

**DIESSE DIAGNOSTICA SENESE SpA**  
53035 Monteriggioni (SI) - Italia - Via delle Rose 10  
Tel +39-0577-587111 Fax +39-0577-318690



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Sede Legale e Amministrazione  
20144 Milano (Italy) Via A. Solari 19 sc. 6  
Tel, +39-02-4859121 r.a, Fax +39-02-48008530

### QUALITY CONTROL CERTIFICATE

**KIT NAME: dsDNA-G - CHORUS**

|                                                                |
|----------------------------------------------------------------|
| <b>Code</b> 86032                                              |
| <b>Lot No.</b> 834               2023-05               2022-05 |

### REAGENTS

| Label | Description           | Lot. | Exp     | No  |
|-------|-----------------------|------|---------|-----|
| D3281 | dsDNA-G               | 834  | 2023-05 | 6X6 |
| B3281 | CALIBRATORE dsDNA-G   | 834  | 2023-05 | 1   |
| C3281 | CONTROL SERUM dsDNA-G | 834  | 2023-05 | 1   |

### CONTROLS

| Found results | Expected results |
|---------------|------------------|
|---------------|------------------|

|                       |      |           |
|-----------------------|------|-----------|
| Calibrator (mOD)      | 470  | ≥ 100     |
| Control Serum (IU/mL) | 74,5 | 43,8-92,3 |

### CALIBRATOR

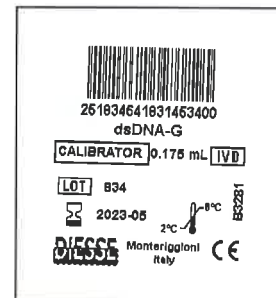
|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 2 | 5 | 1 | 8 | 3 | 4 | 6 | 4 | 1 | 8 | 3 | 1 | 4 | 5 | 3 | 4 | 0 | 0 |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|

Label checked with barcode reader

YES



Operator Signature *Sf*



### CONTROL SERUM

|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 2 | 5 | 1 | 8 | 3 | 4 | 6 | 1 | 6 | 2 | 9 | 0 | 0 | 0 | 0 | 0 | 0 | 1 |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|

Label checked with barcode reader

YES



Operator Signature *Sf*



The test has been carried out in accordance with the package insert

TESTED BY: Simone Baiocchi

*Simone Baiocchi*

APPROVED

RESPONSIBLE HEAD, Quality Control Reagents:

*[Signature]*

DATE

*25.05.2022*

### WARNING: POTENTIAL BIOHAZARDOUS MATERIAL

This kit may contain some reagents made with human serum or plasma. All serum or plasma used has been tested by an FDA approved method and found non-reactive for HIV-1/2, HCV and HBsAg. Because no method can offer complete assurance that HIV-1/2, HCV, HBsAg or other infectious agents are absent, reagents should be handled with maximum attention.