



Notified Body Designated by  
 Europäische Kommission  
 für Gesundheitsprodukte  
 im Zusammenhang mit  
 Medizinprodukten  
 ZLC-05-244.10.08



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 081775 0008 Rev. 02**

**Manufacturer:**

**BMC Medical Co., Ltd.**

Room 110 Tower A Fengyu Building, No. 115 Fucheng Road  
 Haidian

100036 Beijing

PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):**

BMC Medical Co., Ltd.

Room 110 Tower A Fengyu Building, No. 115 Fucheng Road,  
 Haidian, 100036 Beijing, PEOPLE'S REPUBLIC OF CHINA

BMC (Tianjin) Medical Co., Ltd.

3/F, Building No.4, No.1 Xinxing Road, Wujing District, 301700  
 Tianjin, PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):**

**Masks; Tubes; Sleep Apnea Therapy Devices,  
 Respiratory Insufficiency Ventilators and  
 Accessories: CPAP, Auto CPAP, BPAP,  
 Humidifier; Heated Humidifier and Accessories:  
 Humidifier, Water Chamber, Nasal Cannula and  
 Tubes; Sleep Apnea Diagnosis Devices and  
 Accessories: Sleep Screener, Polysomnograph,  
 Portable Sleep Diagnostic System.**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

**Report No.:**

BJ19963043

**Valid from:**

2020-01-20

**Valid until:**

2023-03-31

**Date,**

2020-01-20

Christoph Dicks

Head of Certification/Notified Body

TÜV SÜD  
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT

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