



## DECLARATION OF CONFORMITY

|                              |  |
|------------------------------|--|
| MANUFACTURER:                | DIESSE DIAGNOSTICA SENESE SPA<br>VIA DELLE ROSE 10<br>53035 MONTERIGGIONI (SI),<br>ITALY |
| EUROPEAN REPRESENTATIVE:     | //   |
| GENERIC NAME                 | KIT FOR INSTALLATION OF INTEGRATED AUTOMATED<br>PIPETTOR INSTRUMENT                      |
| PRODUCT:<br>CODE:            | <b>AUTOSAMPLER UPGRADE KIT<br/>81215</b>   |
| TECHNICAL DATA:              | 90-264 V (50-60 Hz); Pwr: 100 VA   |
| CLASSIFICATION:              | IVD NOT IN ANNEX II OR SELF-TESTING IVD  |
| CONFORMITY ASSESSMENT ROUTE: | ANNEX APPLIED N° III EXCLUDING (6)<br>ESSENTIAL REQUIREMENTS OF ANNEX I                  |

WE HEREWITH DECLARE THAT THE DESIGN, TYPE OF MANUFACTURE OF THE IN VITRO MEDICAL DIAGNOSTIC DEVICE DESCRIBED ABOVE AND THE VERSION DISTRIBUTED ON THE MARKET, CONFORMS TO THE 98/79/EEC DIRECTIVE RELEVANT TO THE IN VITRO MEDICAL-DIAGNOSTICS DEVICES (IVD)

### THIS CERTIFICATE WILL LOSE ITS VALIDITY IN THE EVENT OF:

- MODIFICATIONS MADE TO THE MACHINE IN QUESTION WITHOUT OUR AUTHORIZATION
- INCORRECT USE OF THE INSTRUMENT
- TECHNICAL INTERVENTIONS PERFORMED BY UNAUTHORIZED PERSONNEL
- INSTALLATION OF NON-ORIGINAL SPARE PARTS.

### THE PRODUCT CONFORMS, AS A WHOLE AND IN ITS PARTS, WITH THE FOLLOWING STANDARDS AND THEIR AMENDMENTS:

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|----------------------------|---|
| <b>EN 61010-1:2010</b>     | SAFETY REQUIREMENTS FOR ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL, AND LABORATORY USE - PART 1: GENERAL REQUIREMENTS  |
| <b>EN 61010-2-081:2015</b> | SAFETY REQUIREMENTS FOR ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL, AND LABORATORY USE - PART 2-081: PARTICULAR REQUIREMENTS FOR AUTOMATIC AND SEMI-AUTOMATIC LABORATORY EQUIPMENT FOR ANALYSIS AND OTHER PURPOSES |
| <b>EN 61010-2-101:2002</b> | SAFETY REQUIREMENTS FOR ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL, AND LABORATORY USE - PART 2-101: SAFETY REQUIREMENTS FOR IN VITRO DIAGNOSTIC (IVD) MEDICAL EQUIPMENT   |

**EN 61326-1:2013**

ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL AND LABORATORY USE - EMC REQUIREMENTS - PART 1: GENERAL REQUIREMENTS

**EN 61326-2-6:2006**

ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL AND LABORATORY USE - EMC REQUIREMENTS - PART 2-6: PARTICULAR REQUIREMENTS - IN VITRO DIAGNOSTIC (IVD) MEDICAL EQUIPMENT

**AND THEREFORE MEETS THE ESSENTIAL REQUIREMENTS OF THE FOLLOWING COMMUNITY DIRECTIVES AND THEIR AMENDMENTS:**

**LOW VOLTAGE DIRECTIVE (2014/35/EU)**

**ELECTROMAGNETIC COMPATIBILITY DIRECTIVE (2014/30/EU)**

**RESTRICTION OF HAZARDOUS SUBSTANCES IN ELECTRICAL AND ELECTRONIC EQUIPMENT DIRECTIVE – ROHS2 (2011/65/EU)**

NOTIFIED BODY:

NOT NECESSARY

(EC) CERTIFICATE:

N.A.

START OF CE-MARKING:

DECEMBER 2017

REVISION:

2

PLACE, DATE OF ISSUE:

MONTERIGGIONI, 26 NOVEMBER 2020

EXPIRY DATE:

25 MAY 2022

**THE PRESENT DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE AND EXCLUSIVE RESPONSIBILITY OF THE MANUFACTURER.**

SIGNATURE:



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CHIARA MUZZI  
QUALITY & REGULATORY MANAGER

THIS IS THE CERTIFIED COPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA

ISSUED: MONTERIGGIONI,

26/11/2020



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MAGDALENA STOCZKO  
REGULATORY SUPERVISOR