ahn
biotechnologie GmbH

Declaration of Conformity Certificate

We

AHN Biotechnologie GmbH
Uthleber Weg 14
99734 Nordhausen
Germany
Tel. +49 (0) 3631 65242-0
www.cappahn.com

Declare with sole responsibility, that our product/s:

EDMA Code	EDMA Description	Internal Product Name	Classification Rationale per IVDD
21-09	Pipette tips	Expell, ExpellPlus, myTip Pipette Tips, Sterile Tips , Non-Sterile Tips, Low Retention Tips and Filter Tips	Other IVD, Annexe III

meet, the essential requirements of Council Directive 98/79/EC pertaining to in vitro diagnostics. Pathway of conformity per Annex III.

Notified Body: -

The product(s) identified above meet requirements of the IVDD by meeting the following standards

Standard No.
Council Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC

We hereby appoint mdi Europa GmbH, Langenhagener Str. 71, 30855 Langenhagen, Germany to act as European Authorized Representative as explicitly defined in Article 1, § 2(g) of Directive 98/79/EEC.

Signed this day: 05-03-2018, by Magdalena Babut-Carstensen, Compliance Manager

AHN Biotechnologie GmbH
Uthleber Weg 14
99734 Nordhausen
Tel. + 49 3631 46594-04
FAX + 49 3631 46594-10

Expiry Date: -

mdi Europa use only!

The necessary pre-requisites for placing the CE mark on the above mentioned products and marketing them in all Member States of the European Union, have thus been fulfilled.

Signed this day: 05-03-2018

mdi Europa GmbH
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12 Sep 2018

mdiEuropa
THE MEDICAL DEVICE SERVICE-MANAGEMENT

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Digitally signed by Lazari Cristina
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