Gründler
medical

EU Declaration of Conformity

regarding EU directive 2011/65/EU

Manufacturer's Name:	Gründler GmbH
Manufacturer's Address:	Jaspisstraße 23, 72250 Freudenstadt, Germany
Authorized Representative Name:	N/A
Authorized Representative Address:	N/A
Basic UDI-DI:	426237137000952BS
Name of the Device:	Ventilution Patient Set Standard
Production Period:	31.03.2026 – 30.03.2029
Product REF:	REF 000070
Purpose:	Reduction of deadspace
Directive:	2011/65/EU

This declaration of conformity is issued under the sole responsibility of Gründler GmbH. The above-mentioned product is in conformity with Directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment.
All supporting documentation is retained at the premises of the manufacturer. The declaration is valid with the date of the signature.

Signature:

.....
Dr. Christoph Gründler/CEO

Place and date of issue:

Freudenstadt, 2026-03-31

.....
Place and date

Gründler GmbH
Geschäftsführer:
Markus Gründler
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Amtsgericht Stuttgart
HRB 761413

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