

# Test Instructions

## Enzyme Immunoassay for the Detection of Anti-Jo-1 Antibodies

Catalogue-No.: TC 60025

Please read the instructions carefully before testing.

Procedural precautions:

Do not use the reagents beyond the date of expiry.

Reagents from different test kit lots must not be mixed.

### 1. Clinical Use

Anti-Jo-1 antibodies are a special case within the group of antinuclear antibodies (ANA) because their corresponding antigen is located only in the cytoplasm. The Jo-1-autoantigen has been identified as the enzyme histidyl-tRNA synthetase.

Antibodies directed against the Jo-1 antigen are especially found in patients with idiopathic inflammatory myopathia. Anti-Jo-1 antibodies can be detected in 33% of patients with primary polymyositis and in 25% of cases with primary dermatomyositis.

It is remarkable that more than 70% of patients who give positive results in the anti-Jo-1 test are suffering from fibrosing alveolitis and in some cases from polyarthritis. That is why antibodies directed against the Jo-1 antigen are considered as marker antibodies of a subset of myositis with lung disease.

### 2. Principle of the Test

The test is based on the immobilisation of recombinant produced Jo-1 antigen to a solid phase (polystyrene) and subsequent binding of the anti-Jo-1-antibodies. For the detection of antibodies bound in this way a peroxidase-labelled antibody is used which is directed against human IgG. After addition of a substrate solution, a color stain develops, the intensity of which is proportional to the concentration and/or the avidity of the antibodies. For anti-Jo-1 antibody quantification the standard sample has been calibrated with the CDC-serum num. 10.

### 3. Material Provided

- coated microtiter strips (1 x 8), breakable	12 strips
- standards; ready to use,	1 vial each à 750 µl
1: 12.5 U/ml	
2: 25 U/ml	
3: 50 U/ml	
4: 100 U/ml	
5: 200 U/ml	
(contain sodium azide)	
- negative control serum, ready to use, (contains sodium azide)	1 vial 1 ml
- positive control serum, ready to use, (contains sodium azide)	1 vial 1 ml
- washing buffer concentrate (10x), (contain thimerosal)	1 bottle 50 ml
- sample buffer concentrate (5x), (contain sodium azide)	1 bottle 22 ml
- conjugate buffer ready to use,	1 bottle 20 ml
- peroxidase conjugate, anti-human IgG, concentrate (100x),	1 vial 200 µl
- peroxidase substrate solution, (TMB), ready to use,	1 bottle 12 ml
- stopping solution, H <sub>2</sub> SO <sub>4</sub> , ready to use,	1 bottle 12 ml

### 4. Preparation of Reagents

Allow the kit to reach room temperature!

#### 4.1 Preparation of Washing Buffer

If any of the salts have been crystallized they should be resolved before use. Dilute 1 unit washing buffer concentrate with 9 units distilled water. The diluted buffer is stable for 6 weeks stored at 2 - 8 °C.

#### 4.2 Preparation of Sample Buffer

Dilute one unit sample buffer concentrate with 4 units distilled water. If any of the salts have been crystallized they should be resolved before use. The diluted buffer is stable for 6 weeks stored at 2 - 8 °C.

#### 4.3 Preparation of Standards

The standards are ready to use.

#### 4.4 Preparation of Control Sera

The control sera are ready to use.

#### 4.5 Preparation of Sera

Allow sera to reach room temperature (30 min). Dilute sera 1:100 with sample diluent buffer (10 µl serum with 1.0 ml buffer).

#### 4.6 Preparation of Conjugate

The daily required amount of conjugate solution should be prepared freshly. Do not use polystyrene tubes to prepare the conjugate dilution. Dilute conjugate 1:100 with conjugate diluent buffer (for 1 plate: 100 µl conjugate with 10 ml buffer, for 2 strips: 20 µl conjugate with 2 ml buffer). Remaining solution should be disposed.

#### 4.7 Preparation of the Substrate

The TMB-substrate solution is ready to use. Used substrate bottle should be closed carefully. Store substrate solution at 4 - 8 °C protected from light.

#### 4.8 Microtiter Strips

The strips are ready to use. Unused strips should be stored in the sealable original plastic bag at 2 - 8 °C.

#### 4.9 Stopping Solution

H<sub>2</sub>SO<sub>4</sub> (caution!)

### 5. Test Procedure

1. **Pipette 100 µl serum dilution** or (undiluted) standards and control sera into each well, for blanks use ready to use dilution buffer instead of serum dilution, seal wells with adhesive foil
2. **Incubate for 1 hour** at room temperature (RT)
3. **Rinse off the wells 3 x** with min. 200 µl washing buffer per well
4. **Pipette 100 µl of conjugate dilution** into each well, seal wells with adhesive foil

5. **Incubate for 30 minutes** at RT

6. **Rinse off the wells 3 x** with min. 200 µl washing buffer per well

7. **Pipette 100 µl substrate solution** into each well

8. **Incubate for 10 min** at RT in the dark. At a room temperature higher than 25 °C the substrate incubation time should be shortened. The minimum substrate incubation time must be 5 minutes.

9. **Pipette 100 µl stopping reagent** into each well

10. **Measure at 450 nm** within the next 30 min after stopping

### 6. Interpretation of Results:

Calibrate measured absorbance against concentrations/units of standards ( 12.5 U/ml, 25 U/ml, 50 U/ml, 100 U/ml, 200 U/ml), (linear or semilog). Read the units of the examined sera from the standard curve directly.

Results above 25 U/ml (cut off value) are considered positive.

To prove the test function the value of the positive control serum has to be within the range (see label on the vial). The negative control result has to be lower than the cut off value of the test.

### Precautions

For in vitro diagnostic use only.

The human Control Sera and Standards in this kit have been prepared from blood donations which have been tested for Hepatitis B Surface Antigen, anti-HCV- and anti-HIV 1/2 antibodies and shown to be NEGATIVE.

However, as no known test can guarantee the absence of an infectious virus, all reagents and samples must be handled carefully and disposed of in accordance with local legislation.



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