

EU declaration of conformity

Manufacturer according to Regulation 2017/745 Schülke & Mayr GmbH Robert-Koch-Str. 2 22851 Norderstedt Germany

Registration Number acc. to Art. 31 2017/745

DE-MF-000005701

Product name

thermosept® RKF

basic UDI-DI Code acc. to Art. 26 2017/745 Intended Purpose

4032651-BSC00000004-CT

Risk Class according to Regulation 2017/745

VIII annex rule 1

V0799

Standards applied

EN ISO 13485

additional standards see technical documentation

cleaning agent for automated reprocessing of medical devices

Schülke & Mayr GmbH

Conformity Assessment Procedure according to Regulation 2017/745

IV / V annex

Certificate

EN ISO 13485

004567 MP2016

Version

2-0

Schülke & Mayr GmbH herewith declares that the device covered by this declaration is in conformity with the Regulation 2017/745 concerning medical devices.

Schülke & Mayr GmbH declares that Schülke & Mayr GmbH bears the sole responsibility for issuing this Declaration

Norderstedt 09. Nov. 2021

Dr. Uwe Berlekamp Schülke & Mayr GmbH Director Business Line Healthcare ppa.

Jörn Ahlsdorff Schülke & Mayr GmbH **Director Industrial Operations** International Industrial Operations