	<b>ACCUMAX LAB DEVICES PRIVATE LIMITED</b>	MODEL	Sterile Tips , Non-Sterile Tips, Low Retention Tips and Filter Tips
	<b>TECHNICAL CONSTRUCTION FILE</b>	Issue No.	01
		Rev. No.	02
<b>Product Description : Pipette Tips</b>		Date	15.11.2017

# EC Declaration of Conformity

**We** Company : Accumax Lab Devices Private Limited

Address. : Plot No.16, Electronic Park  
(Sez.) , Kolavada Road,  
Gandhinagar-380026  
Gujarat,India.

Country : India.

**Declare with sole responsibility, that our product/s:**

**EDMS-Code**  
21-09

**EDMA-Description**  
Pipette Tips and other Accessories

**Classification**  
Other IVD, Annexe III

meet, (where applicable), the essential requirements of Council Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC pertaining to medical devices.

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



Signed this day.15<sup>th</sup>of November 2017


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 .15<sup>th</sup> of November 2017

22 Nov 2017  
mdi Europa GmbH  
Langenhagener Str. 71 · 30855 Langenhagen  
Phone +49-511. 39 08 95 30  
Fax +49-511. 39 08 95 39

  
**mdi Europa**  
THE MEDICAL DEVICE SERVICE-MANAGEMENT

	<b>ACCUMAX LAB DEVICES PRIVATE LIMITED</b>	MODEL	Mechanical (Regular, PRO, Auto clavable , Smart and Junior Pipette) and Electronic (Digital Micro Pipette ) and Pipette Controller
	<b>TECHNICAL CONSTRUCTION FILE</b>	Issue No.	01
		Rev. No.	02
<b>Product Description : Mechanical and Electronic Micropipette and Pipette Controller</b>		Date	15.11.2017

# EC Declaration of Conformity

**We** Company : Accumax Lab Devices Private Limited

Address. : Plot No.16, Electronic Park  
(Sez.) , Kolavada Road,  
Gandhinagar-380026  
Gujarat,India.

Country : India.

**Declare with sole responsibility, that our product/s:**

**EDMS-Code**  
21-09

**EDMA-Description**  
Electronics and Mechanical Micropipette and  
Pipette Controller (Liquid Handling Equipments),

**Classification**  
Other IVD, Annexe III

meet, (where applicable), the essential requirements of Council Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EEC pertaining to medical devices.

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



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mdi Europa  
THE MEDICAL DEVICE SERVICE-MANAGEMENT