



INSTRUCTION MANUAL

REF 4076

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ANAcyto^{plus} Dot

- 24 x 6 determinations -



IVD *In vitro* diagnostic device

Immunodot for the determination of IgG antibodies to M2, Jo-1, PL-7, PL-12, SRP and ribosomes in human serum or plasma

REF	Catalogue number	LOT	Batch code
	Consult accompanying documents		Manufactured by
	Temperature limitation		Use by
	Consult operating instruction		Biological risk



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INTENDED USE

ANAcyto^{plus} Dot is used for the qualitative determination of IgG autoantibodies to M2, Jo-1, PL-7, PL-12, SRP and ribosomes, in human serum or plasma for the differential diagnosis of autoimmune diseases.

Cytoplasmic staining of HEp2 cells for the detection of antinuclear antibodies (ANA) may be caused by different autoantigens such as M2, t-RNA-synthetases or ribosomes. Since these parameters are pathognomonic for different clinical entities the detection of the specificity of antibodies with cytoplasmic staining is of importance.

The ANAcyto Dot is a reliable and easy-to-use tool for the detection of antibodies recognizing cytoplasmic antigens.

Primary biliary cirrhosis (PBC) is characterized by occurrence of antibodies to mitochondrial antigens. Antibody M2 seems to be the most specific out of nine described so far for PBC. M2 antibodies mainly react with epitopes of the E2 component of the pyruvate-dehydrogenase complex found in mitochondria. M2 has been demonstrated in+ 95% of all PBC patients.

In addition to nuclear antibodies autoantibodies to ribosomal phosphoproteins can be found in up to 20 % of SLE patients. Due to the ready correlation with disease activity these autoantibodies can be used for the follow-up.

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. The criteria of Bohan and Peter are used for the classification of these disorders. Very often autoantibodies to t-RNA transferases like Jo-1, PL-7, and PL-12 are found in myositis patients. Another marker for autoimmune myositis are autoantibodies to Signal recognition particle SRP.

Czaja AL. Natural history, clinical features and treatment of autoimmune hepatitis. *Semin Liver Dis* (1987) 4: 1 – 12

Johnson PJ, McFarlane IA. Meeting Report: International Autoimmune Hepatitis Group. *Hepatology* (1993) 18: 998 - 1005

Tan EM.: Antinuclear antibodies: diagnostic markers for autoimmune diseases and probes for cell biology. *Adv Immunol* 1989 44:93-151

PRINCIPLE of the TEST

ANAcyto^{plus} Dot is a sensitive immunodot assay for the determination of IgG antibodies to M2, Jo-1, PL-7, PL-12, SRP and ribosomes, in human serum or plasma, respectively.

ANAcyto^{plus} Dot includes 24 numbered test strips with 7 dots fixed on a plastic support: Dots are coated with M2, Jo-1, PL-7, PL-12, SRP and ribosomes, Two test dots serve as positive and negative controls.

Patient sera and strips are incubated in the test tray. During the first incubation antibodies of the patient sample bind to the target antigens immobilized on the solid-phase of the strips. Following an incubation period of 30 min unbound serum components are removed by a washing step.

Bound antibodies react specifically with anti-human-IgG conjugated to alkaline phosphatase. Following an incubation period of 30 min excessive conjugate is separated from the solid-phase immune complexes by an additional washing step.

Alkaline phosphatase converts the colourless substrate solution into a dark purple precipitating dot. After 10 - 12 minutes while shaking the reaction is stopped by a washing step.

Strips are dried for at least 30 min by pressing the reactive side onto absorbent paper. Results are regarded to be positive if the colouration of the test dot is more intense than the colouration of the negative control.

PATIENT SAMPLES

Specimen collection and storage

Blood is taken by venipuncture. Serum is separated after clotting by centrifugation. Plasma can also be used.

The samples may be kept at 2 - 8 °C for up to three days. Long-term storage requires - 20 °C.

Repeated freezing and thawing should be avoided. If samples are to be used for several assays, initially aliquot samples and keep at - 20 °C.

Preparation before use

Allow all components to reach room temperature prior to use in the assay. Take care to agitate serum samples gently in order to ensure homogeneity.

TEST COMPONENTS for 24 x 6 determinations

A	Dot strips	24
Ag	24 numbered strips coated with 6 specific antigens	24 dot strips for the determination of six antibody specificities each
	M2 (mitochondrial membran enzymes: PDC, OGDC und BCOADC; human recombinant)	
	Jo-1 (Histidyl-tRNA-synthetase, bovin)	
	PL-7 (Thronyl-tRNA-synthetase, human recombinant)	
	PL-12 (Alanyl-tRNA-synthetase; human recombinant)	
	SRP (human recombinant)	
	Ribosomes (P0; human recombinant)	
	- Positive control	
	- Negative control	
B	Wash buffer	40 ml
BUF	sufficient for 400 ml solution	concentrate capped blue
WASH	10x	
C	Sample diluent	40 ml
DIL	(coloured yellow)	capped yellow
D	Conjugate	40 ml
CONJ	anti-human-IgG (goat) coupled with alkaline phosphatase (coloured red)	ready for use capped red
E	Substrate	40 ml
SOLN	nitroblue tetrazolium with bromo-chloro-indolyl-phosphate	ready for use capped black
NBT/BCIP	(black bottle)	
F	Incubation tray for 8 strips	3 x

Materials required

- micropipette 100 - 1000 µl
- micropipette 10 - 100 µl
- pipette tips
- graduated cylinders
- distilled or deionised water
- plate shaker
- plastic pincers
- paper towel

Size and storage

ANAcyto^{plus} Dot has been designed for 24 x 6 determinations.

The expiry date of each component is reported on its respective label, that of the complete kit on the box label.

Upon receipt all components of the ANAcyto^{plus} Dot have to be kept at 2 - 8 °C, preferably in the original kit box.

After opening all kit components are stable for at least 2 months, provided proper storage.

Preparation before use

Allow all components to reach room temperature prior to use in the assay.

Prepare a sufficient amount of wash solution by diluting the concentrated washing buffer 10 times (1 + 9) with deionised or distilled water. For example, dilute 10 ml of the concentrate with 90 ml of deionised or distilled water.

For each test strip 15 ml of washing buffer are requested.

The wash solution prepared is stable at 2 - 8 °C up to 30 days.

All other components are ready for use and so stable until the expiry date.

Avoid exposure of the substrate solution to light.

After each filling of wells with solution, agitate the incubation tray manually to ensure strips are completely immersed and to remove air bubbles which may be trapped under the strip.

ASSAY PROCEDURE

1. Bring all reagents to room temperature (RT) (18-25°C) before use. Mix gently without causing foam.
2. Place the strips with the reactive side up (labels on top) into the respective well. Dispense 2 ml of wash solution (made of B) into the respective wells.
3. Cover tray, incubate **10 min** at RT (18-25°C) while shaking.
4. Discard wash solution. (Discard the solution in the wells by slowly inverting the plate. Dry the edges of the tray with absorbent paper in order to remove the remaining fluid.)
5. Add **1.5 ml** sample diluent (C) and **10 µl** patient serum or plasma to the respective wells.
6. Cover tray and incubate **30 min** at RT (18-25°C) while shaking.
7. Decant or aspirate, wash each well **three times three minutes** with **1.5 ml** wash solution (made of B) while shaking. (Discard the solution in the wells by slowly inverting the plate. Dry the edges of the tray with absorbent paper in order to remove the remaining fluid.)
8. Add 1.5 ml conjugate (D) to each well
9. Cover tray and incubate **30 min** at RT (18-25°C) while shaking.
10. Decant or aspirate, wash each well **three times three minutes** with **1.5 ml** wash solution (made of B) while shaking. (Discard the solution in the wells by slowly inverting the plate. Dry the edges of the tray with absorbent paper in order to remove the remaining fluid.)
11. Add **1.5 ml** of substrate (E) to each well.
12. Cover plate, incubate **10-12 min** while shaking.
13. Decant or aspirate, wash each well **once three minutes** with wash solution (made of B) while shaking to stop the reaction. (Discard the solution in the wells by slowly inverting the plate. Dry the edges of the tray with absorbent paper in order to remove the remaining fluid.)
14. Collect the strips from the wells and dry the membranes by pressing briefly the reactive side of the strip onto absorbent paper. After approximately 30 min the strips are to be interpreted.

EVALUATION OF RESULTS

Evaluation:

Results should be interpreted only after strips have been dried for at least 30 minutes.

The **positive control** must be positive in all cases. The colouration of the dot ensures that the test has been run correctly and the kit components are not degraded. If the positive control dot shows no colouration the results cannot be interpreted.

The **negative control** demonstrates the extent of non-specific antibody binding of the sample in the test. The colouration of the dot corresponds to the minimal intensity above which a sample is considered positive.

The test dots are coated with autoantigens and detect specific antibody binding of the sample in the test. The colour intensity of the test dot depends on the titer of specific antibody binding in the sample. The patient sample is positive concerning a certain antibody if the test dot colouration is stronger (more intense) than the negative control.

Test example

1	ANAcyto ^{plus} Dot	result:
●	- Positive control	positive
●	- M2	positive
○	- Jo-1	
○	- PL-7	
○	- PL-12	
○	- SRP	
○	- ribosomes	
○	- Negative control	

Positive result:

A sample is considered to be positive for autoantibodies to M2, Jo-1, PL-7, PL-12, SRP and ribosomes if the colouration of the test dot is more intense than the colouration of the negative control.

The colour intensity of the negative dot depends on the test conditions (e.g. incubation times, temperature, washing efficiency) and on the composition of each individual sample. It might be uncoloured even if the test has been run in optimal conditions.

Negative result:

A sample is considered to be negative for autoantibodies to M2, Jo-1, PL-7, PL-12, SRP and ribosomes if the colouration of the test dot is less intense than the colouration of the negative control.

Limitations of Method

Healthy individuals should be tested negative by the ANAcyto^{plus} Dot. However, M2, Jo-1, PL-7, PL-12, SRP and ribosomes autoantibody positive apparently healthy individuals do occur.

Any clinical diagnosis should not be based on the results of in vitro diagnostic methods alone. Physicians are supposed to consider all clinical and laboratory findings possible to state a diagnosis.

CHARACTERISTIC ASSAY DATA

Sensitivity and specificity

Clinically defined populations (confirmed positive with disease specific reference methodologies) have been used for checking the sensitivity. Specificity was checked with control groups that embrace a normal healthy population as well as clinically defined control groups.

Sensitivity:

M2, Jo-1, PL-7, PL-12, SRP, Ribosomes > 99%

Specificity:

M2, Jo-1, PL-7, PL-12, SRP, Ribosomes > 99%

Reproducibility

The dot assay is a qualitative test and the precision is evaluated in terms of variation of the visual colour of the test. Three control sera (high, medium, low positive) were assayed for intraassay and interassay imprecision in a statistically relevant repetition.

REMARKS:

INCUBATION SCHEME

ANAcyto^{plus} Dot (4076)

1.	Bring all reagents and the requested number of strips to room temperature (18-25°C)
2.	Place the strips with the reactive side upside in the tray and dispense 2 ml of wash solution (made of B) into the respective wells
3.	Seal plate and incubate while shaking 10 minutes, room temperature (18-25°C)
4.	Discard wash solution
5.	Pipette 1.5 ml sample diluent (C) and 10 µl patient serum or plasma (1 + 150) into each well
6.	Incubate while shaking 30 minutes, room temperature (18-25°C)
7.	Decant, wash strips while shaking 3 x 3 minutes with 1.5 ml (made of B)
8.	Pipette 1.5 ml conjugate (D) in the respective well
9.	Incubate while shaking 30 minutes, room temperature (18-25°C)
10.	Decant, wash strips while shaking 3 x 3 minutes with 1.5 ml (made of B)
11.	Pipette 1.5 ml substrate (E)
12.	Incubate while shaking 10 - 12 minutes, room temperature (18-25°C)
13.	Decant, wash strips to stop reaction while shaking 1 x 3 minutes with 1.5 ml (made of B)
14.	Dry membranes by pressing the strip onto absorbent paper. After approximately 30 min the strips are ready to be interpreted.

SAFETY PRECAUTIONS

- **This kit is for in vitro use only.** Follow the working instructions carefully. GA GENERIC ASSAYS GmbH and its authorized distributors shall not be liable for damages indirectly or consequentially brought about by changing or modifying the procedure indicated. The kit should be performed by trained technical staff only.
- The expiration dates stated on the respective labels are to be observed. The same relates to the stability stated for re-constituted reagents.
- Do not use or mix reagents from different lots.
- Do not use reagents from other manufacturers.
- Avoid time shift during pipetting of reagents.
- All reagents should be kept at 2 - 8 °C prior use in the original shipping container.
- Some of the reagents contain small amounts of bromonitrodioxane (< 0.01 % w/w), methylisothiazolones (< 20 ppm) or sodium azide (< 0.05 %) as a preservative. They must not be swallowed or allowed to come into contact with skin or mucosa.
- Since the kit contains potentially hazardous materials the following precautions should be observed:
 - Do not smoke, eat or drink while handling kit material,
 - Always use protective gloves,
 - Never pipette material by mouth,
 - Wipe up spills promptly, washing the affected surface thoroughly with a decontaminant.
- In any case GLP should be applied with all general and individual regulations to the use of this kit.