	DECLARATION OF CONFORMITY
MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA STRADA DEI LAGHI 39 53035 MONTERIGGIONI (SIENA), ITALY
EUROPEAN REPRESENTATIVE:	//
GENERIC NAME	AUTOMATIC INSTRUMENT FOR ESR ANALYSIS
PRODUCT: CODE:	CUBE 30 touch 10395
TECHNICAL DATA:	110-230 Vac (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) 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 MEASUREMENT, CONTROL, AND LABORATORY USE - PART 1: GENERAL REQUIREMENTS
- EN 61010-2-081:2015 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 61326-1:2013		EQUIPMENT FOR MEASUREMENT, CONTROL AND USE - EMC REQUIREMENTS - PART 1: GENERAL TS
EN 61326-2-6:2006	LABORATORY	EQUIPMENT FOR MEASUREMENT, CONTROL AND USE - EMC REQUIREMENTS - PART 2-6: PARTICULAR TS - IN VITRO DIAGNOSTIC (IVD) MEDICAL EQUIPMENT
AND THEREFORE MEETS THE ESSENTIAL REQUIREMENTS OF THE FOLLOWING COMMUNITY DIRECTIVES AND THEIR AMENDMENTS:		
LOW VOLTAGE DIRECT	IVE (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	:	FEBRUARY 2018
REVISION:		4
PLACE, DATE OF ISSUE:		MONTERIGGIONI, 9 SEPTEMBER 2021
EXPIRY DATE:		25 MAY 2022
THE PRESENT EC DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE AND EXCLUSIVE RESPONSIBILITY OF THE MANUFACTURER.		
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SIGNATURE:		CHIARA MUZZI REGULATORY AFFAIRS MANAGER
THIS IS THE CERTIFIED COPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA		
ISSUED: MONTERIGGIONI,	09/09/2021	M. Stocks
		MAGDALENA STOCZKO REGULATORY SUPERVISOR