



Tracoe Medical GmbH | Reichelsheimer Str. 1/3 | 55268 Nieder-Olm

To whom it may concern

**Tracoe Medical GmbH**

Reichelsheimer Str. 1/3  
55268 Nieder-Olm

Tel: +49 6136 9169-0  
Fax: 06136 9169-200

info-tracoe@atosmedical.com  
www.tracoe.com

Geschäftsführer:  
Dr. Hans Kränzler

Amtsgericht Mainz, HRB 44548  
Steuer-Nr.: 08/658/71878

Your Reference

Our Reference

Date

May 08, 2024

## Declaration of Manufacturer Stating the Fulfilment of Extension Conditions

The undersigned, acting as Managing Director of Tracoe Medical GmbH, hereby confirms that Tracoe Medical GmbH, located Reichelsheimer Str. 1/3, 55268 Nieder-Olm, Germany, has obtained an extension of the validity of its conformity assessment certificates issued by the Notified Body TÜV SÜD Product Service GmbH (NB #0123) issued under the Medical Devices Directive 93/42/CEE (MDD), in accordance with the provisions set by the Regulation EU 2023/607 amending Regulations (EU) 2017/745 (MDR).

The medical devices in scope of this extension are those listed in the Notified Body Confirmation Letter #CL 036993 0027 issued by TÜV SÜD and dated 09 April 2024.

Consistent with the provisions set out in Article 120 of the MDR, as amended, this extension entails the following conditions:

- those devices continue to comply with the MDD;
- there are no significant changes in the design and intended purpose;
- the devices do not present an unacceptable risk to the health or safety of patients, users or other persons, or to other aspects of the protection of public health;
- Tracoe Medical GmbH has put in place a quality management system in accordance with applicable provisions of the MDR;
- Tracoe Medical GmbH has lodged a formal application and has signed a mutual agreement with the Notified Body TÜV SÜD for continued surveillance under MDD and conformity assessment under the MDR, as applicable.

In accordance with the provisions set out in Article 120(3) of MDR, the transition timeline that applies to the devices manufactured by Tracoe and that are covered by this extension is 31 December 2028. Declarations of Conformity applicable to these products that were previously issued by Tracoe under the MDD remain valid up to 31 December 2028 are thereby also extended.

Declarations of Conformity issued prior to the publication of the Regulation EU 2023/607 and applicable to these devices may have been previously issued with a specified expiration date. Where such expiration date is present on the Declaration of Conformity, it should also be regarded as extended to 31 December 2028.

Attachment

- Confirmation Letter #CL 036993 0027, 7 pages.

Best regards

  
Dr. Hans-Marcus Kränzler  
Managing Director

Mitglied im

  
AKTIONSBÜNDNIS  
PATIENTENSICHERHEIT



## KONFORMITÄTSERKLÄRUNG DECLARATION OF CONFORMITY

Entsprechend DIN EN ISO / IEC 17050-1 According to DIN EN ISO / IEC 17050-1

Die TRACOE medical GmbH, Reichelsheimer Str. 1/3, in 55268 Nieder-Olm, wendet ein Qualitätsmanagementsystem an, das den Anforderungen der DIN EN ISO 13485:2016 entspricht.

TRACOE medical GmbH erklärt in alleiniger Verantwortung, dass die folgenden Medizinprodukte allen Anforderungen der Richtlinie 93/42/EWG entsprechen, und deren Umsetzung in nationale Gesetze. Das Konformitätsverfahren gemäß Anhang II ohne (4) der Richtlinie wurde durchgeführt in Zusammenarbeit mit der Benannten Stelle:

TRACOE medical GmbH, Reichelsheimer Str. 1/3, in 55268 Nieder-Olm, Germany is maintaining a quality management system, which meets the requirements of DIN EN ISO 13485:2016.

TRACOE medical GmbH declares in its own responsibility that the following medical devices meet all requirements of the Directive 93/42/EEC and the implementation of the Directive according to national laws. The conformity assessment procedure according to Annex II excluding (4) of the Directive has been carried out in cooperation with our Notified Body:

**TÜV SÜD Product Service GmbH (0123)**  
**Ridlerstraße 65**  
**80339 München**

### TRACOE *vario*

#### Tracheostomiekannülen Tracheostomy Tubes

REF 450, REF 455, REF 460, REF 465 (-06,-07,-08,-09,-10,-11) seit 05/2003	REF 450, REF 455, REF 460, REF 465 (-06,-07,-08,-09,-10,-11) since 05/2003
---	--

REF 470 (-07,-08,-09,-10) seit 07/2004	REF 470 (-07,-08,-09,-10) since 07/2004
---	--

REF 451 (-07,-08,-09,-10) seit 04/2006	REF 451 (-07,-08,-09,-10) since 04/2006
---	--

REF 461, REF 471 (-07,-08,-09,-10) seit 12/2011	461, REF 471 (-07,-08,-09,-10) since 12/2011
---	--

REF 464 (-07,-08,-09,-10) seit 03/2016	REF 464 (-07,-08,-09,-10) since 03/2016
--	---

<b>Klassifizierung nach Richtlinie 93/42/EWG, Anhang IX als Medizinprodukt Klasse IIb</b>	<b>Classification according to Directive 93/42/EEC, Annex IX, as Medical Devices, class IIb</b>
---	---

**EG-Zertifikat Nr. G1 036993 0016 Rev. 01 EC-Certificate No. G1 036993 0016 Rev. 01****Gültig bis 2024-05-26 Valid until 2024-05-26**11.03.2020

Nieder-Olm

T. Jurisch**Dr. Thomas Jurisch**  
General Manager



Add value.  
Inspire trust.

TÜV SÜD Product Service GmbH · Ridlerstr. 65 · 80339 Munich · Germany

TRACOE medical GmbH  
Reichelsheimer Str. 1/3  
55268 Nieder-Olm

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
	713181269	Bettina.Hoeffeler@tuvsud.co		2024-04-09	1 of 7

**TÜV SÜD Product Service GmbH**  
**Confirmation Letter**  
**CL 036993 0027 Rev. 00**

**Reference: 713181269**

To whom it may concern,

**Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 (in the following referenced as MDR) as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.**

With this letter TÜV SÜD Product Service GmbH, designated under MDR and identified by the number 0123 on NANDO, confirms that we have received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the above stated manufacturer with the following SRN Number:

SRN Number: DE-MF-000006938

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below.

- Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive.

- If devices covered by certificates issued under Directive 90/385/EEC (AIMDD) or Directive

**Registered Office: Munich**  
Trade Register Munich HRB 85 742  
UniCredit Bank AG · BIC HYVEDEMMXXX  
IBAN DE13 7002 0270 0048 8522 11  
VAT ID No. DE129484267  
Information pursuant to § 2 [1] DL-InfoV  
(Germany) at tuvsud.com/imprint

**Supervisory Board:**  
Holger Lindner (Chairman)  
**Board of Management:**  
Walter Reithmaier (CEO)  
Patrick van Welij

**TÜV SÜD Product Service GmbH**  
Ridlerstr. 65  
80339 Munich  
Germany

tuvsud.com/ps  
Hotline: +49 89 50084-747

**TÜV®**



93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that

- the manufacturer signed the written agreement under MDR by the date of MDD/AIMDD certificate expiry; or
- provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively.

The transition timelines in accordance Article 120 (3) of MDR that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120 (3c) of MDR, are shown below:

- 26 May 2026 for Class III custom-made implantable devices
- 31 December 2027 for Class III devices and Class IIb implantable devices (except sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition, measuring function
- 31 December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

We reserve the right to invoice any issuance, copies, amendments and / or changes of the confirmation letter according to effort.

For confirmation letter validity see [www.tuvsud.com/ps-cert?q=cert:CL\\_036993\\_0027\\_Rev.00](http://www.tuvsud.com/ps-cert?q=cert:CL_036993_0027_Rev.00)

In case of inquiries please contact [medical\\_devices@tuvsud.com](mailto:medical_devices@tuvsud.com).

On behalf of the Notified Body TÜV SÜD Product Service GmbH,  
2024-04-09

TÜV SÜD Product Service GmbH  
Medical and Health Services

TÜV SÜD Product Service GmbH  
Medical and Health Services

Handwritten signature of Bettina Hoeffler in black ink.

Bettina Hoeffler  
Conformity Assessment Responsible (CARE)

Handwritten signature of Claus Matthias Mumme in black ink.

Claus Matthias Mumme  
Application Reviewer





**Table 1: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:**

Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
<b>Device 1</b> 4035324TWIST_PLUSSR  <b>TRACOE twist plus Tracheostomy Tube</b>  <b>TRACOE twist plus extract Tracheostomy Tube</b>  <b>TRACOE twist plus Spare Inner Cannula</b>  <b>TRACOE experec Set twist plus</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: Certificate G1 036993 0016 Rev. 01 NB 0123
<b>Device 2</b> 4035324TWISTX8  <b>TRACOE twist Tracheostomy Tube</b>  <b>TRACOE twist Laryngectomy Tube</b>  <b>TRACOE twist extract Tracheostomy Tube</b>  <b>TRACOE twist Spare Inner Cannula</b>  <b>TRACOE twist short Spare Inner Cannula</b>  <b>TRACOE experec Set twist</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 3</b> 4035324VARIOT9  <b>TRACOE vario Tracheostomy Tube</b>  <b>TRACOE vario XL Tracheostomy Tube</b>  <b>TRACOE vario extract Tracheostomy Tube</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
TRACOE vario extract XL Tracheostomy Tube  TRACOE experec Set vario  TRACOE experec Set vario XL			
<b>Device 4</b> 4035324SILCOSOFTEL  TRACOE silcosoft Tracheostomy Tube  TRACOE silcosoft PL Tracheostomy Tube	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 5</b> 4035324MINIW4  TRACOE mini Extension Piece  TRACOE mini Tracheostomy Tube	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 6</b> 4035324COMFORTNT  TRACOE comfort Tracheostomy Tube  TRACOE comfort XL Tracheostomy Tube  TRACOE comfort plus Tracheostomy Tube  TRACOE care Paraffin oil	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input checked="" type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device <input type="checkbox"/> Class I reusable surgical instruments	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 7</b> 4035324AERISM6	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input type="checkbox"/> Class IIb / Class IIb implantable (exempted)	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123





Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
<b>TRACOE aeris Ballon-Dilatationskatheter</b>	<input checked="" type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device <input type="checkbox"/> Class I reusable surgical instruments		
<b>Device 8</b> <b>4035324EXPERC_DIL_SET46</b>  <b>TRACOE experc Dilation Set</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input checked="" type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device <input type="checkbox"/> Class I reusable surgical instruments	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 9</b> <b>4035324COMPR_STERILE3R</b>  <b>TRACOE purofoam Tracheal Compress</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input checked="" type="checkbox"/> Class IIa <input type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device <input type="checkbox"/> Class I reusable surgical instruments	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1 036993 0016 Rev. 01 NB 0123
<b>Device 10</b> <b>4035324SPEAK_VALVESRC</b>  <b>Occlusion Cap</b>  <b>Stopper</b>  <b>Speaking Valves Tracoe Phone assist I</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input checked="" type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1S 036993 0014 Rev.01 NB 0123



Device name or Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified during application review)	If the MDR device is a substitute device, identification of the corresponding MDD/AIMDD device	MDD/AIMDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
	<input type="checkbox"/> Class I reusable surgical instruments		
<b>Device 11</b> <b>4035324SMARTTC</b>  <b>TRACOE smart Cuffmanager</b>	<input type="checkbox"/> Class III <input type="checkbox"/> Class IIb implantable (non-exempted) <input type="checkbox"/> Class IIb / Class IIb implantable (exempted) <input type="checkbox"/> Class IIa <input checked="" type="checkbox"/> Class I devices in sterile condition <input type="checkbox"/> Class I devices with measuring function <input type="checkbox"/> Class III implantable custom-made-device <input type="checkbox"/> Class I reusable surgical instruments	<input checked="" type="checkbox"/> N/A	<input checked="" type="checkbox"/> Certification as follows: G1S 036993 0014 Rev.01 NB 0123



### Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024-04-09	713181269	Initial issue