RF latex

Qualitative and semiguantitative determination by latex agglutination on slide of Rheumatoid Factor in serum

2 x 5 ml **REF 6102**

Also available:

RF latex kit complete with controls and accessories **REF 6002**

PRINCIPLE

The rheumatoid disease is a systemic morbid disease associated with a chronic inflammatory process. From a diagnostic perspective it is characterized by VES, α -2 gamma globulins and CRP increase and by the presence of rheumatoid factors.

In this test the rheumatoid factors, if present in the serum, agglutinate with latex particles sensitized with human immunoglobulins to form an easily visible agglomerate.

REAGENTS

Kits components: **REAGENT 1** (liquid) **REF 6102** 1 x 5 ml

Latex polystyrene particles in suspension sensitized with human immunoglobulins, ready to use.

WARNING: the reagents contain sodium azide (< 0.1%) as preservative. Handle with caution and avoid ingestion and contact with skin.

SLIDE

STABILITY: sealed and stored at 2-8°C, reagents are stable up to the expiration date printed on the label.

SAMPLE

STABILITY: 2 days at 2-8°C, 1 month at -20°C.

PREPARATION OF THE REAGENTS

Let the reagents reach the test temperature before use. Shake the Reagent 1 accurately before use. Make sure all latex particles are in suspension and not deposited on the bottom of the vial.

MANUAL PROCEDURE OF THE QUALITATIVE TEST

Pour the reagents in the proper areas of the slide, as the following table:

	Sample	Pos. Control	Neg. Control
Sample	50 µl		
Positive control		1 drop	
Negative control			1 drop
Reagent 1	1 drop	1 drop	1 drop

Mix to spread the liquid on the whole area of the slide. Shake the slide gently (round movement) for 2 minutes and then evaluate the results.

READING OF THE RESULTS

The presence of any agglutination in the sample shows the rheumatoid factor concentrations of at least 8 IU/ml. On positive sera, it is recommended to perform the title with the semiquantitative test as further on described.

MANUAL PROCEDURE WITH THE SEMIQUANTITATIVE **TEST**

Dilute the sample with saline solution as follows:

Dilution	1:2	1:4	1:8	1:16	1:32
Sensitivity equal UI/ml	16	32	64	128	256

Proceed on each dilution same as for the qualitative test.

RESULTS

The last dilution which shows any agglutination corresponds to the sample title.

REFERENCE VALUES

RF is usually not found in serum.

NOTES

- Longer reaction times may give false positive results. 1.
- 2. Always compare the results with the controls.
- All reagents have been deactivated and tested for HIV, HBsAg and HCV antibodies. However, they should be treated as potentially infectious.
- After an accurate wash, slides can be reused maximum three times.

REFERENCES

Waaler E. Acta Path. Microb. Scand. 17,1-2, (1940) Muller W. "The Serology of Rheumatoid Arthritis" Gottingen 97 (1962)

KEY SYMBOLS

IVD	In Vitro diagnostic medical device
LOT	batch number
REF	catalog number
X	temperature limits
Ω	use by
\triangle	caution
[]i	read instructions for use





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MANUFACTURER



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