



## 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*

**product name**

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**Robotic Tips (non-sterile)**

**J.T.Baker & ritter**


Types: **Conductive and non-conductive** Basic-UDI-DI: **4033789RoTip-Co/NCo-NStNR**  
Risk Class: **A (according to IVDR Annex VIII, rule 5a products for general laboratory use, accessories)**

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



Schwabmünchen, 2023-12-14 (yyyy-mm-dd)

  
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Thorsten Kopp  
Managing Director

  
\_\_\_\_\_  
Judith Kroll  
RRRC

This declaration applies to non-sterile IVD robotic tips (conductive and non-conductive) marked with CE and from production date 25<sup>th</sup> of May 2022.