

(A Unit of KOS Healthcare)



Dr. Vinay Chopra
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Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist

NAME : Miss. JANYA CHHABRA

AGE/ GENDER : 16 YRS/FEMALE PATIENT ID : 1823698

COLLECTED BY : REG. NO./LAB NO. : 012504090007

 REFERRED BY
 : 09/Apr/2025 07:27 AM

 BARCODE NO.
 : 01528639
 COLLECTION DATE
 : 09/Apr/2025 07:35AM

 CLIENT CODE.
 : KOS DIAGNOSTIC LAB
 REPORTING DATE
 : 09/Apr/2025 04:07PM

CLIENT ADDRESS : 6349/1, NICHOLSON ROAD, AMBALA CANTT

Test Name Value Unit Biological Reference interval

HAEMATOLOGY COMPLEMENT COMPONENT - C3

COMPLEMENT COMPONENT - C3 116 mg/dL 90.0 - 180.0

bv NEPHLOMETRY

INTERPRETATION:-

C3 plays a central role in the activation of <u>complement system</u>. Its activation is required for both <u>classical</u> and <u>alternative complement activation</u> pathways. People with C3 deficiency are susceptible to bacterial infections.

Low levels indicate activation by one or both pathways.

Complement C3 levels may be useful in following the activity of immune complex diseases as most of them show decreased C3 levels.

In the classical pathway, <u>C3-convertase</u>, known as C4b2a, catalyzes the <u>proteolytic</u> cleavage of C3 into <u>C3a</u> and <u>C3b</u>. While in the alternative pathway this effect is induced by C3bBb. C3a is an<u>anaphylotoxin</u> and the precursor of some cytokines such as <u>ASP</u>, and C3b serves as an <u>opsonizing</u> agent. <u>Factor I</u> can cleave C3b into C3c and C3d, the latter of which plays a role in enhancing <u>B cell</u> responses.

Measurement of serum C3 levels are used in the assessment of children suffering from repeated severe bacterial infections and in the work up of some types of kidney disease such as <u>post-infectious glomerulonephritis</u> and <u>shunt nephritis</u>.

INCREASED IN - many inflammatory conditions as an acute-phase reactant, active phase of rheumatic diseases (eg, rheumatoid arthritis, SLE), acute viral hepatitis, myocardial infarction, cancer, diabetes mellitus, pregnancy, sarcoidosis, amyloidosis, thyroiditis.

DECREASED BY - decreased synthesis (protein malnutrition, congenital deficiency, severe liver disease), increased catabolism (immune complex disease, membranoproliferative glomerulonephritis [75%], SLE, SjAgren syndrome, rheumatoid arthritis, DIC, paroxysmal nocturnal hemoglobinuria, autoimmune hemolytic anemia, gram-negative bacteremia), increased loss (burns, gastroenteropathies).



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COMPLEMENT COMPONENT - C4

COMPLEMENT COMPONENT - C4 20.4 mg/dL 9.0 - 36.0

by NEPHLOMETRY

INTERPRETATION

C4 is a component of the classic complement pathway. Depressed levels usually indicate classic pathway activation. Low C4 accompanies acute attacks of hereditary angioedema (HAE), and C4 is used as a first-line test for the disease. C1 esterase inhibitor levels are not indicated for the evaluation of hereditary HAE unless C4 is low.

INCREASED:

1. Various malignancies (not clinically useful).

DECREASED:

- 1. Decreased synthesis (congenital deficiency),
- 2.Increased catabolism (SLE, rheumatoid arthritis, proliferative glomerulonephritis, HAE
- 3.Increased loss (burns, protein-losing enteropathies).



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Test Name Value Unit Biological Reference interval

IMMUNOPATHOLOGY/SEROLOGY ANTI dsDNA ANTIBODIES (QUANTITATIVE)

ANTI dsDNA ANTIBODIES (QUANTITATIVE)

0.48 AI

by ELISA (ENZYME-LINKED IMMUNOSORBENT ASSAY)

Negative: <0.9 Low Positive: 0.9-1.10 High Positive: >1.10

INTERPRETATION:

NOTE

- 1. Autoimmune reactivities are not by themselves diagnostic, but must be correlated with other laboratory & clinical findings **COMMENTS**
- 1. Anti ds-DNA antibodies are detected more frequently and at higher titres in Systemic lupus erythematosus (SLE) patients with Lupus nephritis.
- 2. 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.



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CLIENT ADDRESS : 6349/1, NICHOLSON ROAD, AMBALA CANTT

Value Unit Test Name **Biological Reference interval**

CLINICAL PATHOLOGY URINE ROUTINE & MICROSCOPIC EXAMINATION

PHYSICAL EXAMINATION

QUANTITY RECIEVED 10 ml by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

PALE YELLOW PALE YELLOW COLOUR

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

HAZY **CLEAR** TRANSPARANCY

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY >=1.0301.002 - 1.030

SPECIFIC GRAVITY by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

CHEMICAL EXAMINATION

ACIDIC REACTION

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

PROTEIN NEGATIVE (-ve) 3+ by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

SUGAR Negative NEGATIVE (-ve)

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

5.5 5.0 - 7.5pН

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY **BILIRUBIN** NEGATIVE (-ve) Negative

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

NITRITE Negative NEGATIVE (-ve) by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY.

UROBILINOGEN Normal EU/dL 0.2 - 1.0by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

KETONE BODIES Negative NEGATIVE (-ve)

by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY **BLOOD** NEGATIVE (-ve)

Negative by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

ASCORBIC ACID NEGATIVE (-ve) NEGATIVE (-ve) by DIP STICK/REFLECTANCE SPECTROPHOTOMETRY

MICROSCOPIC EXAMINATION



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Test Name	Value	Unit	Biological Reference interval
RED BLOOD CELLS (RBCs) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)	/HPF	0 - 3
PUS CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	1-2	/HPF	0 - 5
EPITHELIAL CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	3-4	/HPF	ABSENT
CRYSTALS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)
CASTS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)
BACTERIA by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)
OTHERS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	NEGATIVE (-ve)		NEGATIVE (-ve)
TRICHOMONAS VAGINALIS (PROTOZOA) by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT	ABSENT		ABSENT
RECHECKED			

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Test Name Value Unit Biological Reference interval

SPECIAL INVESTIGATIONS

ANTI NUCLEAR ANTIBODY/FACTOR (ANA/ANF) - WITH REFLEX TO TITRES: IFA (HEP-2)

ANTI NUCLEAR ANTIBODY (ANA) - IFA, HEp2

NEGATIVE (-ve)

NEGATIVE (-ve)

by IFA (IMMUNO FLUORESCENT ASSAY)

INTERPRETATION:

- 1. Immunofluorescence microscopy using human cellular extracts like Hep-2 cells is sensitive for detection of serum antibodies that react specifically with various cellular proteins and nucleic acid.
- 2. Test conducted on serum

3. Patients are reported as per international consensus ANA Patterns (ICAP)

INTERNATIONAL GUIDELINES FOR GRADING			
GRADE	REMARKS		
Negative (-ve)	No fluorescence		
<u>1+</u>	Minimum fluorescence		
<u>2+</u>	Mildly positive		
<u>3+</u>	Significantly positive		
4+	Strongly positive		

COMMENTS:

Anti Nuclear antibody (ANA / ANF) is a group of autoantibodies directed against constituents of cell nuclei including DNA, RNA & various nuclear proteins. These autoantibodies are found with high frequency in patients with connective tissue disorders specially SLE. Since positive ANA results have been reported in healthy individuals, these reactivities are not by themselves diagnostic but must be correlated with other laboratory and clinical findings.

PATTERN (ICAP)	ICAP CODE	ANTIGEN ASSOCIATION	DISEASE ASSOCIATION			
NUCLEAR PATTERNS						
Homogenous	AC-1	dsDNA, nucleosomes, histones	SLE, Drug-induced lupus, Juvenile idiopathic			



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Γest Name		Value	Unit B	iological Reference interval
			arthritis	7
Speckled	AC-2,4,5	hnRNP, U1RNP, Sm, SS-A/Ro (Ro 80), SS-B/La, RNA polymerase III, Mi-2, Ku	MCTD, SLE, DM, SSc/PN overlap	Л
Dense fine speckled	AC-2	DFS70/LEDGF	Rare in SLE, Sjogren's syndrome, SSc	
Fine speckled	AC-4	SS-A/Ro (Ro 80), T1F1ß, SS-B/La,Mi-2,T1F1γ, Ku, RNA helicase A, replication protein A	Sjogren's syndrome, SL DM,SSc/PM overlap	E,
Large/Coarse speckled	AC-5	hnRNP, U1RNP, Sm, RNA polymerase III	MCTD, SLE, SSc	
Centromere	AC-3	CENP-A/B	Limited cutaneous SSc PBC	,
Discrete nuclear dots	AC-6,7			
Multiple nuclear dots	Ac-6	Sp-100, PML proteins, MJ/NXP-2	PBC, SARD, PM/DM	
Few nuclear dots	Ac-7	P80-coilin, SMN	Sjogren's syndrome, SL SSc, PM, asymptomati individuals	
Nucleolar	AC-8,9,10			
Nucleolar homogenous	AC-8	PM/ScI-75, PM/ScI-100, Thi/To,B23/nucleophosmin, nucleolin, No55/SC65	SSc, SSc/PM overlap	
Nucleolar clumpy	AC-9	U3-smoRNP/fibrillarin	SSc	
Nucleolar punctate	Ac-10	RNA polymerase 1, hUBF/NOR-90	SSc, Sjogren's syndrom	е
Nuclear envelope	AC-11,12			
Smooth nuclear envelope	AC-11	Lamin A,B,C or lamin associated proteins	SLE, Sjogren's syndrom Seronegative arthritis	
			T	



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Test Name		Value	Unit	Biological Reference interval
Punctate nuclear envelope	AC-12	Nuclear pore complex proteins (gp210)	PBC	
Pleomorphic	AC-13,14			
PCNA-like	AC-13	PCNA	SLE, other conditi	ons
CENP-F like	AC-14	CENP-F	Cancer, other cond	itions
	СҮТС	OPLASMIC PATTERNS		
Fibrillar	AC-15,16,17			
Linear/actin	AC-15	Actin, non-muscle myosin, MCTD	MCTD, Chronic ac hepatitis, Liver cirrh Myasthenia gravis, Crohn's dise PBC, Long term hemodialysis, rar SARD other than M	nosis, ease, e in
Filamentous/microtubules	AC-16	Vimentin, cytokeratins	Infections or inflammatory condi Long term hemodia Alcoholic liver dise SARD, Psoriasis, he controls	llysis, ease,
Segmental	AC-17	Alpha-actin, vinuculin, tropomyosin	Myasthenia grav Crohn's disease Ulcerative colit	2,
Speckled	AC-18,19,20			
Discrete dots/GW body like	AC-18	SGW182, Su/Ago2,	PBC, SARD, neurolo and autoimmun conditions	ngical e
Dense fine speckled	AC-19	PL-7, PL-12, ribosomal P proteins	Anti-synthetase syndrome, PM/DM, Juvenile SLE, Neuropsychiatric	SLE,



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Test Name		Value	Unit Biolo	ogical Reference interval
Fine speckled	AC-20	Jo-1/histidyl-Trna synthetase	Anti-synthetase syndrome, PM/DM, limited SSc, Idiopathic pleural effusion	
Reticular/AMA (Mitochondrial)	AC-21	PDC-E2/M2, BCOADC-E2 OGDC- E2, E1a subunit of PDC, E3BP/proteinX	Common in PBC, SSc, rare in other SARD	
Polar/ Golgi like	AC-22	Giantin/macrogolgin, golgin-97, golgin-245	Rare in Sjogren's syndrome, SLE, RA, MCTD,GPA, Idiopathic cerebellar ataxia, Paraneoplastic cerebellar degeneration,viral infections	
Rods and rings	AC-23	IMPDH2, others	HCV patients post IFN/Ribavirin therapy,rare in SLE, Hashimoto's and healthy controls	
		MITOTIC PATTERNS		
Centrosome	AC-24	Pericentrin, ninein, Cep250, Cep110	Rare in SSc, Raynaud's phenomenon, infections (viral and mycoplasma)	
Spindle fibres	AC-25	HsEg5	Rare in Sjogren's syndrome, SLE, other SARD	
NuMA like	AC-26	Centrophilin	Sjogren's syndrome, SLE, other	
Intracellular bridges	AC-27	Aurora kinase B, CENP-E,MSA-2, KIF-14, MKLP-1	Rare in SSc, Raynaud's phenomenon, malignancy	
	1		1	1



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Test Name		Value	Unit Bio	logical Reference interval
Mitotic chromosome coat	AC-28	Modified histone H3, MCA-1	Rare in Discoid lupus	
			erythematous, Chronic	
		· ·	lymphocytic leukemia,	
			Sjogren's syndrome, and	
			Polymyalgia rheumatica	

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End Of Report



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