	DECLARATION OF CONFORMITY
MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA STRADA DEI LAGHI 39 53035 MONTERIGGIONI (SI), ITALY
EUROPEAN REPRESENTATIVE:	//
GENERIC NAME	AUTOMATIC INSTRUMENT FOR ESR ANALYSIS
PRODUCT: CODE:	VES-MATIC CUBE 200 FOR BECKMAN COULTER 10370/BC
TECHNICAL DATA:	90-264 Vac (50-60 Hz); Pwr: 265 VA
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 61010-1:2001 SAFETY REQUIREMENTS FOR ELECTRICAL EQUIPMENT FOR MEASUREMENT, CONTROL, AND LABORATORY USE - PART 1: GENERAL REQUIREMENTS
- EN 61010-2-081:2002 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
- 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 61326-1:2006		EQUIPMENT FOR MEASUREMENT, CONTROL AND USE - EMC REQUIREMENTS - PART 1: GENERAL IS
EN 61326-2-6:2006	LABORATORY	EQUIPMENT FOR MEASUREMENT, CONTROL AND USE - EMC REQUIREMENTS - PART 2-6: PARTICULAR IS - 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	:	NOVEMBER 2005
REVISION:		8
PLACE, DATE OF ISSUE		MONTERIGGIONI, 25 MAY 2022
EXPIRY DATE:		25 MAY 2027
THE PRESENT EC DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE AND EXCLUSIVE RESPONSIBILITY OF THE MANUFACTURER.		
		Chhi
SIGNATURE:		
		REGULATORY AFFAIRS MANAGER
THIS IS THE CERTIFIED COPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA		
ISSUED: MONTERIGGIONI,	25/05/2022	M. Stocks
		MAGDALENA STOCZKO REGULATORY SUPERVISOR