



Instructions For Use

Valid as of 01.11.2023

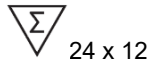
DotDiver Myositis 12

REF 5093

Dot immunoassay for the determination of IgG antibodies against nuclear and cytoplasmic antigens in human serum



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1 Intended Purpose

The DotDiver Myositis 12 is a qualitative dot immunoassay for the determination of IgG antibodies against nuclear and cytoplasmic antigens (Jo-1, PL-7, PL-12, EJ, SRP, Mi-2, MDA-5, TIF1- γ , SSA/Ro52kD, SAE1, SAE2 and NXP-2) in human serum.

The DotDiver Myositis 12 is intended as an aid in the diagnosis of autoimmune myositis diseases in conjunction with other clinical and laboratory findings.

The immunoassay is designed for semi-automated use with the DotDiver instrument.

The immunoassay is designed for professional *in vitro* diagnostic use.

2 Diagnostic Relevance

Polymyositis and dermatomyositis are idiopathic inflammatory muscle disorders with an unknown etiology. They are characterized by muscle weakness, atrophy of peripheral muscles, biochemical and electromyographic signs of muscle damage and characteristic skin alterations.

Autoantibodies to t-RNA synthetases like Jo-1, PL-7, PL-12 and EJ are characteristic markers in myositis patients. Antibodies to the signal recognition particle SRP are also associated with polymyositis patients. Mi-2 autoantibodies are detected almost exclusively in patients with dermatomyositis and indicate a relatively mild clinical course. Other specific autoantibody in dermatomyositis is directed to TIF1- γ and associated with aggressive skin lesions. High anti-TIF1- γ titers in adults are increasing the risk of cancer-associated dermatomyositis.

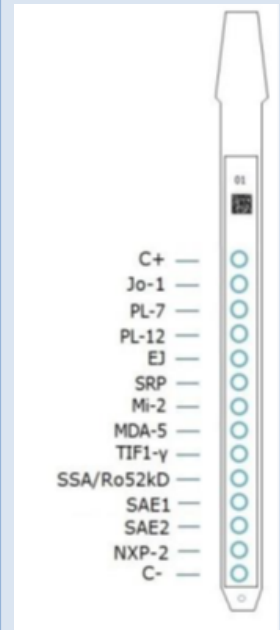
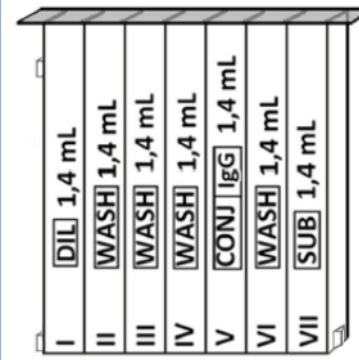
Antibodies against MDA5 are characteristic for amyopathic dermatomyositis with interstitial lung disease. In juvenile and adult dermatomyositis antibodies to NXP-2 are found in cases with severe skin lesions, calcinosis and muscle contractures, a tumor association is also described. SAE antibodies are appearing in patients with dermatomyositis too, systemic diseases are rare in these cases.

Autoantibodies directed to SS-A/Ro 52 kD are not associated to any clinical entity, but often are found as accompanying antibody in rheumatic diseases.

3 Test Principle

Dot immunoassays are frequently used for the determination of specific antibodies directed against multiple antigens. The test strips are coated with various antigens in consistent intervals. If antibodies are present in the patient's sample, they bind to the respective antigens. A secondary antibody conjugated with the enzyme alkaline phosphatase detects the generated immune complexes. A colorless substrate is converted into a colored, insoluble product. The signal intensity of the precipitated reaction product is proportional to the antibody activity in the sample.

4 Test Components

Component	Description
Test strips STRIP , 24 (3 x 8) pieces 	24 (3 x 8) breakable test strips on plastic supports (ready-to-use), each strip coated with dots of highly purified <ul style="list-style-type: none"> - Jo-1 - PL-7 - PL-12 - EJ - SRP - Mi-2 - MDA-5 - TIF1-γ - SSA / Ro 52 kDa - SAE1 - SAE2 and - NXP-2 as well as positive and negative control
Cartridges CART , 24 pieces with 7 compartments 	24 cartridges (ready-to-use) sealed with aluminum foil, each compartment contains reagents
Sample diluent DIL , 1 x 1.4 mL	Colored solution (contains ProClin 300)

Wash buffer WASH , 4 x 1.4 mL	Colorless solution (contains ProClin 300)
Conjugate IgG CONJ , 1 x 1.4 mL	Colored solution of polyclonal anti-human IgG antibody conjugated to alkaline phosphatase (contains ProClin 300)
Substrate BCIP / NBT SUB , 1 x 1.4 mL	Substrate solution of bromo-chloro-indolyl-phosphate and nitroblue tetrazolium (ready-to-use; contains sodium azide)
QC Certificate 1 piece	-
Instructions for Use 1 piece	-

5 Materials required but not provided

- Common laboratory equipment
- Precision pipette (10 µL) and disposable pipette tips
- DotDiver2.0 instrument or, alternatively, DotDiver instrument with additional scanner and Blot GALaxy evaluation software or equivalents
- Adsorbent paper or paper towel

6 Storage and Stability

Upon receipt, all test components must be stored at 2 °C to 8 °C, preferably in the original kit box. If stored properly in their original containers, all components are stable until their expiry date.

7 General Information

This product is for *in vitro* diagnostic use only. The instructions for use of the immunoassay as well as of the DotDiver device must be read carefully. They are valid only for the present product with the given composition and must be strictly followed to ensure reliable test results. Deviations can lead to erroneous test results. Components must not be exchanged by test reagents of different lots or of other manufacturers. Exposure of reagents to strong light must be avoided throughout the entire test procedure and storage.

8 Preparation

8.1 Preparation of Reagents

All components including the test strips must be brought to room temperature (RT: 18 °C to 25 °C) before use for at least 30 min. All liquid components must be mixed gently to ensure homogeneity.

8.1.1 Test Strips

The breakable test strips are provided on plastic supports within sealed aluminium pouches. Unused test strips should always be stored refrigerated and protected from moisture.

8.1.2 Cartridges

The cartridges are ready-to-use. They contain ready-to-use reagents in sufficient amounts for the analysis of one sample. Unused cartridges should always be stored refrigerated.

8.2 Preparation of Samples

8.2.1 Sample Material

The use of freshly collected serum from blood taken by venipuncture is recommended. The use of icteric, lipemic, hemolytic or bacterially contaminated samples should be

avoided. Insoluble substances must be removed from the sample by centrifugation. Samples must not be thermally inactivated.

8.2.2 Sample Storage

Samples may be kept at 2 °C to 8 °C up to three days. Long-term storage requires -20 °C. Repeated freezing and thawing should be avoided. For multiple use, samples should be aliquoted and kept at -20 °C.

9 Test Performance

9.1 Procedure

Start the DotDiver device. Place the test strips into the clamp and the cartridges into the corresponding positions of the cartridge holder and insert both into the instrument. Close the cover and follow the indications on the DotDiver screen.

Processing, analysis and evaluation of the immunoassay performed by the DotDiver2.0 are described below (Protocol 02):

Step	Description
1. Piercing	Piercing of well 1 of the cartridges by the instrument
2. Open cover	Open the cover of the instrument
3. Addition of samples	Add 10 µL of undiluted samples into well 1 of the cartridges and application of sample information
4. Close cover	Close the cover of the instrument
5. Piercing	Piercing of all other wells of the cartridges by the instrument
6. Pre-wash	Incubation of strips for 1 min. in well 2 (wash buffer) of the cartridge.
7. Sample Incubation	Incubation of strips for 30 min. in well 1 (dilution buffer) of the cartridge.
8. Wash	Incubation of strips for 2 min. in well 2 (wash buffer) of the cartridge.
9. Wash	Incubation of strips for 2 min. in well 3 (wash buffer) of the cartridge.
10. Wash	Incubation of strips for 2 min. in well 6 (wash buffer) of the cartridge.
11. Conjugate Incubation	Incubation of strips for 10 min. in well 5 (conjugate) of the cartridge.
12. Wash	Incubation of strips for 2 min. in well 4 (wash buffer) of the cartridge.
13. Wash	Incubation of strips for 2 min. in well 3 (wash buffer) of the cartridge.
14. Wash	Incubation of strips for 2 min. in well 2 (wash buffer) of the cartridge.
15. Substrate Incubation	Incubation of strips for 10 min. in well 7 (substrate) of the cartridge.
16. Wash	Incubation of strips for 2 min. in well 6 (wash buffer) of the cartridge.
17. Drying	Automated drying of test strips for about 25 min. by integrated fan.
18. Analysis	Automated imaging of the test strips by integrated camera
19. Evaluation	Automated evaluation of the test strips by integrated software

9.2 Automation

Processing, analysis and evaluation of the immunoassay is performed completely by the DotDiver2.0 device. For semi-automated use of the immunoassay with the DotDiver device, prepare test strips and cartridges according to the instructions for use of the instrument and follow the indications on the DotDiver screen. Processing of the immunoassay must be performed

analogous to the described procedure. Processed strips must be dried for approximately 30 min. prior scanning and evaluation of images by use of the Blot GALaxy evaluation software or equivalent.

10 Test Evaluation

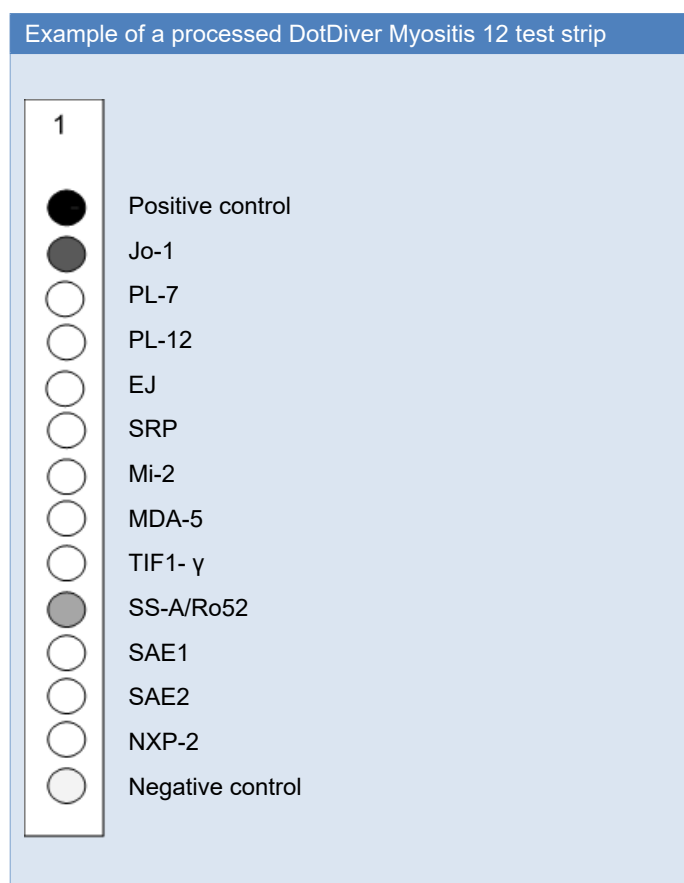
10.1 Metrological Traceability

The immunoassay is calibrated using internal reference samples.

10.2 Evaluation

The qualitative evaluation of is performed by comparison of the antigen specific signal intensities of the patient's sample with the negative control dot on the test strip. The negative control dot serves as a cut off control for qualitative evaluation.

The test strips are coated with antigens. The color intensity is proportional to the specific antibody activity in the sample. A sample is considered to be positive for a specific antigen, if the coloration of the antigen dot shows a more intense coloration than the negative control dot on the test strip. A sample is considered to be negative for a specific antigen, if the coloration of the antigen dot shows a less intense or equal coloration compared to the negative control dot on the test strip.



10.3 Criteria of Validity

Test runs are only valid if the following criteria of validity are fulfilled:

- Intensity negative control < intensity positive control
- The positive control must be evaluated positive.

The coloration of these controls ensures that the test has been performed correctly. If these criteria are not met, the test is not valid and must be repeated.

10.4 Troubleshooting

In case of an invalid test run, the expiry dates and storage conditions, incubation times and temperatures, and precise calibration of all instruments used should be verified. If no reason for an invalid test run could be identified, please contact the supplier or manufacturer of the product.

10.5 Reference Ranges

As a result of different seroprevalences in individual regions, each laboratory should verify the reference ranges by own analysis and adapt, if necessary.

10.6 Interpretation of Test Results

A positive test result indicates the presence of specific antibodies. A negative result indicates the absence of specific antibodies, but does not exclude the possibility of an autoimmune reaction. In case of a borderline test result, a reliable evaluation is not possible.

10.7 Limitations of the Method

The interpretation of test results must always be considered in combination with the clinical picture of the patient. The diagnosis should not be based on the results of a sole diagnostic method. All clinical and laboratory findings should be evaluated to state a diagnosis. For confirmation, further investigations should be carried out.

11 Performance Characteristics

11.1 Analytical Performance Characteristics

11.1.1 Precision

The precision of test results was assessed by the determination of the intra- and interassay variation by the analysis of multiple samples with different antibody activities. No differences in the qualitative evaluation have been observed.

11.2 Diagnostic Performance Characteristics

11.2.1 Diagnostic Sensitivity and Specificity

The sensitivity and specificity were assessed by the analysis of characterized samples (confirmed positive or negative for specific antibodies by reference laboratories and/or methodologies).

	Sensitivity	Specificity
Jo-1	> 99 %	> 99 %
PL-7	> 99 %	> 99 %
PL-12	> 99 %	> 99 %
EJ	> 99 %	> 99 %
SRP	> 99 %	> 99 %
Mi-2	> 99 %	> 99 %
MDA-5	> 99 %	> 99 %
TIF1-γ	> 99 %	> 99 %
SS-A/Ro-52	-	-
SAE1	-	-
SAE-2	-	-
NXP2	-	-

12 Warnings and Precautions

The product is designed exclusively for *in vitro* diagnostic use by qualified, authorized and trained personnel. All test components and human samples should be handled with care as potentially hazardous. Good laboratory practices (GLP) and all relevant regulations should be adhered to.

In case the product is damaged or product information including labelling is wrong or incorrect, please contact the manufacturer or supplier.

This product contains preparations of human and / or animal origin. Any material derived from human body fluids or organs used for the preparation of components were tested and found negative for HBsAg (Hepatitis B-Virus-surface Antigen) and anti-HIV as well as anti-HCV antibodies. However, all components

and all patient samples should be handled as potentially hazardous in accordance with national laws and appropriate guidelines on biological safety.

As the product contains potentially hazardous materials, the following precautions should be followed: Do not smoke, eat or drink while handling kit material or samples. Avoid direct contact to kit material or samples by wearing protective gloves laboratory coat and safety glasses. Never pipette material by mouth. Wipe up spills promptly and wash the affected surface thoroughly with a decontaminant. Wash hands thoroughly after use.

Some of the reagents contain ProClin (< 0.0015 %) as a preservative, and must not be swallowed or allowed to come into contact with skin or mucosa.

Some of the reagents contain sodium azide (< 0.1 %) as a preservative and must not be swallowed or allowed to come into contact with skin or mucosa. The possible formation of heavy metal azides in the drainage has to be prevented by sufficient rinsing with water.

The information in the safety data sheet on possible hazards, first aid measures, measures in the event of the unintentional release of large quantities, handling and storage, personal protective equipment, information on disposal as well as information on toxicology must be observed.

Any serious incident that has occurred in relation to the device shall be reported to the manufacturer and the competent authority of the member state in which the user and/or the patient is established.





















13 Disposal

For decontamination and disposal the recommendations of the CDC as well as the relevant local and national environmental guidelines and regulations should be adhered to. Samples, potentially contaminated materials and infectious waste must be decontaminated, e.g. by autoclaving for 20 min. at 121 °C.

14 References

- Casciola-Rosen L, Mammen AL: Myositis autoantibodies. *Curr. Opin. Rheumatol.* 2012, 24, 3 – 11.
- Ghirardello A, Borella E, Beggio M, Franceschini F, Fredi M, Doria A: Myositis autoantibodies and clinical phenotypes. *Autoimmun. Highlights* 2014, 5, 69 – 75.
- Gunawardena H, Betteridge ZE, McHugh NJ: Myositis-specific autoantibodies: their clinical and pathogenic significance in disease expression. *Rheumatology* 2009, 48, 607 – 12.
- Peixoto D, Costa J, Ferretti M, Malattia C, Martini A: New autoantibodies and their clinical associations in juvenile myositis – a systematic review. *Acta Rheumatol. Port.* 2013, 38, 234 – 41.

15 Symbols

	Manufacturer
	CE marking of conformity
	<i>In vitro</i> diagnostic medical device
	Catalogue number
	Unique device identifier
	Batch code
	Temperature limit
	Use-by date
	Consult instructions for use
	Contains sufficient for <n> tests
	Do not re-use
	Caution
	Biological risk
	Keep away from sunlight
	Test strip
	Cartridge
	Sample diluent
	Conjugate
	Wash buffer
	Substrate

16 Changes

Changes in current Instructions for Use	
Current Version	006/11.2023
Summary of Changes	No change, harmonisation of language variants.