EU Certificate

Production Quality Assurance
REGULATION (EU) 2017/745 on Medical Devices, Annex XI, Part A

TÜVRheinland®

Registration No.:	DZ 2174537-1
Manufacturer:	**Goodwood Medical Care Ltd.** 3-919 Jiulicun, Yongzheng Street, Jinzhou District, Dalian 116100 Liaoning P.R. China
EUDAMED Single Registration No.:	CN-MF-000016564
Products:	Products of Class I, Sterile: A99 - DEVICES FOR ADMINISTRATION, WITHDRAWAL AND COLLECTION - OTHERS - Sterile Tipped Applicators A1101 - SAMPLING COLLECTION NEUTRAL SWABS - Disposable sampling swabs - Transport Swab V9001 - TONGUE DEPRESSORS, SINGLE-USE - Sterile Tongue Depressors U089002 - DEVICES FOR GYNAECOLOGICAL CYTOLOGY - Cervical Scrapers - Cytology Brushes The scope of certification is limited to the aspects relating to establishing, securing and maintaining sterile conditions
Authorised representative(s):	CMC Medical Devices&Drugs S.L. C/ Horacio Lengo No18, CP 29006, Málaga, Spain

Certificate history		
Revision:	Description:	Issue date:
0	Initial Revision	2023-01-06

The Notified Body hereby declares that the requirements of Annex XI, Part A of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a production quality assurance, which is subject to periodic surveillance, defined by Annex XI, Part A, Section 7 of the aforementioned regulation. If class III devices or class IIb devices are covered by this certificate, an EU type-examination certificate in accordance with Annex X of the aforementioned regulation is required before placing them on the market.

Report No.:	244384642-200
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Issue date:	2023-01-06

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-091

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Herbert Zhong
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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