EC-declaration of conformity
for In-vitro Diagnostic Device

according to Regulation (EU) 2017/746 IVDR (Article 48)

We, the ***manufacturer***

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

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

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*)

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

CE

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

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 25th of May 2022.

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Registergericht: Augsburg | Handelsregister: HRB 6735 | UST-ID Nr.: DE 127495980