Certificate of EU Medical Device Notification

This is to certify that, according Regulation (EU) 2017/745 of THE EUROPEAN PARLIAMENT AND OF THE COUNCIL on medical devices,

Humiss Beratung GmbH
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SRN: DE-AR-000023447

has fulfilled all notification responsibility and duty as the European Authorized representative of:

Manufacturer: Shanghai Uretech Film Material Co., Ltd
Address: Building 2, 1st Floor, No. 88, Lane 819, Duhui Road, Minhang District, Shanghai, 201108,China

The manufacturer has provided with all the appropriate declaration according the Regulation (EU) 2017/745 requirements including the EC Declaration of Conformity confirming that the medical device, as stipulated here below, is fulfilling the applicable requirements of Regulation (EU) 2017/745.

Product(s):	Dental film
Notification number:	DE/CA20/00198897
Model(s):	Uremono 661, Uremono 662
Classification:	Class I, rule 1

Where then manufacturer affixes the CE marking to the product listed, they must ensure that all the requirements of the appropriate EU regulation(s) have and continue to be met.

The notification of aforementioned device(s) has been completed by European Authorized representative in Germany, the Germany Competent Authority has notified the manufacturer's medical device above and has allocated registration.

CE

Humiss Beratung GmbH

This is only a CE mark sample which is only use for reference.

Signature of Executive Director
James St. WU
Title: General manager

Cert. №: D-YQ-MD-231220140
Issue date: Dec.20,2023
Valid until: Dec.19,2028