraction Systems, and Oxygen Concentrators.

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). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 086916 0024 Rev. 05

Report No.: GZ2311701

Valid from: 2023-02-17
Valid until: 2026-02-16

Date, 2022-11-22

Christoph Dicks
Head of Certification/Notified Body

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

TÜV®

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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

TÜV SÜD Product Service

Certificate

No. Q5 086916 0024 Rev. 05

Applied Standard(s):

EN ISO 13485:2016
Medical devices - Quality management systems - Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies):

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.
12th Floor, Baiwang Research Building, No.5158 Shahe West Road, Xili, Nanshan, 518055 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

Distribution of Syringe Pumps, Infusion Pumps, Infusion Workstations, Infusion Central Monitoring Systems, DVT Preventive Pumps, Infusion Docking Stations, Thromboelastograph Instruments and Reagents, Vein Illuminators, Enteral Feeding Pumps, Video Laryngoscopes, Chemiluminescence Immunoassay Analyzers and Reagents, Hemostasis Reagents, Colloidal Gold Reagents, Blood Grouping Systems, Spirometers, Nucleic Acid Extraction Systems, and Oxygen Concentrators.

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.
Building 7, Shenzhen International Innovation Valley, Dashi Road 1, Nanshan, 518055 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

Design and Development, and Distribution of Syringe Pumps, Infusion Pumps, Infusion Workstations, Infusion Central Monitoring Systems, DVT Preventive Pumps, Infusion Docking Stations, Thromboelastograph Instruments and Reagents, Vein Illuminators, Enteral Feeding Pumps, Video Laryngoscopes, Chemiluminescence Immunoassay Analyzers and Reagents, Hemostasis Reagents, Colloidal Gold Reagents, Blood Grouping Systems, Spirometers, Nucleic Acid Extraction Systems, and Oxygen Concentrators.

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.
Building C, Jiale Science and Technology Industrial Park, Matian Street, Guangming, 518106 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

Production of Syringe Pumps, Infusion Pumps, Infusion Workstations, Infusion Central Monitoring Systems, DVT Preventive Pumps, Infusion Docking Stations, Thromboelastograph Instruments and Reagents, Vein Illuminators, Enteral Feeding Pumps, Video Laryngoscopes, Chemiluminescence Immunoassay Analyzers and Reagents, Hemostasis Reagents, Colloidal Gold Reagents, Blood Grouping Systems, Spirometers, Nucleic Acid Extraction Systems, and Oxygen Concentrators.

-

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

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Inspire trust.

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MEDCAPTAIN MEDICAL
TECHNOLOGY CO., LTD.
12th Floor, Baiwang Research Building
No.5158 Shahe West Road
Xili, Nanshan
518055 SHENZHEN, GUANGDONG
PEOPLE'S REPUBLIC OF CHINA

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
86916	713336204 | 713257146	medical_devices@tuvsud.com	N/A	2024-05-15	1 of 6

TÜV SÜD Product Service GmbH
Confirmation Letter
CL 086916 0056 Rev. 00

Reference: 713336204 | 713257146

To whom it may concern,

Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 (in the following referenced as MDR) as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

With this letter TÜV SÜD Product Service GmbH, designated under MDR and identified by the number 0123 on NANDO, confirms that we have received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the above stated manufacturer with the following SRN Number:

SRN Number: CN-MF-000005925

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below.

- Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive.
- Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but TÜV SÜD Product Service GmbH has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

If devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that:

- the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or
- provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively.

The transition timelines in accordance Article 120 (3) of MDR that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120 (3c) of MDR, are shown below:

- 26 May 2026 for Class III custom-made implantable devices
- 31 December 2027 for Class III devices and Class IIb implantable devices (except sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition, measuring function.
- 31 December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

The issuance of the first confirmation letter is free of charge. We reserve the right to invoice further copies, amendments and / or changes of the confirmation letter according to effort.

For certificate validity see www.tuvsud.com/ps-cert?q=CL 086916 0056 Rev. 00

In case of inquiries please contact medical_devices@tuvsud.com.

On behalf of the Notified Body TÜV SÜD Product Service GmbH,

15th May 2024.

TÜV SÜD Product Service GmbH
Medical and Health Services

Ben Xu
Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH
Medical and Health Services

Tunde Junaid
Application Reviewer

Table 1: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under MDR application)	**MDR Device classification (as proposed by the manufacturer and verified during application review)**	**If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device**	**MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification**
Syringe Pump 69268026MP3000000000001KB	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Syringe Pump 69268026HP30000001NE	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Syringe Pump 69268026SYS500000000001MW	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Pump 69268026MP6000000000001P6	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
	☐ Class I devices with measuring function ☐ Class III implantable custom-made-device		granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Pump 69268026SYS601000000001PY	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Pump 69268026SYS700000000001Q8	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Workstation 69268026MP8000000000001RQ	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Workstation 69268026HP8000000000001JD	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or

Device name or Basic UDI-DI (under MDR application)	**MDR Device classification (as proposed by the manufacturer and verified during application review)**	**If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device**	**MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification**
	☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	Individual Article number:	☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Workstation 69268026HP80MRI00000001UU	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Infusion Workstation 69268026MS1000000000001LS	☐ Class III ☐ Class IIb implantable (non-exempted) ☒ Class IIb / Class IIb implantable (exempted) ☐ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
DVT Preventive Pump 69268026TP2000000000001UB	☐ Class III ☐ Class IIb implantable (non-exempted) ☐ Class IIb / Class IIb implantable (exempted) ☒ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☒ N/A or ☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or ☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#
Enteral Feeding Pump 69268026EP6000000000001BE	☐ Class III ☐ Class IIb implantable (non-exempted) ☐ Class IIb / Class IIb implantable (exempted)	☒ N/A or	☒ Certification as follows: Certificate G1 086916 0023 Rev.01; NB0123 or

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
	☒ Class IIa ☐ Class I devices in sterile condition ☐ Class I devices with measuring function ☐ Class III implantable custom-made-device	☐ Identification of the corresponding device under MDD/AIMDD Individual Article number:	☐ Evidence that a competent authority of a Member State had granted acc. MDR, Art.59 (1) or Art.97 (1) Evidence #1; CA# Evidence #2; CA#

Table 2: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive: N/A

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification

Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024/05/15	713336204 | 713257146	Initial issue

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 086916 0023 Rev. 01

Manufacturer:

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.
12th Floor, Baiwang Research Building
No.5158 Shahe West Road
Xili, Nanshan
518055 Shenzhen, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Syringe Pump, Infusion Pump, Infusion Workstation, DVT Preventive Pump, Enteral Feeding Pump

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G1 086916 0023 Rev. 01

Report No.: GZ2111701

Valid from: 2021-05-18
Valid until: 2024-05-26

Date, 2021-05-18

Christoph Dicks
Head of Certification/Notified Body

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

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

TÜV®

CERTIFICATE OF QUALITY MANAGEMENT SYSTEM CERTIFICATION

Certificate No.016XZ23Q30394R0M
Unified social credit code:91440300573106363N

This is to Certify that the Quality Management System of

MEDCAPTAIN MEDICAL TECHNOLOGY CO.,LTD

is in conformity with
GB/T19001-2016 idt ISO9001:2015 Standard, applies to

SYRINGE PUMPS, INFUSION PUMPS, INFUSION WORKSTATIONS,INFUSION CENTRAL MONITORING SYSTEMS,DVT PREVENTIVE PUMPS, THROMBOELASTOGRAPH INSTRUMENTS AND REAGENTS, VEIN IIIUMINATORS, ENTERAL FEEDING PUMPS, VIDEO LARYNGOSCOPES, CHEMILUMINESCENCE IMMUNOASSAY ANALYZERS AND REAGENTS, HEMOSTASIS REAGENTS, BLOOD GROUPING SYSTEM, SPIROMETERS,NUCLEIC ACID EXTRACTION SYSTEMS, AND OXYGEN CONCENTRATORS(WITHIN THE SCOPE OF QUALIFICATION)
REGISTERED ADDRESS:12TH FLOOR, BAIWANG RESEARCH BUILDING NO.5158 SHAHE WEST ROAD XILI, NANSHAN 518055 SHENZHEN,GUANGDONG PEOPLE'S REPUBLIC OF CHINA
OPERATION ADDRESS:12TH FLOOR, BAIWANG RESEARCH BUILDING NO.5158 SHAHE WEST ROAD XILI, NANSHAN 518055 SHENZHEN,GUANGDONG PEOPLE'S REPUBLIC OF CHINA (DISTRIBUTION); BUILDING 7, SHENZHEN INTERNATIONAL INNOVATION VALLEY, DASHI ROAD 1, NANSHAN, SHENZHEN,GUANGDONG, PEOPLE'S REPUBLIC OF CHINA(DESIGN AND DEVELOPMENT, AND DISTRIBUTION); BUILDING C,JIALE AND TECHNOLOGY INDUSTRIAL PARK, MATIAN STREET, GUANGMING, SHENZHEN, GUANGDONG,PEOPLE'S REPUBLIC OF CHINA(PRODUCTION)

Date of Initial Issuance: Mar. 3, 2023
Date of Expiration: Mar. 2, 2026

BCC Inc.
President:

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The information of this certificate available for inquiry on CNCA's website: www.cnca.gov.cn.

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