CE

EU Declaration of Conformity for medical devices
In accordance to Regulation (EU) 2017/745 of the European Parliament and the Council

Manufacturer	Profümed Karlheinz Lohr e.K. Erzstraße 36 09618 Brand-Erbisdorf
Product	Zellstofftupfer 4 x 5 cm, 8 x 10 cm
Risk Class according to annex VIII	Class I (Rule IV)
Fulfilled technical standards	Manufactured according to product specifications
Conformity assessment procedure	According to annex IV of the regulation
Basis UDI-DI	4251334101000NV

The Manufacturer declares that the above products / group of products are in compliance with the applicable Regulation (EU) 2017/745. The Manufacturer has carried out the conformity assessment procedure and labeled the products with the appropriate CE marking and reported to the authority.

The technical documentation is compiled and is kept by the manufacturer.

26.05.2020 BED

Date and Place of issue of the Declaration of conformity

Georg Scheffer, CEO