



EU Declaration of Conformity

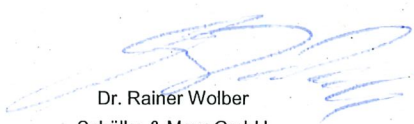
Manufacturer according to Regulation 2017/745	Schülke & Mayr GmbH Robert-Koch-Str. 2 22851 Norderstedt Germany		
Registration Number acc. to Art. 31 2017/745	DE-MF-000005701		
Product Name	gigasept® FF new		
Basic UDI-DI Code acc. to Art. 26 2017/745	4032651BSC00000036AF Z12011385		
Intended Purpose	disinfectant for manual reprocessing of medical devices		
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
Certificates	Annex	IX	004567 MDR2017Q
	EN ISO 13485		004567 MP2016
Version	1-0		

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 05.11.2024
ppa.

05.11.2024
ppa.


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


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