	DECLARATION OF CONFORMITY
MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA STRADA DEI LAGHI 39
	53035 MONTERIGGIONI (SI) ITALY
EUROPEAN REPRESENTATIVE:	//
PRODUCT: CODE:	ENZY-WELL RUBELLA IgG 91030, 91176
CLASSIFICATION:	IVD IN ANNEX II B
CONFORMITY ASSESSMENT ROUTE:	ANNEX APPLIED N° IV EXCLUDING (4, 6)
WE HEREWITH DECLARE THAT THE ABOVE MENTIONED PRODUCTS MEET THE PREVISIONS OF THE COUNCIL DIRECTIVE 98/79EC FOR IN VITRO DIAGNOSTIC MEDICAL DEVICE. ALL SUPPORTING DOCUMENTATION IS RETAINED UNDER THE PREMISES OF THE MANUFACTURER.	
NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH
	ZERTIFIZIERSTELLE RIDLERST. 65 – 80339 MÜNCHEN
	GERMANY
	No. 0123
(EC) CERTIFICATE:	VI 056726 0002 Rev. 00
START OF CE-MARKING:	SEPTEMBER 2003
REVISION:	14
PLACE, DATE OF ISSUE:	MONTERIGGIONI, 11 MAY 2022
EXPIRY DATE:	26 MAY 2025
THE PRESENT EC DECLARATION OF CONFORMITY IS ISSUED UNDER THE SOLE AND EXCLUSIVE RESPONSIBILITY OF THE MANUFACTURER.	
	Chli
SIGNATURE:	CHIARA MUZZI REGULATORY AFFAIRS MANAGER
	REGULATORT AFFAIRS MANAGER
THIS IS THE CERTIFIED COPY OF THE ORIGINAL DOCUMENT STORED IN ARCHIVE OF DIESSE DIAGNOSTICA SENESE SPA	
	M. Stocks
ISSUED: MONTERIGGIONI, 11/05/2022	MAGDALENA STOCZKO REGULATORY SUPERVISOR