



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

*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)

Types: **Conductive and non-conductive**

Risk Class: A (according to IVDR Annex VIII, rule 5a products for general laboratory use,
accessories)

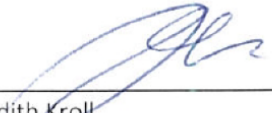
*complies with all the requirements of Regulation (EU) 2017/746 (IVDR) Annex II and III and is
marked with:*



Schwabmünchen, 2022-05-25 (yyyy-mm-dd)



Johannes v. Stauffenberg
Vice President & Managing Director



Judith Kroll
PRRC