

**DIESSE DIAGNOSTICA SENESE SpA**  
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Sede Legale e Amministrazione  
20144 Milano (Italy) Via A. Solari 19 sc. 6

### QUALITY CONTROL CERTIFICATE

**KIT NAME: CARDIOLIPIN-G - CHORUS**

|                |         |
|----------------|---------|
| <b>Code</b>    | 86046   |
| <b>Lot No.</b> | 845-A   |
|                | 2023-06 |

### REAGENTS

| Label | Description                 | Lot. | Exp     | No  |
|-------|-----------------------------|------|---------|-----|
| D3311 | CARDIOLIPIN-G               | 845  | 2023-06 | 6X6 |
| B3311 | CALIBRATORE CARDIOLIPIN-G   | 845  | 2023-06 | 1   |
| C3311 | CONTROL SERUM CARDIOLIPIN-G | 845  | 2023-06 | 1   |

### CONTROLS

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

|                         |      |           |
|-------------------------|------|-----------|
| Calibrator (mOD)        | 360  | ≥ 100     |
| Control Serum (GPLU/mL) | 29,1 | 21,4-40,6 |

### CALIBRATOR

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

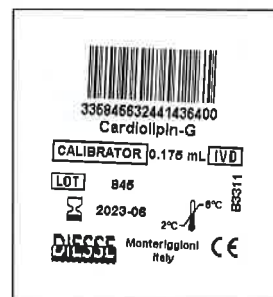
Label checked with barcode reader

YES



Operator Signature

*Simone Baiocchi*



### CONTROL SERUM

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

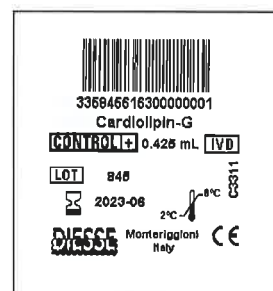
Label checked with barcode reader

YES



Operator Signature

*Simone Baiocchi*



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

TESTED BY: Simone Baiocchi

*Simone Baiocchi*

APPROVED

RESPONSIBLE HEAD, Quality Control Reagents

DATE

*26.07.2022*

### 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.