

	Laboratory investiga	поп кероп		
Patient Name	Centre OP/IP No/UHID			
Age/Gender				
MaxID/Lab ID	Collection Date/Time			
Ref Doctor	Reporting Date/Time			
	Hematology Specia	al	SIN No:B2B3873045	
	Max SLE Profile		3117 100.0203073043	
Test Name	Result	Unit	Bio Ref Interval	
ANA By Immunofluorescence, Serur	n			
Anti Nuclear Antibodies	Negative		Negative	
Primary Dilution	1:40			

Laboratory Investigation Report

## Interpretation

Anti Nuclear Antibody IFA, HEP2000, Serum Immunofluorescence

(Syn: Anti-Nuclear Antibody)

ANA immunofluorescence is the gold standard test for screening for autoimmune antibodies and has higher sensitivity as compared to ANA ELISA. False ANA positivity may be seen in - certain viral infections (Hepatitis C, Parvovirus and many other), bacterial infections (Tuberculosis), parasitic infection (schistosomiasis), certain malignancies and medications.ANA Immunofluorescence results need to be corroborated with clinical features and other laboratory findings for definitive evidence of auto-immune disorder.

Advise: -

- A repeat ANA testing is recommended after 12 weeks after an acute episode of infection.
- ANA LIA should be added in cases with positive ANA Immunofluorescence result to know which extractable nuclear antigen is present in the patients, which helps in classifying patients for specific autoimmune disorder.

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Poonam. S. Das, M.D. Principal Director-Max Lab & Blood Bank Services

Dr. Dilip Kumar M.D. Associate Director & Manager Quality

02-

Dr. Nitin Dayal, M.D. Principal Consultant & Head, Haematopathology

Test Performed at :910 - Max Hospital - Saket M S S H, Press Enclave Road, Mandir Marg, Saket, New Delhi, Delhi 110017 Booking Centre :5124 - Anuja Jaswal, Booth No 53, Phase 5, Sector 59, 9815446080 The authenticity of the report can be verified by scanning the Q R Code on top of the page

Max Lab Limited (A Wholly Owned Subsidiary of Max Healthcare Institute Ltd.)

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Conditions of Reporting: 1. The tests are carried out in the lab with the presumption that the specimen belongs to the patient name as identified in the bill/test request form. 2. The test results relate specifically to the sample received in the lab and are presumed to have been generated and transported per specific instructions given by the physicians/laboratory. 3. The reported results are for the information and interpretation by the referring doctor only. 4. Some tests are referred to other laboratories to provide a wider test menu to the customer. 5. Max Healthcare shall in no event be lable for accidental damages loss, or destruction of specimen which is not attributable to any direct and mala fide act or omission of Max Healthcare or its employees. Liability of Max Healthcare for deficiency of services, or other errors and omissions shall be limited to fee paid by the patient for the relevant laboratory services.



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-	Laboratory Investigat	ion Report		
Patient Name	Centre OP/IP No/UHID Collection Date/Time			
Age/Gender MaxID/Lab ID				
Ref Doctor				
	Re	porting Date/Time		
	Serology Special		SIN No:B2B3873045	
	Max SLE Profile			
Test Name	Result	Unit	Bio Ref Interval	
ANA - LIA,Serum LIA				1
dsDNA	Negative			
Nucleosome	Negative			
Histones	Negative			
SmD1	Negative			
PCNA	Negative			
PO (RPP)	Negative			
SS-A/Ro60	Negative			
SS-A/Ro52	Negative			
SS-B/La	Negative			
CENP-B	Negative			
Scl70	Negative			
U1-snRNP	Negative			
AMA M2	Negative			
Jo-1	Negative			
PM-Scl	Negative			
Mi-2	Negative			
Ku	Negative			
DFS70	Negative			

## Interpretation

- 1. The test provides a differential diagnosis using 17 different autoantibodies which is an additional diagnostic survey of autoimuune diseases like SLE, Mixed connective tissue diseases, Rheumatoid arthritis, Sjorgensyndrome, Progressive systemic sclerosis and CREST syndrome.
- 2. A definitive clinical diagnosis should not be made by result of a single test only, but should be made by taking clinical history and other laboratory findings in to account.

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MaxID/Lab ID	Collection Date/Time			
Ref Doctor	Reporting Date/Time			
	Serology Spec	ial		
	Max SLE Profi	le	SIN No:B2B3873045	
Test Name	Result	Unit	Bio Ref Interval	
Anti dsDNA Antibody (Double Stran FEIA	nded), Serum			
Anti-dsDNA FEIA	0.3	IU/i	mL	
Ref. Range				

Negative <10 Equivocal 10 - 15 Positive>15

## Comments:

The determination of ANA is important for the clinical diagnosis of connective tissue disease, ds DNA antibodies. It represents one of the diagnostic criteria for SLE.

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Principal Director-

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MC-2714

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Ref Doctor	Reporting Date/Time			
	Hematology Specia	I	SIN No:B2B3873045	
	Max SLE Profile		514 10.02050/5045	
Test Name	Result	Unit	Bio Ref Interval	
Anti dsDNA by Immunofluorescer	nce (IF), Serum			
Ds DNA Antibody	Negative			
Titre	1:10			

**Interpretation** Anti ds-DNA antibodies are detected more frequently and at higher titres in Systemic lupus erythematosus (SLE) pateitns with Lupus nephritis Presence of these antibodies or an increase in titre correlate with an increased risk of Lupus nephritis flare. Hence it is useful to monitor Anti ds-DNA antibody levels and initiate appropriate therapy when titres increase.

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

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