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TÜVRheinland®

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

Boen Healthcare Co., Ltd.
Unit 602, International Center
No. 535, Shenxu Road
215021 Suzhou, Jiangsu
China

has established and applies a quality management system for medical devices for the following scope:

Manufacture and Distribution of Medical Devices

(see attachment for products included)

Proof has been furnished that the requirements specified in

EN ISO 13485:2016

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date:	2019-08-07
Certificate Registration No.:	SX 60138020 0001
An audit was performed. Report No.:	15092074 004
This Certificate is valid until:	2022-02-27

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

Date 2019-08-07

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail.cert-validity@de.tuv.com http://www.tuv.com/safety

EC Certificate

TÜVRheinland®

Production Quality Assurance
MDD Annex V

Registration No.: DD 2063008-1

Manufacturer:

Boen Healthcare Co., Ltd.
Unit 602, International Center, No. 535, Shenxu Road,
Suzhou,
215021 Jiangsu
P.R. China

Medical Brushes, Disposable Vaginal Speculums, Disposable Gynecological Sets,
Disposable Dressing Kits, Disposable Colostomy Bags, Disposable Umbilical Cord Clamps, Disposable Urine Drainage Bags, Sterile Wooden Tongue Depressors, Non Woven Surgical Drapes, Non Woven Surgical Gowns, X-ray Detectable Gauze Swabs (Sponges), Gauze Balls and Lap Sponges in Sterilization Packing, Gauze Swabs (Sponges), Gauze Balls Gauze Bandages and Non Woven Wound Care Products, Medical Elastic Bandages, First Aid Kits and Its Related Products, Disposable Nasal Speculums, Disposable Ear Checkers, Disposable Oral Cavity Kits and Implements, Sterile Urine Meters;
Aspects of manufacture concerned with conformity of products with metrological requirements: Sphygmomanometers, Mercury-free Clinical Thermometers

Replaces Approval, Registration No.: DD 60142274 0001

Report No.: 15092074 009

Effective date: 2020-11-18

Expiry date: 2024-05-26

Issue date: 2020-11-18

TÜV Rheinland LGA Products GmbH

Jason Pan
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

EC Certificate

TÜVRheinland®

Production Quality Assurance
MDD Annex V

Registration No.: DD 2063008-1

Manufacturer: Boen Healthcare Co., Ltd.
Unit 602, International Center, No. 535, Shenxu Road,
Suzhou,
215021 Jiangsu
P.R. China

Products: Nasal Oxygen Cannulae, Suction Catheters, Stomach Tubes, Feeding Tubes, Suction Connecting Tubes with Yankauer, Sterile Latex Surgical Gloves, Disposable Surgical Blades & Scalpels With Plastic Handle, Sterile Blood Lancets, Disposable Syringes, Disposable Infusion Sets, Disposable Transfusion Sets, Intravenous Needles for Single Use, Sterile Hypodermic Needles for Single Use, Disposable Tracheal Tubes (Standard & Reinforced), Disposable Oxygen Masks, Non-Rebreathing Masks, Aerosol Masks, Closed Suction Catheters, Tracheostomy Tubes, Laryngeal Mask Devices, Disposable Air Cushion Face Masks, Disposable Breathing Circuits, Oropharyngeal Airways, Venturi Masks, Self-destruction Safety Syringes, Blood Collecting Needles, Foley Catheters, Disposable Acupuncture Needles, Three-way Stopcocks (with Extension Tube), Nelaton Catheters, Insulin Needles for Single Use, Wound Drainage System with and without Trocars, Needle Free Connectors, Digital Thermometers, Humidifier Jar (Bubble Humidifier Jar), Enteral Feeding Sets (Bag);
Aspects of manufacture concerned with securing and maintaining sterile conditions: Sterile Hemostasis Adhesive Dressing Series (Sterile Wound Plaster, Liquid Transfusion Plaster and Adhesive Dressing), Disposable

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Report No.: 15092074 009

Effective date: 2020-11-18

Expiry date: 2024-05-26

Issue date: 2020-11-18

TÜV Rheinland LGA Products GmbH

Jason Pan
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

TÜVRheinland®

Doc. 2/2, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: SX 60138020 0001
Report No.: 15092074 004

Organization:
Boen Healthcare Co., Ltd.
Unit 602, International Center
No. 535, Shenxu Road
215021 Suzhou, Jiangsu
China

Scope:

Products:
Gauze Swabs (Sponges), Gauze Balls Gauze Bandages and Non Woven Wound Care Products, Medical Elastic Bandages, First Aid Kits and Its Related Products, Tracheostomy Tubes, Laryngeal Mask Devices, Disposable Air Cushion Face Masks, Disposable Breathing Circuits, Oropharyngeal Airways, Venturi Masks, Self-destruction Safety Syringes, Blood Collecting Needles, Foley Catheters, Disposable Acupuncture Needles, Three-way Stopcocks (with Extension Tube), Nelaton Catheters, Insulin Needles for Single Use, Humidifier Jar (Bubble Humidifier Jar), Wound Drainage System with and without Trocars, Sterile Urine Meters, Needle Free Connectors, Disposable Nasal Speculums, Disposable Ear Checkers, Enteral Feeding Sets (Bag), Disposable Oral Cavity Kits and Implements, Mercury-free Clinical Thermometers, Digital Thermometers

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Date: 2019-08-07

Fuxiu Sheng

TÜVRheinland®

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: SX 60138020 0001
Report No.: 15092074 004

Organization:
Boen Healthcare Co., Ltd.
Unit 602, International Center
No. 535, Shenxu Road
215021 Suzhou, Jiangsu
China

Scope:

Products:
Sterile Hemostasis Adhesive Dressing Series (Sterile Wound Plaster, Liquid Transfusion Plaster and Adhesive Dressing), Sphygmomanometers, Disposable Medical Brushes, Disposable Vaginal Speculums, Disposable Gynecological Sets, Disposable Dressing Kits, Disposable Colostomy Bags, Disposable Umbilical Cord Clamps, Nasal Oxygen Cannulae, Suction Catheters, Stomach Tubes, Feeding Tubes, Suction Connecting Tubes with Yankauer, Disposable Urine Drainage Bags, Sterile Wooden Tongue Depressors, Sterile Latex Surgical Gloves, Disposable Surgical Blades & Scalpels With Plastic Handle, Sterile Blood Lancets, Disposable Syringes, Disposable Infusion Sets, Disposable Transfusion Sets, Intravenous Needles for Single Use,Sterile Hypodermic Needles for Single Use, Disposable Tracheal Tubes(Standard & Reinforced), Disposable Oxygen Mask, Non-Rebreathing Masks, Aerosol Masks, Closed Suction Catheters, Non Woven Surgical Drapes, Non Woven Surgical Gowns, X-ray Detectable Gauze Swabs (Sponges), Gauze Balls and Lap Sponges in sterilization Packing

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

Date: 2019-08-07

TÜV Rheinland LGA Products GmbH TÜVRheinland® Zertifizierungsstelle

Fuxiu Sheng

TÜVRheinland®

Doc. 2/2, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: SX 60138020 0001
Report No.: 15092074 004

Organization:
Boen Healthcare Co., Ltd.
Unit 602, International Center
No. 535, Shenxu Road
215021 Suzhou, Jiangsu
China

Scope:

Products:
Gauze Swabs (Sponges), Gauze Balls Gauze Bandages and Non Woven Wound Care Products, Medical Elastic Bandages, First Aid Kits and Its Related Products, Tracheostomy Tubes, Laryngeal Mask Devices, Disposable Air Cushion Face Masks, Disposable Breathing Circuits, Oropharyngeal Airways, Venturi Masks, Self-destruction Safety Syringes, Blood Collecting Needles, Foley Catheters, Disposable Acupuncture Needles, Three-way Stopcocks (with Extension Tube), Nelaton Catheters, Insulin Needles for Single Use, Humidifier Jar (Bubble Humidifier Jar), Wound Drainage System with and without Trocars, Sterile Urine Meters, Needle Free Connectors, Disposable Nasal Speculums, Disposable Ear Checkers, Enteral Feeding Sets (Bag), Disposable Oral Cavity Kits and Implements, Mercury-free Clinical Thermometers, Digital Thermometers

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Date: 2019-08-07

Fuxiu Sheng

EC Certificate

TÜVRheinland®

Production Quality Assurance
MDD Annex V

Registration No.: DD 2063008-1

Manufacturer: Boen Healthcare Co., Ltd.
Unit 602, International Center, No. 535, Shenxu Road,
Suzhou,
215021 Jiangsu
P.R. China

Products: Nasal Oxygen Cannulae, Suction Catheters, Stomach Tubes, Feeding Tubes, Suction Connecting Tubes with Yankauer, Sterile Latex Surgical Gloves, Disposable Surgical Blades & Scalpels With Plastic Handle, Sterile Blood Lancets, Disposable Syringes, Disposable Infusion Sets, Disposable Transfusion Sets, Intravenous Needles for Single Use, Sterile Hypodermic Needles for Single Use, Disposable Tracheal Tubes (Standard & Reinforced), Disposable Oxygen Masks, Non-Rebreathing Masks, Aerosol Masks, Closed Suction Catheters, Tracheostomy Tubes, Laryngeal Mask Devices, Disposable Air Cushion Face Masks, Disposable Breathing Circuits, Oropharyngeal Airways, Venturi Masks, Self-destruction Safety Syringes, Blood Collecting Needles, Foley Catheters, Disposable Acupuncture Needles, Three-way Stopcocks (with Extension Tube), Nelaton Catheters, Insulin Needles for Single Use, Wound Drainage System with and without Trocars, Needle Free Connectors, Digital Thermometers, Humidifier Jar (Bubble Humidifier Jar), Enteral Feeding Sets (Bag);
Aspects of manufacture concerned with securing and maintaining sterile conditions: Sterile Hemostasis Adhesive Dressing Series (Sterile Wound Plaster, Liquid Transfusion Plaster and Adhesive Dressing), Disposable

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Report No.: 15092074 009

Effective date: 2020-11-18

Expiry date: 2024-05-26

Issue date: 2020-11-18

TÜV Rheinland LGA Products GmbH

Jason Pan
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

EC Certificate

TÜVRheinland®

Production Quality Assurance
MDD Annex V

Registration No.: DD 2063008-1

Manufacturer:

Boen Healthcare Co., Ltd.
Unit 602, International Center, No. 535, Shenxu Road,
Suzhou,
215021 Jiangsu
P.R. China

Medical Brushes, Disposable Vaginal Speculums, Disposable Gynecological Sets,
Disposable Dressing Kits, Disposable Colostomy Bags, Disposable Umbilical Cord Clamps, Disposable Urine Drainage Bags, Sterile Wooden Tongue Depressors, Non Woven Surgical Drapes, Non Woven Surgical Gowns, X-ray Detectable Gauze Swabs (Sponges), Gauze Balls and Lap Sponges in Sterilization Packing, Gauze Swabs (Sponges), Gauze Balls Gauze Bandages and Non Woven Wound Care Products, Medical Elastic Bandages, First Aid Kits and Its Related Products, Disposable Nasal Speculums, Disposable Ear Checkers, Disposable Oral Cavity Kits and Implements, Sterile Urine Meters;
Aspects of manufacture concerned with conformity of products with metrological requirements: Sphygmomanometers, Mercury-free Clinical Thermometers

Replaces Approval, Registration No.: DD 60142274 0001

Report No.: 15092074 009

Effective date: 2020-11-18

Expiry date: 2024-05-26

Issue date: 2020-11-18

TÜV Rheinland LGA Products GmbH

Jason Pan
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

TÜVRheinland®

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

Boen Healthcare Co., Ltd.
Unit 602, International Center
No. 535, Shenxu Road
215021 Suzhou, Jiangsu
China

has established and applies a quality management system for medical devices for the following scope:

Manufacture and Distribution of Medical Devices

(see attachment for products included)

Proof has been furnished that the requirements specified in

EN ISO 13485:2016

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date:	2019-08-07
Certificate Registration No.:	SX 60138020 0001
An audit was performed. Report No.:	15092074 004
This Certificate is valid until:	2022-02-27

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

Date 2019-08-07

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail.cert-validity@de.tuv.com http://www.tuv.com/safety

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Stool Container**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Sputum Specimen Container**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD** **Unit 602, International Center, No.535, Shenxu Road,** **Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Centrifuge Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD** **(Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /** **EG Annex III (expect point 6) of IVDD 98/79/EC** **Annexe III (sauf le point 6) de l'IVDD 98/79 / CE** **Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**

Registrier-Nr.: /
Registration No.: /
N°d’enregistrement: /
Numero di registrazione:

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date /
Lieu, date / Luogo, data

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD** **Unit 602, International Center, No.535, Shenxu Road,** **Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Glass Test Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD** **(Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /** **EG Annex III (expect point 6) of IVDD 98/79/EC** **Annexe III (sauf le point 6) de l'IVDD 98/79 / CE** **Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**

Registrier-Nr.: /
Registration No.: /
N°d’enregistrement: /
Numero di registrazione:

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date /
Lieu, date / Luogo, data

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Eppendorf Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

BOEN HEALTHCARE CO.,LTD 苏州博恩贸易有限公司 3205011038530

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD** **Unit 602, International Center, No.535, Shenxu Road,** **Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Non Vacuum Blood Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD** **(Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /** **EG Annex III (expect point 6) of IVDD 98/79/EC** **Annexe III (sauf le point 6) de l'IVDD 98/79 / CE** **Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**

Registrier-Nr.: /
Registration No.: /
N°d’enregistrement: /
Numero di registrazione:

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date /
Lieu, date / Luogo, data

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Microscope Slide**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Microscope Cover Glass**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Micro Blood Collection with Capillary Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD** **Unit 602, International Center, No.535, Shenxu Road,** **Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Mechanical Autoclavable Pipette**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD** **(Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /** **EG Annex III (expect point 6) of IVDD 98/79/EC** **Annexe III (sauf le point 6) de l'IVDD 98/79 / CE** **Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	
Benannte Stelle: / Notified Body: / Organisme notifié: / Organismo notificato:	

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data	Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Serological Pipette**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

BOEN HEALTHCARE CO.,LTD 苏州博恩贸易有限公司 3205011038530

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: /
Name and address of the manufacturer: /
Nom et adresse du fabricant: /
Nome e indirizzo del fabbricante:

BOEN HEALTHCARE CO., LTD
Unit 602, International Center, No.535, Shenxu Road,
Suzhou, 215021, Jiangsu, China

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that /
Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: /
the medical device: /
le dispositif médical: /
il dispositivo medico:

Graduated Pasteur Pipette

der Klasse: /
of class: /
de la classe: /
di classe:

Common/Others IVD
(Devices of NOT Annex II and NOT self-test)

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: /
Conformity assessment procedure: /
Procédure d’évaluation de la conformité: /
Procedura di valutazione della conformità:

Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /
EG Annex III (expect point 6) of IVDD 98/79/EC
Annexe III (sauf le point 6) de l'IVDD 98/79 / CE
Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE

Registrier-Nr.: /
Registration No.: /
N°d’enregistrement: /
Numero di registrazione:

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

Ort, Datum / Place, date /
Lieu, date / Luogo, data

General Manager

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Gilson Pipette Tips**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	
Benannte Stelle: / Notified Body: / Organisme notifié: / Organismo notificato:	

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data	Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Filtered Pipette Tips**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

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Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Vacuum Blood Collection Tube**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD** **(Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II
(IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

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soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 /** **EG Annex III (expect point 6) of IVDD 98/79/EC** **Annexe III (sauf le point 6) de l'IVDD 98/79 / CE** **Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**

Registrier-Nr.: /
Registration No.: /
N°d’enregistrement: /
Numero di registrazione:

Benannte Stelle: /
Notified Body: /
Organisme notifié: /
Organismo notificato:

CE

Suzhou, 201.05.26

General Manager

Ort, Datum / Place, date /
Lieu, date / Luogo, data

Name und Funktion / Name and function /
Nom et fonction / Nome e funzione

Stool Container

Stool Container

Designed to collect and store solid or liquid samples, including food, medicine, urine and feces, opaque containers ideal for photosensitive samples or when it is required not to show the content.

With screw cap and spoon

Cat. No.	Description	Qty/case (pcs)
611201	30ml stool container, screw cap, PP, with label, sterile, individual pack	1000
611202	30ml stool container, screw cap, PP, with label, non-sterile	1000
611203	30ml stool container, screw cap, PP, writing surface, sterile, individual pack	1000
611204	30ml stool container, screw cap, PP, writing surface, non-sterile	1000

Sputum Specimen Container

Sputum Specimen Container

Manufactured from virgin ultra-clear polypropylene ensures good samples visibility
Cap Material: Polypropylene
Ideal for Sputum Specimen
Non sterile

Cat. No.	Description	Qty/case (pcs)
613201	Sputum specimen container 30ml, bulk pack	1000
613202	Sputum specimen container 30ml, 10pcs/sleeve bag	1000

50ml Plastic Centrifuge Tube

50ml Plastic Centrifuge Tube

Used for the general laboratory centrifugation
Made of PP material, chemical stability
Easy to read and view tube contents
One-hand operation, easy to coil cover, sealing well.
Self-standing bottom, conical bottom or round bottom with bayonet.

Cat. No.	Description	Qty/case (pcs)
660801	50ml self-standing bottom screw cap centrifuge tube	800
660802	50ml conical bottom screw cap centrifuge tube	800
660803	50ml round bottom with bayonet centrifuge tube	800
660804	50ml self-standing bottom screw cap centrifuge tube, sterile	800
660805	50ml conical bottom screw cap centrifuge tube, sterile	800
660806	50ml round bottom with bayonet centrifuge tube, sterile	800

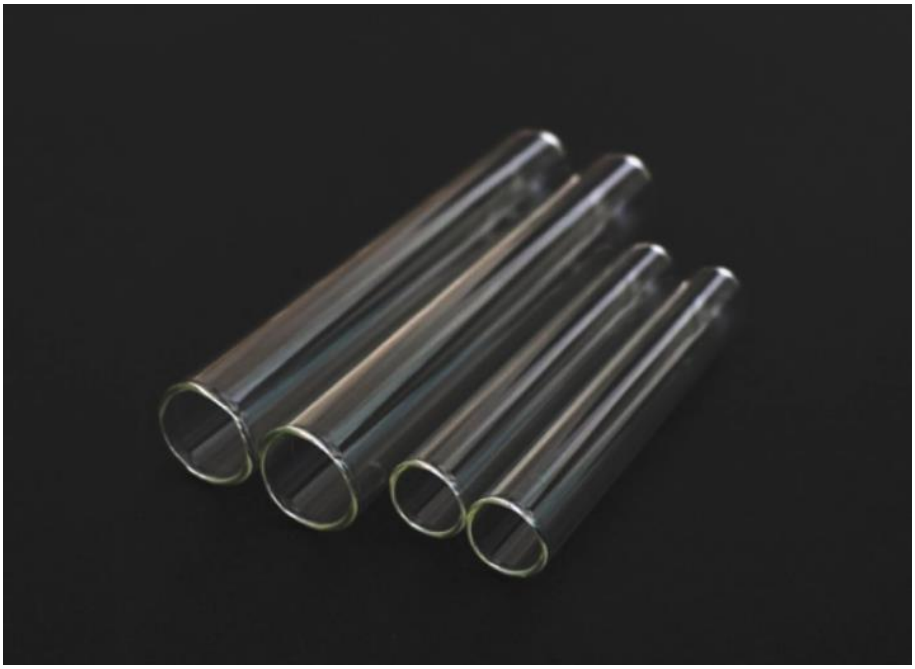
Plastic Test Tube

Plastic Test Tube

Plastic test tube, also known as a culture tube or sample tube.
Disposable round bottom tubes no rim, for laboratory use.
Multiple size and type to meet various test requirements.
Material: PS/ PP.

Cat No.	Description	Qty/Case
670501	Test tube, PS/PP, 12x60mm	5000
670502	Test tube, PS/PP, 12x75mm	5000
670503	Test tube, PS/PP, 13x75mm	5000
670504	Test tube, PS/PP, 12x100mm	4000
670505	Test tube, PS/PP, 13x100mm	4000
670506	Test tube, PS/PP, 15x100mm	3000
670507	Test tube, PS/PP, 16x100mm	2800
670508	Test tube, PS/PP, conical bottom 16x102mm	2500
670509	Test tube, PS/PP, 17x100mm	240

Glass Test Tube



Glass Test Tube

Test tube, glass test tube, glass culture tube made of borosilicate glass. The tubes provide excellent chemical resistance and durability.
Made of superior quality 3.3 borosilicate glass tubes, suitable for all usual laboratory applications, can withstand high temperature.

Material: 3.3 borosilicate glass or borosilicate glass.
Mouth: plain mouth
Bottom: round bottom
Size: 10x50mm, 10x75mm, 11x70mm, 12x75mm, 12x100mm, 13x100mm, 12x110mm, 12x120mm, 15x100mm, 15x125mm, 16x125mm, 16x100mm, 16x160mm, 20x150mm, 18x150mm, 20x200mm or customized.

Cat No.	Size	Wall thick(mm)	Qty/Case (PCS)
Glass test tube, borosilicate glass, plain, round bottom			
670701	Ø10x50mm,2.5ml	0.8-1.0mm	2000
670702	Ø10x75mm,3.0ml	0.8-1.0mm	2000
670703	Ø11x70mm,3.0ml	0.8-1.0mm	2000
670704	Ø12x75mm,5.0ml	0.8-1.0mm	2000
670705	Ø12x100mm,7.0ml	0.8-1.0mm	2000
670706	Ø13x100mm,7.0ml	0.8-1.0mm	2000
670707	Ø12x110mm,8.0ml	0.8-1.0mm	1000
670708	Ø12x120mm,8.0ml	0.8-1.0mm	1000
670709	Ø15x100mm,10ml	0.8-1.0mm	1000

670710	Ø15x125mm,12ml	0.8-1.0mm	1000
670711	Ø16x125mm,14ml	0.8-1.0mm	1000
670712	Ø16x100mm,10ml	1.0mm	1000
670713	Ø16x150mm,16ml	1.0mm	1000
670714	Ø16x160mm,17ml	1.0mm	1000
670715	Ø20x150mm,20ml	1.0-1.2mm	600
670716	Ø18x200mm,25ml	1.0-1.2mm	600
670717	Ø20x200mm,25ml	1.0-1.2mm	600
Glass test tube, 3.3 borosilicate glass, plain, round bottom			
670801	Ø10x50mm,2.5ml	0.8-1.0mm	2000
670802	Ø10x75mm,3.0ml	0.8-1.0mm	2000
670803	Ø11x70mm,3.0ml	0.8-1.0mm	2000
670804	Ø12x75mm,5.0ml	0.8-1.0mm	2000
670805	Ø12x100mm,7.0ml	0.8-1.0mm	2000
670806	Ø13x100mm,7.0ml	0.8-1.0mm	2000
670807	Ø12x110mm,8.0ml	0.8-1.0mm	1000
670808	Ø12x120mm,8.0ml	0.8-1.0mm	1000
670809	Ø15x100mm,10ml	0.8-1.0mm	1000
670810	Ø15x125mm,12ml	0.8-1.0mm	1000
670811	Ø16x125mm,14ml	0.8-1.0mm	1000
670812	Ø16x100mm,10ml	1.0mm	1000
670813	Ø16x150mm,16ml	1.0mm	1000
670814	Ø16x160mm,17ml	1.0mm	1000
670815	Ø20x150mm,20ml	1.0-1.2mm	600
670816	Ø18x200mm,25ml	1.0-1.2mm	600
670817	Ø20x200mm,25ml	1.0-1.2mm	600

Eppendorf Tube

Eppendorf Tube

Made of PP material, resistance to most chemicals regents and autoclavable.
With snap cap, tight-fitting lid provides a leak-resistant seal.
Flat top, and with large frosted writing area for easy sample identification.
Molded graduation for volume estimation.

Cat. No.	Description	Qty/Case (Pcs)
660201	1.5ml microcentrifuge tube, snap cap, 500pcs/bag	10000
660202	1.5ml microcentrifuge tube, snap cap, 500pcs/bag, sterile	10000
660401	2ml microcentrifuge tube, snap cap, 500pcs/bag	10000
660402	2ml microcentrifuge tube, snap cap, 500pcs/bag, sterile	10000

Non Vacuum Blood Tube

Non Vacuum Blood Tube

Cat. No.	Description	Qty/case (pcs)
631601	Non Vacuum Blood Tube, Plain, 1ml	1200
631602	Non Vacuum Blood Tube, Plain, 2ml	1200
631603	Non Vacuum Blood Tube, Plain, 3ml	1200
631604	Non Vacuum Blood Tube, Plain, 4ml	1200
631605	Non Vacuum Blood Tube, Plain, 5ml	1200
631606	Non Vacuum Blood Tube, EDTA K3, 1ml	1200
631607	Non Vacuum Blood Tube, EDTA K3, 2ml	1200
631608	Non Vacuum Blood Tube, EDTA K3, 3ml	1200
631609	Non Vacuum Blood Tube, EDTA K3, 4ml	1200
631610	Non Vacuum Blood Tube, EDTA K3, 5ml	1200
631611	Non Vacuum Blood Tube, 3.2% Sodium Citrate, 1ml	1200
631612	Non Vacuum Blood Tube, 3.2% Sodium Citrate, 2ml	1200
631613	Non Vacuum Blood Tube, 3.2% Sodium Citrate, 3ml	1200
631614	Non Vacuum Blood Tube, 3.2% Sodium Citrate, 4ml	1200
631615	Non Vacuum Blood Tube, 3.2% Sodium Citrate, 5ml	1200
631616	Non Vacuum Blood Tube, Heparin Lithium, 1ml	1200
631617	Non Vacuum Blood Tube, Heparin Lithium, 2ml	1200
631618	Non Vacuum Blood Tube, Heparin Lithium, 3ml	1200
631619	Non Vacuum Blood Tube, Heparin Lithium, 4ml	1200

631620	Non Vacuum Blood Tube, Heparin Lithium, 5ml	1200
631621	Non Vacuum Blood Tube, Fluoride, 1ml	1200
631622	Non Vacuum Blood Tube, Fluoride, 2ml	1200
631623	Non Vacuum Blood Tube, Fluoride, 3ml	1200
631624	Non Vacuum Blood Tube, Fluoride, 4ml	1200
631625	Non Vacuum Blood Tube, Fluoride, 5ml	1200
631626	Non Vacuum Blood Tube, Heparin Sodium, 1ml	1200
631627	Non Vacuum Blood Tube, Heparin Sodium, 2ml	1200
631628	Non Vacuum Blood Tube, Heparin Sodium, 3ml	1200
631629	Non Vacuum Blood Tube, Heparin Sodium, 4ml	1200
631630	Non Vacuum Blood Tube, Heparin Sodium, 5ml	1200
631631	Non Vacuum Blood Tube, Clot Activator, 1ml	1200
631632	Non Vacuum Blood Tube, Clot Activator, 2ml	1200
631633	Non Vacuum Blood Tube, Clot Activator, 3ml	1200
631634	Non Vacuum Blood Tube, Clot Activator, 4ml	1200
631635	Non Vacuum Blood Tube, Clot Activator, 5ml	1200
631636	Non Vacuum Blood Tube, ESR 1:4, 1ml	1200
631637	Non Vacuum Blood Tube, ESR 1:4, 2ml	1200
631638	Non Vacuum Blood Tube, ESR 1:4, 3ml	1200
631639	Non Vacuum Blood Tube, ESR 1:4, 4ml	1200
631640	Non Vacuum Blood Tube, ESR 1:4, 5ml	1200
631641	Non Vacuum Blood Tube, Gel & Clot Activator, 1ml	1200
631642	Non Vacuum Blood Tube, Gel & Clot Activator, 2ml	1200
631643	Non Vacuum Blood Tube, Gel & Clot Activator, 3ml	1200
631644	Non Vacuum Blood Tube, Gel & Clot Activator, 4ml	1200

Vacuum Blood Collection Tube

Vacuum Blood Collection Tube, Gel & Clot Activator Tube

Material: PET, Glass
Color: Yellow

Cat. No.	Specification	Volume	Additive	Qty/case (Glass)	Qty/case (PET)
630201	13×75mm	3ml	Gel & Clot Activator	1800	1800
630202	13×75mm	4ml	Gel & Clot Activator	1800	1800
630203	13×75mm	5ml	Gel & Clot Activator	1800	1800
630204	13X100mm	5ml	Gel & Clot Activator	1200	1200
630205	13×100mm	6ml	Gel & Clot Activator	1200	1200
630206	13×100mm	8ml	Gel & Clot Activator	1200	1200
630207	16x100mm	8ml	Gel & Clot Activator	1200	1200
630208	16×100mm	9ml	Gel & Clot Activator	1200	1200
630209	16×100mm	10ml	Gel & Clot Activator	1200	1200

Microscope Slide

Microscope Slide

Slide with 90° Corner, 45° Clipped Corner, both beveled edges and clipped corners.
50pcs/box and 72pcs/box are available.

Cat. No.	Description	Dim. (mm)	Thickness (mm)	Qty/Case
7101	Ground Edges	25.4×76.2 (1''×3'')	1-1.2	50
7102	Cut Edges	25.4×76.2 (1''×3'')	1-1.2	50
7103	Single Cavity, Ground Edges	25.4×76.2 (1''×3'')	1-1.2	50
7104	Double Cavities, Ground Edges	25.4×76.2 (1''×3'')	1-1.2	50
7105	Ground Edges, Frosted One End One Side	25.4×76.2 (1''×3'')	1-1.2	50
7105-1	Cut Edges, Frosted One End One Side	25.4×76.2 (1''×3'')	1-1.2	50
7106	Ground Edges, Frosted Two Ends One Side	25.4×76.2 (1''×3'')	1-1.2	50
7107	Ground Edges, Frosted One End Both Sides	25.4×76.2 (1''×3'')	1-1.2	50
7107-1	Cut Edges, Frosted One End Both Sides	25.4×76.2 (1''×3'')	1-1.2	50
7109	Ground Edges, Color Frosted One End One Side, White, Orange, Yellow, Green, Pink, Blue	25.4×76.2 (1''×3'')	1-1.2	50

Microscope Cover Glass

Microscope Cover Glass

Hinged-lid box wrapped in Individual Vacuum Tropical Pack. 200pcs/box. Or package of 10×100pcs/Vacuum Tropical Pack.

Cat. No.	Description	Thickness (mm)		Qty/Case (bxs)
		No.1	No.2	
620801	12×12mm	0.13-0.17	0.17-0.25	100
620802	14×14mm	0.13-0.17	0.17-0.25	100
620803	16×16mm	0.13-0.17	0.17-0.25	100
620804	18×18mm	0.13-0.17	0.17-0.25	100
620805	20×20mm	0.13-0.17	0.17-0.25	100
620806	22×22mm	0.13-0.17	0.17-0.25	100
620807	24×24mm	0.13-0.17	0.17-0.25	100
620808	24×32mm	0.13-0.17	0.17-0.25	100
620809	24×40mm	0.13-0.17	0.17-0.25	100
620810	24×50mm	0.13-0.17	0.17-0.25	100
620811	24×60mm	0.13-0.17	0.17-0.25	100

620812	∮ 12 mm	0.13-0.17	0.17-0.25	200
620813	∮ 13 mm	0.13-0.17	0.17-0.25	200
620814	∮ 16 mm	0.13-0.17	0.17-0.25	200
620815	∮ 18 mm	0.13-0.17	0.17-0.25	200
620816	∮ 19 mm	0.13-0.17	0.17-0.25	200
620817	∮ 20 mm	0.13-0.17	0.17-0.25	200
620818	∮ 22 mm	0.13-0.17	0.17-0.25	200

Micro Blood Collection with Capillary Tube

Micro Blood Collection with Capillary Tube

Mirco blood collection tube is mainly use for peripheral blood collection, it is specially suitable for small children and geriatriatric patient. The inner surface has an excellent hemorepellency where it is necessary for a good preparation of high quality blood sample.

Specification:
Material: medical grade PP
Tube Size: 8x45mm
Volume:0.25ml, 0.5ml
Type: Micro Blood Collection Tube

Features:
a. accurate test result
b. excellent performance additive
c. shelf time 24 months

Packages:100pcs/tray, 1000pcs/ctn

Cat. No.	Description	Qty/Case (Pcs)
631401	Micro Fluoride Oxalate tube, grey cap, 0.5ml	1000
631402	EDTA K2 tube, purple cap, 0.5ml	1000
631403	EDTA K3 tube, purple cap, 0.5ml	1000

631404	Lithium Heparin tube, Lithium Heparin, green cap, 0.5ml	1000
631405	Lithium Heparin tube, Sodium Heparin, green cap, 0.5ml	1000
631406	Micro plain tube, no additive, red cap, 0.5ml	1000
631407	Micro Fluoride Oxalate tube, grey cap, 0.25ml	1000
631408	EDTA K2 tube, purple cap, 0.25ml	1000
631409	EDTA K3 tube, purple cap, 0.25ml	1000
631410	Lithium Heparin tube, Lithium Heparin, green cap, 0.25ml	1000
631411	Lithium Heparin tube, Sodium Heparin, green cap, 0.25ml	1000
631412	Micro plain tube, no additive, red cap, 0.25ml	1000

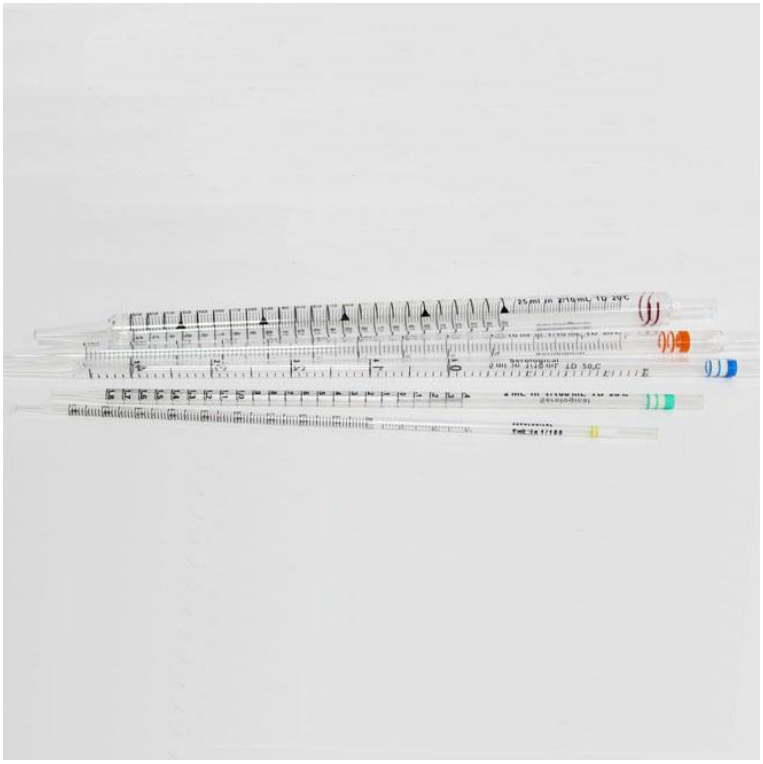
Mechanical Autoclavable Pipette

Mechanical Autoclavable Pipette

Adjustable single-channel Mechanical Pipette with a manual volume control for transporting a measured volume of liquid.
Fully autoclavable
Ergonomic design provides excellent operating experience
Easy-to-read volume display
The pipettes cover the volume range from 0.1μL to 5mL
Easy calibration and maintenance

Cat. No.	Description	Qty/case (pcs)
640901	Mechanical Autoclavable Pipette, single-channel, 0.1-2.5μL	130
640902	Mechanical Autoclavable Pipette, single-channel, 0.5-10μl	130
640903	Mechanical Autoclavable Pipette, single-channel, 2-20μl	130
640904	Mechanical Autoclavable Pipette, single-channel, 5-50μl	130
640905	Mechanical Autoclavable Pipette, single-channel, 10-100μl	130
640906	Mechanical Autoclavable Pipette, single-channel, 20-200μl	130
640907	Mechanical Autoclavable Pipette, single-channel, 50-200μl	130
640908	Mechanical Autoclavable Pipette, single-channel, 100-1000μl	130
640909	Mechanical Autoclavable Pipette, single-channel, 200-1000μl	130
640910	Mechanical Autoclavable Pipette, single-channel, 1000-5000μl	130
640911	Mechanical Autoclavable Pipette, single-channel, 500-5000μl	130

Serological Pipettes

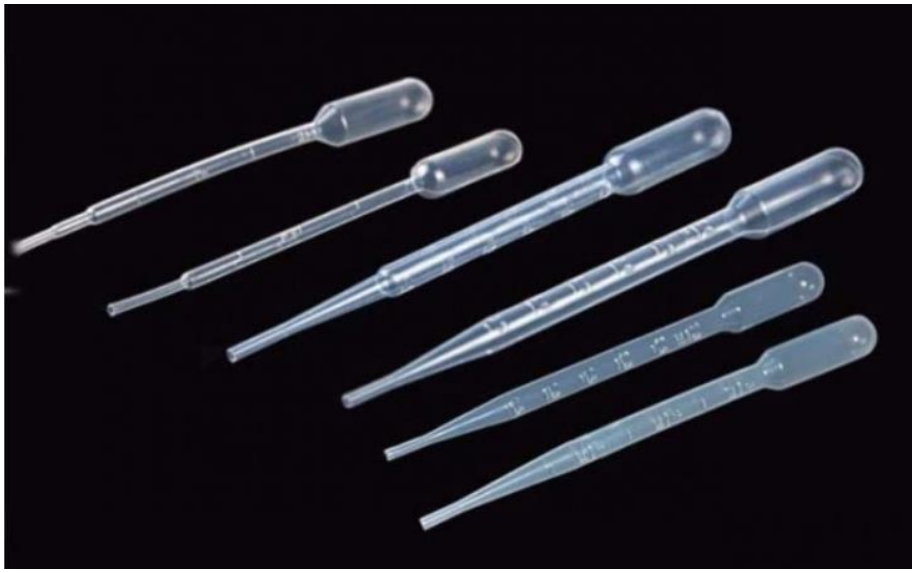


Serological Pipettes

Serological Pipette, designed for quantitatively transferring and dispensing exact volumes of liquid.
Made of PS material.
Accurate graduation, easy to read, color-codes by size for identification.
Available with 6 capacity of 1, 2, 5, 10, 25 and 50ml
Sterile or non-sterile
Package type: individual sterile pack, bulk pack.

Cat No.	Description	Qty/Case
640601	Serological Pipettes, PS, 1ml, yellow band color	4000
640602	Serological Pipettes, PS, 2ml, green band color	3200
640603	Serological Pipettes, PS, 5ml, blue band color	1800
640604	Serological Pipettes, PS, 10ml, red band color	1200
640605	Serological Pipettes, PS, 25ml, purple band color	800
640606	Serological Pipettes, PS, 50ml, black band color	800

Graduated Pasteur Pipette



Graduated Pasteur Pipette

Pasteur pipette, made from medical grade LDPE.
Graduated pasteur pipette, with moulded graduations make it easy to perform quick measured liquid transfer.
Transfer pipette, with variety of volume for choices. 1ml, 3ml, 5ml.
Available for E.O. sterilization or non sterile.
Bulk and individual packing or other.

Cat No.	Description	Qty/Case
640701	Pasteur pipette 1ml, with graduation, bulk pack	30000
640702	Pasteur pipette 1ml, with graduation, sterile, individual pack	30000
640703	Pasteur pipette 3ml, with graduation, bulk pack	5000
640704	Pasteur pipette 3ml, with graduation, sterile, individual pack	5000
640705	Pasteur pipette 5ml, with graduation, bulk pack	2000
640706	Pasteur pipette 5ml, with graduation, sterile, individual pack	2000
740707	Pasteur pipette 0.5ml, with graduation, bulk pack	10000
740708	Pasteur pipette 0.5ml, with graduation, sterile, individual pack	10000
740709	Pasteur pipette 2ml, with graduation, bulk pack	6000
740710	Pasteur pipette 2ml, with graduation, sterile, individual pack	6000

Gilson Pipette Tips

Gilson Pipette Tips

Designed for use in a wide variety of pipetting applications, compatibility with Gilson Pipettors.
PP material.
Size:10ul,200ul,1000ul,5000ul
Various color optional easy to identify
Non sterile.

Cat No.	Description	Qty/Case(bags)
640101	Gilson pipette tips,10ul, clear, 1000tips/bag	100
640102	Gilson pipette tips,200ul, yellow, 1000tips/bag	50
640103	Gilson pipette tips,1000ul, blue, 500tips/bag	50
640104	Gilson pipette tips,5000ul, clear, 300tips/bag	30

Filtered Pipette Tips

Filtered Pipette Tips

Filter pipette tips are used for protection against contamination.
Pipette tips with filter barrier protects pipettors and samples from contamination.
RNase, DNase and Pyrogen Free.
Filtered tips,Filter Tips Pipette, universal and fit most popular brands of pipettors.

Sterile pipette tips.
PP Material, clear color.
Volume:10ul-1250ul.
Package Style: Bulked: 1000tips/bag, 10bags/ctn
Racked: 96tips/rack, 10racks/pk, 50racks/ctn

Cat No.	Description	Qty/Case(pcs)
640401	Racked filter tips,96tips/rack, sterile, 10ul	4800
640402	Racked filter tips,96tips/rack, sterile, 10ul long	4800
640403	Racked filter tips,96tips/rack, sterile, 20ul	4800
640404	Racked filter tips,96tips/rack, sterile, 50ul	4800
640405	Racked filter tips,96tips/rack, sterile, 100ul	4800

640406	Racked filter tips,96tips/rack, sterile, 200ul	4800
640407	Racked filter tips,96tips/rack, sterile, 1000ul	4800
640408	Racked filter tips,96tips/rack, sterile, 1250ul (1000ul long)	4800
640409	Racked filter tips,96tips/rack, sterile, 200ul long	4800
640410	Racked filter tips,96tips/rack, sterile, 300ul	4800
640411	Bulked filter tips,1000tips/bag, sterile, 10ul	10000
640412	Bulked filter tips,1000tips/bag, sterile, 10ul long	10000
640413	Bulked filter tips,1000tips/bag, sterile, 20ul	10000
640414	Bulked filter tips,1000tips/bag, sterile, 50ul	10000
640415	Bulked filter tips,1000tips/bag, sterile, 100ul	10000
640416	Bulked filter tips,1000tips/bag, sterile, 200ul	10000
640417	Bulked filter tips,1000tips/bag, sterile, 1000ul	10000
640418	Bulked filter tips,1000tips/bag, sterile, 1250ul (1000ul long)	10000
640419	Bulked filter tips,1000tips/bag, sterile, 200ul long	10000
640420	Bulked filter tips,1000tips/bag, sterile, 300ul	10000

Urine Sediment Tube

Urine Sediment Tube

Urine Tube, PS, Sediment, Ø20xH103mm,12ml, Snap Cap
With flared opening, With sediment bulb, whale type
Graduated
Centrifuge spin rate:1500 RPM
Non sterile

Cat. No.	Description	Qty/Case
613001	Urine tube, PS, sediment, 12ml, tube only	2000
613002	Urine tube, PS, sediment, 12ml, with snap cap	2000

Vacuum Blood Collection Tube

Vacuum Blood Collection Tube, EDTA Tube

Material: PET, Glass

Color: Violet

Cat. No.	Specification	Volume	Additive	Qty/case (Glass)	Qty/case (PET)
630801	13×75mm	1ml	EDTAK2	1800	1800
630802	13×75mm	2ml	EDTAK2	1800	1800
630803	13×75mm	3ml	EDTAK2	1800	1800
630804	13×75mm	4ml	EDTAK2	1800	1800
630805	13×75mm	5ml	EDTAK2	1800	1800
630806	13×100mm	5ml	EDTAK2	1200	1200
630807	13×100mm	6ml	EDTAK2	1200	1200
630808	13×100mm	7ml	EDTAK2	1200	1200
630809	16×100mm	10ml	EDTAK2	1200	1200

630901	13×75mm	1ml	EDTAK3	1800	1800
630902	13×75mm	2ml	EDTAK3	1800	1800
630903	13×75mm	3ml	EDTAK3	1800	1800
630904	13×75mm	4ml	EDTAK3	1800	1800
630905	13×75mm	5ml	EDTAK3	1800	1800
630906	13×100mm	5ml	EDTAK3	1200	1200
630907	13×100mm	6ml	EDTAK3	1200	1200
630908	13×100mm	7ml	EDTAK3	1200	1200
630909	16×100mm	10ml	EDTAK3	1200	1200
631001	13×75mm	2ml	EDTANA2	1800	1800
631002	13×100mm	10ml	EDTANA2	1200	1800