

**DIESSE DIAGNOSTICA SENESE SpA**  
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### QUALITY CONTROL CERTIFICATE

**KIT NAME: 88000-W2 WESTERN RAGWEED**

|         |          |         |         |
|---------|----------|---------|---------|
| Code    | 88000-W2 |         |         |
| Lot No. | 890      | 2023-11 | 2023-01 |

### REAGENTS

| Label | Description                            | Lot | Exp     | No  |
|-------|----------------------------------------|-----|---------|-----|
| D3151 | 88000-W2 WESTERN RAGWEED               | 890 | 2023-11 | 2X6 |
| B3150 | CALIBRATORE 88000-W2 WESTERN RAGWEED   | 890 | 2024-02 | 1   |
| C3150 | CONTROL SERUM 88000-W2 WESTERN RAGWEED | 890 | 2024-02 | 1   |

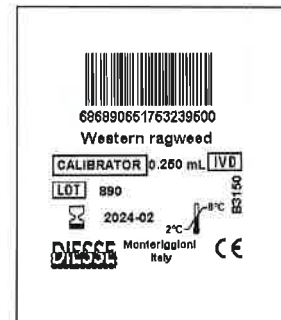
### CONTROLS

|                      | Found results | Expected result |
|----------------------|---------------|-----------------|
| Calibrator (mOD)     | 232           | ≥ 100           |
| Control Serum (KU/L) | 13,0          | 8,8-20,3        |

### CALIBRATOR

|   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |   |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| 6 | 8 | 6 | 8 | 9 | 0 | 6 | 5 | 1 | 7 | 5 | 3 | 2 | 3 | 9 | 5 | 0 | 0 |
|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|

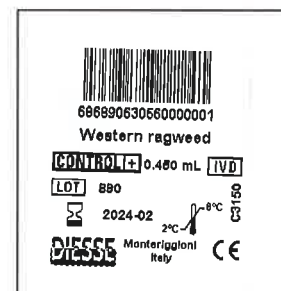
Label checked with barcode reader YES ☒ Operator Signature *SB*



### CONTROL SERUM

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

Label checked with barcode reader YES ☒ Operator Signature *SB*



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 *18.01.2023*

### 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.