



RUO in the USA

Revised 23 May 2011 rm (Vers. 2.1)

Please use only the valid version of the package insert provided with the kit.

This kit is intended for Research Use Only.

Not for use in diagnostic procedures.

NAME AND INTENDED USE 1

ENA-4-Profile ELISA is an indirect solid phase enzyme immunometric assay (ELISA). It is designed for determination of IgG class autoantibodies directed against the extractable nuclear antigens SS-A (Ro), SS-B (La), Sm and RNP/Sm in human serum or plasma.

PRINCIPLE OF THE TEST 2

Extractable nuclear antigens SS-A (Ro), SS-B (La), Sm, RNP/Sm, are bound to microwells. Antibodies against these antigens, if present in diluted serum or plasma, bind to the respective antigens. Washing of the microwells removes unspecific serum and plasma components. Horseradish peroxidase (HRP) conjugated anti-human IgG immunologically detects the bound patient antibodies forming a conjugate/antibody/antigen complex. Washing of the microwells removes unbound conjugate. An enzyme substrate in the presence of bound conjugate hydrolyses to form a blue colour. The addition of an acid stops the reaction forming a yellow end-product. The intensity of this yellow colour is measured photometrically at 450 nm. The amount of colour is directly proportional to the concentration of IgG anti-bodies present in the original sample.

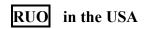
WARNINGS AND PRECAUTIONS 3

- 1. Do not interchange kit components from different lots.
- Components containing human serum were tested and found negative for HBsAg, HCV, HIV1 and HIV2 by FDA 2. approved methods. No test can guarantee the absence of HBsAg, HCV, HIV1 or HIV2, and so all human serum based reagents in this kit must be handled as though capable of transmitting infection.
- Avoid contact with the TMB (3,3',5,5'-Tetramethyl-benzidine). If TMB comes into contact with skin, wash 3. thoroughly with water and soap.
- 4. Avoid contact with the Stop Solution which is acid. If it comes into contact with skin, wash thoroughly with water and seek medical attention.
- Some kit components (i.e. Controls, Sample buffer and Buffered Wash Solution) contain Sodium Azide as 5. preservative. Sodium Azide (NaN₃) is highly toxic and reactive in pure form. At the product concentrations (0.09%), though not hazardous. Despite the classification as non-hazardous, we strongly recommend using prudent laboratory practices (see 8., 9., 10.)
- Some kit components contain Proclin 300 as preservative. When disposing reagents containing Proclin 300, flush 6. drains with copious amounts of water to dilute the components below active levels.
- Wear disposable gloves while handling specimens or kit reagents and wash hands thoroughly afterwards. 7.

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- Do not pipette by mouth. 8.
- 9. Do not eat, drink, smoke or apply makeup in areas where specimens or kit reagents are handled.
- 10. Avoid contact between the buffered Peroxide Solution and easily oxidized materials; extreme temperature may initiate spontaneous combustion.

Observe the guidelines for performing quality control in medical laboratories by assaying controls and/or pooled sera. During handling of all kit reagents, controls and serum samples observe the existing legal regulations.

CONTENTS OF THE KIT 4

Package size 4 x 24 determ.

Qty.1	Divisible microplate consisting of 3 modules of 8 wells each, coated with individual highly purified antigens: SS-A, SS-B, Sm, RNP/Sm. Ready to use.							
	The modules are colour coded: SS-A red	SS-B Sm RNP / Sm	blue yellow green					
	Anti-ENA controls in a serum/buffer matrix (PBS, $NaN_3 < 0.1\%$ (w/w)). Ready to use.							
1 vial, 2 ml	Positive Control (1) (100 U/ml).							
1 vial, 2 ml	Negative Control (2) (0 U/ml).							
4 vials, 1.5 ml each	Cut-off Calibrators (25 U/ml), specific for each individual antigen.							
1 vial, 20 ml	Sample buffer (Tris, NaN ₃ <0.1% (w/w)), yellow, concentrate (5x).							
1 vial, 15 ml	Enzyme conjugate solution (PBS, Proclin 300 <0.5% (v/v)), (light red) containing polyclonal rabbit anti-human IgG, labelled with horseradish peroxidase. Ready to use.							
1 vial, 15 ml	TMB substrate solution. Ready to use.							
1 vial, 15 ml	Stop solution (contains acid). Ready to use.							
1 vial, 20 ml	Wash solution (PBS, NaN ₃ <0.1% (w/w)), concentrate (50x).							

5 STORAGE AND STABILITY

- Store the kit at 2-8 °C. 1.
- 2. Keep microplate wells sealed in a dry bag with desiccants.
- 3. The reagents are stable until expiration of the kit.
- Do not expose test reagents to heat, sun or strong light during storage and usage. 4.

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Diluted sample buffer and wash buffer are stable for at least 30 days when stored at 2-8 °C. 5.

MATERIALS REQUIRED 6

Equipment

- Microplate reader capable of endpoint measurements at 450 nm
- Multi-Channel Dispenser or repeatable pipette for 100 μl
- Vortex mixer
- Pipettes for 10 μ l, 100 μ l and 1000 μ l
- Laboratory timing device
- Data reduction software

Preparation of reagents

- Distilled or deionised water
- Graduated cylinder for 100 and 1000 ml
- Plastic container for storage of the wash solution

7 SPECIMEN COLLECTION, STORAGE AND HANDLING

- 1. Collect whole blood specimens using acceptable medical techniques to avoid hemolysis.
- 2. Allow blood to clot and separate the serum by centrifugation.
- Test serum should be clear and non-hemolysed. Contamination by hemolysis or lipemia is best avoided, but does not 3. interfere with this assay.
- 4. Specimens may be refrigerated at 2-8 °C for up to five days or stored at -20 °C up to six months.
- Avoid repetitive freezing and thawing of serum samples. This may result in variable loss of autoantibody activity. 5.
- Testing of heat-inactivated sera is not recommended. 6.

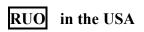
8 **PROCEDURAL NOTES**

- 1. Do not use kit components beyond their expiration dates.
- 2. Do not interchange kit components from different lots.
- 3. All materials must be at room temperature (20-28 °C).
- Have all reagents and samples ready before start of the assay. Once started, the test must be performed without 4. interruption to get the most reliable and consistent results.
- 5. Perform the assay steps only in the order indicated.
- 6. Always use fresh sample dilutions.
- 7. Pipette all reagents and samples into the bottom of the wells.
- To avoid carryover contamination, change the tip between samples and different kit controls. 8.

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- 9. It is important to wash microwells thoroughly and remove the last droplets of wash buffer to achieve best results.
- 10. All incubation steps must be accurately timed.
- 11. Control sera or pools should routinely be assayed as unknowns to check performance of the reagents and the assay.
- 12. Do not re-use microplate wells.

For all controls, the respective concentrations are provided on the labels of each vial. Using these concentrations a calibration curve may be calculated to read off the patient results semi quantitatively.

PREPARATION OF REAGENTS 9

9.1 **Preparation of sample buffer**

Dilute the contents of each vial of the sample buffer concentrate (5x) with distilled or deionised water to a final volume of 100 ml prior to use.

Store refrigerated: stable at 2-8 °C for at least 30 days after preparation or until the expiration date printed on the label.

9.2 **Preparation of wash solution**

Dilute the contents of each vial of the buffered wash solution concentrate (50x) with distilled or deionised water to a final volume of 1000 ml prior to use.

Store refrigerated: stable at 2-8 °C for at least 30 days after preparation or until the expiration date printed on the label.

9.3 Sample preparation

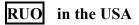
Dilute all samples 1:100 with sample buffer before assay. Therefore combine 10 μ l of sample with 990 μ l of sample buffer in a polystyrene tube. Mix well.

Controls are ready to use and need not be diluted.

10 TEST PROCEDURE

- 1. Prepare a sufficient number of microplate modules to accommodate controls and prediluted patient samples.
- 2. Pipette 100 µl of controls and prediluted patient samples in duplicate into the wells.





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	SS-A	SS-B	Sm	RNP/Sm	
	1	2	3	4	
A	NC	NC	NC	NC	
В	CalC	CalC	CalC	CalC	NC Negative control
С	PC	РС	РС	PC	CalC Calibration control
D	P1	P1	P1	P1	PC Positive control P1, P2 Patient sample
Ε	P2	P2	P2	P2	11,121 uttent sumpte
F	P3	P3	P3	P3	
G	P4	P4	P4	P4	
Η	P5	P5	P5	P5	

- Incubate for 30 minutes at room temperature (20-28 °C). 3.
- Discard the contents of the microwells and wash 3 times with 300 µl of wash solution. 4.
- Dispense 100 µl of enzyme conjugate into each well. 5.
- 6. Incubate for 15 minutes at room temperature.
- 7. Discard the contents of the microwells and wash 3 times with 300 µl of wash solution.
- 8. Dispense 100 μ l of TMB substrate solution into each well.
- 9. Incubate for 15 minutes at room temperature.
- 10. Add 100 μ l of stop solution to each well of the modules and incubate for 5 minutes at room temperature.
- 11. Read the optical density at 450 nm and calculate the results. Bi-chromatic measurement with a reference at 600-690 nm is recommended.

The developed colour is stable for at least 30 minutes. Read optical densities during this time.

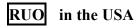
11 REFERENCES / LITERATURE

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