



**EC-declaration of conformity for In-vitro Diagnostic Device**  
according to Regulation (EU) 2017/746 IVDR (Article 48)

*We, the manufacturer*

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Ritter GmbH (part of Avantor)  
Kaufbeurer Straße 55, 86830 Schwabmünchen  
**SRN:** DE-MF-000036772

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*hereby declare under our sole responsibility that the accessory for an in-vitro diagnostic medical device (EU 2017/746 Article 2 Paragraph 4) with the*

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<b>product name:</b>	Robotic Tips
<b>brands:</b>	- J.T.Baker - Ritter
<b>types:</b>	- Conductive, non-sterile - Non-conductive, non-sterile
<b>Basic- UDI-DI:</b>	4033789RoTip-Co/NCo-NStNR
<b>Risk Class:</b>	A (according to IVDR Annex VIII, rule 5a products for general laboratory use, accessories)

**and the intended purpose:**

The robotic tips from Ritter GmbH are used exclusively for the low-contamination dispensing and dosing of liquids in the context of analytical, diagnostic, biotechnological, or microbiological tests. They are intended for single use only and may be used only by qualified and trained specialised/laboratory personnel. The robotic tips are considered accessories for automated liquid handling platforms, which are themselves marketed as in vitro diagnostics (IVD) or general laboratory devices. Together with the necessary reagents or assay kits, these enable the intended use.

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*complies with all the requirements of Regulation (EU) 2017/746 (IVDR) Annex II and III, is marked with:*



Schwabmünchen, 2025-02-20 (yyyy-mm-dd)

Michael Kazinski- Höfer  
Managing Director

  
Judith Kröfl  
PRRC