DIESSE	DECLARATION OF CONFORMITY
MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA VIA DELLE ROSE 10 53035 MONTERIGGIONI (SI), ITALY
EUROPEAN REPRESENTATIVE:	//
GENERIC NAME	AUTOMATIC INSTRUMENT FOR ESR ANALYSIS
PRODUCT: CODE:	VES-MATIC EASY 10376
TECHNICAL DATA:	9V DC; 2 A
CLASSIFICATION:	IVD NOT IN ANNEX II OR SELF-TESTING IVD
CONFORMITY ASSESSMENT ROUTE:	ANNEX APPLIED N° III EXCLUDING (6) 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:

EN 61010-1:2010	SAFETY	REQUIREMENTS	FOR	ELECTRICAL	EQUIPMENT	FOR
	MEASURE	MENT, CONTROL,	AND LA	BORATORY USE	E - PART 1: GEN	IERAL
	REQUIRE	MENTS				

- EN 61010-2-081:2015SAFETYREQUIREMENTSFORELECTRICALEQUIPMENTFORMEASUREMENT, CONTROL, ANDLABORATORYUSE-PART2-081:PARTICULARREQUIREMENTSFORAUTOMATICANDSEMI-AUTOMATICLABORATORYEQUIPMENTFORANALYSISANDOTHERPURPOSES
- **EN 61010-2-101:2002** 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	G: JANUARY 2007				
REVISION:	5				
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.					
	Ch hi				
SIGNATURE:	CHIARA MUZZI QUALITY & REGULATORY MANAGER				
THIS IS THE CERTIFIED C	OPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA				
ISSUED: MONTERIGGIONI,	26/11/2020 MACDALENA STOCZKO				

MAGDALENA STOCZKO REGULATORY SUPERVISOR