



## EC DECLARATION OF CONFORMITY

MANUFACTURER:

DIESE DIAGNOSTICA SENESE SPA  
STRADA DEI LAGHI 39  
53035 MONTERIGGIONI (SI)  
ITALY

EUROPEAN REPRESENTATIVE:

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PRODUCT:

**CHORUS RUBELLA IgG AVIDITY CONTROL  
SERUM**

CODE:

**81524**

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

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

SIGNATURE:

  
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CHIARA MUZZI  
REGULATORY AFFAIRS MANAGER

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

ISSUED: MONTERIGGIONI, 11/05/2022

  
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MAGDALENA STOCZKO  
REGULATORY SUPERVISOR