<b>DIESSE</b> EC DECLARATION OF CONFORMITY	
MANUFACTURER:	DIESSE DIAGNOSTICA SENESE SPA
	STRADA DEI LAGHI 39
	53035 MONTERIGGIONI (SI)
	ITALY
EUROPEAN REPRESENTATIVE:	//
PRODUCT:	CHORUS RUBELLA IgG AVIDITY
CODE:	81095
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:	V1 056726 0002 Rev. 00
START OF CE-MARKING:	FEBRUARY 2004
REVISION:	14

EXPIRY DATE:

PLACE, DATE OF ISSUE:

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

26 MAY 2025

MONTERIGGIONI, 11 MAY 2022

SIGNATURE:

CHIARA MUZZI REGULATORY AFFAIRS MANAGER

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

M. Stoules

ISSUED: MONTERIGGIONI, 11/05/2022

MAGDALENA STOCZKO REGULATORY SUPERVISOR