

| MANUFACTURER: | DIESSE DIAGNOSTICA SENESE SPA STRADA DEI LAGHI 39 53035 MONTERIGGIONI (SI), ITALY |
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| EUROPEAN REPRESENTATIVE: | // |
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| CLASSIFICATION: | IVD NOT IN ANNEX II OR SELF-TESTING IVD |
| CONFORMITY ASSESSMENT ROUTE: | ANNEX APPLIED N° III |
| 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. | |
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| 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 |

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ISSUED: MONTERIGGIONI, 24/05/2022

> MAGDALENA STOCZKO REGULATORY SUPERVISOR

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