EU Declaration of Conformity

schülke

EC 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	**gigasept® powerSET3 disinfection wipe**
Basic UDI-DI	4032651BSC00000014A5
Code acc. to Art. 26 2017/745	D050103
Intended Purpose	Disinfectant of medical device surfaces at the endpoint of reprocessing
Risk Class according to Regulation 2017/745	II b Annex VIII rule 16
Standards applied	EN ISO 13485 additional standards see technical documentation Schülke & Mayr GmbH
Notified Body	DQS Medizinprodukte GmbH August-Schanz-Str. 21 60433 Frankfurt am Main Germany No.: 0297
Conformity Assessment Procedure according to Regulation 2017/745	Annex IX Chapter I, II section 4 and III
Certificate	Annex IX 004567 MDR2017Q
Version	1-1

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 31.08.2024
ppa.

Dr. Rainer Wolber
Schülke & Mayr GmbH
Director Innovation & Regulatory Affairs

31.08.2024
ppa.

Lars Lemke
Schülke & Mayr GmbH
Chief Operating Officer

This Declaration is valid until an updated version has been issued, but not longer than 03.05.2028