



| | Dr. Vinay Chopra | | Dr. Yugan | |
|--|--|-------------------|--|--------------------------------------|
| | MD (Pathology & Micr Chairman & Consultan | | | (Pathology) t Pathologist |
| NAME : | Mr. ABHISHEK MALIK | | | |
| AGE/ GENDER : | 41 YRS/MALE | | PATIENT ID | : 1749590 |
| COLLECTED BY : | SURJESH | | REG. NO./LAB NO. | : 012502080019 |
| REFERRED BY : | | | REGISTRATION DATE | : 08/Feb/2025 10:21 AM |
| | 01525139 | | COLLECTION DATE | : 08/Feb/2025 10:24AM |
| | KOS DIAGNOSTIC LAB | | REPORTING DATE | : 08/Feb/2025 11:27AM |
| CLIENT ADDRESS : | 6349/1, NICHOLSON ROAD, AMB/ | ALA CANTT | | |
| Test Name | | Value | Unit | Biological Reference interval |
| | SWASTI | HYA WEI | LLNESS PANEL: 1.0 | n |
| | | | DOD COUNT (CBC) | |
| RED BLOOD CELLS (J | RBCS) COUNT AND INDICES | | · · | |
| HAEMOGLOBIN (HB) | | 15.3 | gm/dL | 12.0 - 17.0 |
| by CALORIMETRIC RED BLOOD CELL (RB | C) COUNT | 5.12 ^H | Millions | /cmm 3.50 - 5.00 |
| by HYDRO DYNAMIC FOC | USING, ELECTRICAL IMPEDENCE | | | |
| PACKED CELL VOLUM by CALCULATED BY AUT | E (PCV) OMATED HEMATOLOGY ANALYZER | 45.4 | % | 40.0 - 54.0 |
| MEAN CORPUSCULAR | | 88.6 | fL | 80.0 - 100.0 |
| • | OMATED HEMATOLOGY ANALYZER 2. HAEMOGLOBIN (MCH) | 29.8 | pg | 27.0 - 34.0 |
| - | OMATED HEMATOLOGY ANALYZER | 007 | | 0.00 |
| | CHEMOGLOBIN CONC. (MCHC) | 33.7 | g/dL | 32.0 - 36.0 |
| | ION WIDTH (RDW-CV) omated hematology analyzer | 15.2 | % | 11.00 - 16.00 |
| RED CELL DISTRIBUT | ION WIDTH (RDW-SD) | 50.7 | fL | 35.0 - 56.0 |
| by CALCULATED BY AUT | OMATED HEMATOLOGY ANALYZER | 17.3 | RATIO | BETA THALASSEMIA TRAIT: < |
| by CALCULATED | | 17.0 | in the second se | 13.0 |
| | | | | IRON DEFICIENCY ANEMIA: >13.0 |
| GREEN & KING INDEX | K | 26.23 | RATIO | BETA THALASSEMIA TRAIT:< |
| by CALCULATED | | | | 65.0 |
| | | | | IRON DEFICIENCY ANEMIA: > 65.0 |
| WHITE BLOOD CELLS | <u>S (WBCS)</u> | | | |
| TOTAL LEUCOCYTE C | OUNT (TLC) y sf cube & microscopy | 8430 | /cmm | 4000 - 11000 |
| NUCLEATED RED BLC | OOD CELLS (nRBCS) | NIL | | 0.00 - 20.00 |
| by AUTOMATED 6 PART F NUCLEATED RED BLC | HEMATOLOGY ANALYZER | NIL | % | < 10 % |
| | OD CELLS (IRBCS) % OMATED HEMATOLOGY ANALYZER | INIL | 70 | < 1 U 70 |
| | | | | |





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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.







Dr. Vinay Chopra Dr. Yugam Chopra MD (Pathology & Microbiology) MD (Pathology) Chairman & Consultant Pathologist **CEO & Consultant Pathologist** NAME : Mr. ABHISHEK MALIK AGE/ GENDER : 41 YRS/MALE **PATIENT ID** :1749590 **COLLECTED BY** : SURJESH :012502080019 REG. NO./LAB NO. **REFERRED BY REGISTRATION DATE** :08/Feb/2025 10:21 AM : **BARCODE NO. COLLECTION DATE** :08/Feb/2025 10:24AM :01525139 CLIENT CODE. : KOS DIAGNOSTIC LAB **REPORTING DATE** :08/Feb/202511:27AM **CLIENT ADDRESS** : 6349/1, NICHOLSON ROAD, AMBALA CANTT Value Unit **Biological Reference interval** Test Name CORRECTED TOTAL LEUCOCYTE COUNT (C-TLC) 8430 4000 - 11000 /cmm by MICROSCOPY ON EDTA SMEAR **DIFFERENTIAL LEUCOCYTE COUNT (DLC) NEUTROPHILS** 57 % 50 - 70 by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY LYMPHOCYTES 26% 20 - 40 by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY 12^H EOSINOPHILS % 1 - 6by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY 5 MONOCYTES % 2 - 12by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY BASOPHILS 0 % 0 - 1 by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY **ABSOLUTE LEUKOCYTES (WBC) COUNT** ABSOLUTE NEUTROPHIL COUNT 4805 2000 - 7500 /cmm by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY ABSOLUTE LYMPHOCYTE COUNT 800 - 4900 2192 /cmm by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY ABSOLUTE EOSINOPHIL COUNT 1012^H 40 - 440 /cmm by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY ABSOLUTE MONOCYTE COUNT 422/cmm 80 - 880 by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY PLATELETS AND OTHER PLATELET PREDICTIVE MARKERS. PLATELET COUNT (PLT) 150000 - 450000 221000 /cmm by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE PLATELETCRIT (PCT) 0.29 % 0.10 - 0.36 by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE MEAN PLATELET VOLUME (MPV) 13^H fL 6.50 - 12.0 by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE 30000 - 90000 PLATELET LARGE CELL COUNT (P-LCC) /cmm 106000^H by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE 48^H % PLATELET LARGE CELL RATIO (P-LCR) 11.0 - 45.0 by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE PLATELET DISTRIBUTION WIDTH (PDW) 15.0 - 17.0 16.2%

by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD



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| Test Name | Value | Unit | Biological Reference interval | | | |





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| CLIENT CODE. | : KOS DIAGN | OSTIC LAB | REI | PORTING DATE | :08/Feb/202512:45PM | |
| CLIENT ADDRESS | : 6349/1, NIC | HOLSON ROAD, A | AMBALA CANTT | | | |
| Test Name | | | Value | Unit | Biological Reference interval | |
| ERYTHROCYTE SE by red cell aggre INTERPRETATION: | | RATE (ESR) | 11 | NTATION RATE (ES mm/1st h | | |





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| | | | & Microbiology) nsultant Pathologis | | (Pathology) Pathologist | |
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| CLIENT CODE. | : KOS DIAGNOS | TIC LAB | | REPORTING DATE | : 08/Feb/2025 12:08PM | |
| CLIENT ADDRESS | : 6349/1, NICH | OLSON ROAD, | AMBALA CANTT | | | |
| Test Name | | | Value | Unit | Biological Reference interval | |
| | | CLINI | CAL CHEMIS | TRY/BIOCHEMIST | 'RY | |
| | | | GLUCOSE | FASTING (F) | | |
| GLUCOSE FASTIN | G (F): PLASMA Se - peroxidase (G | DD-POD) | 108.56 ^H | mg/dL | NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.0 | |

IN ACCORDANCE WITH AMERICAN DIABETES ASSOCIATION GUIDELINES:

A fasting plasma glucose level below 100 mg/dl is considered normal.
 A fasting plasma glucose level between 100 - 125 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients.

test (after consumption of 75 gms of glucose) is recommended for all such patients. 3. A fasting plasma glucose level of above 125 mg/dl is highly suggestive of diabetic state. A repeat post-prandial is strongly recommended for all such patients. A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.





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| | | | | |
| Fest Name | | Value | Unit | Biological Reference interval |
| | | LIPID PROFI | LE · BASIC | |
| HOLESTEROL TOTAL | : SERUM | 213.95 ^H | mg/dL | OPTIMAL: < 200.0 |
| by CHOLESTEROL OXIDA | | 213.95" | ilig/ uL | BORDERLINE HIGH: 200.0 - |
| | | | | 239.0 |
| | | | | HIGH CHOLESTEROL: > OR = 240.0 |
| RIGLYCERIDES: SERU | | 151.67 ^H | mg/dL | OPTIMAL: < 150.0 |
| by GLYCEROL PHOSPHAT | E OXIDASE (ENZYMATIC) | | | BORDERLINE HIGH: 150.0 - 199.0 |
| | | | | HIGH: 200.0 - 499.0 |
| | | | | VERY HIGH: > OR = 500.0 |
| IDL CHOLESTEROL (I by SELECTIVE INHIBITION | DIRECT): SERUM | 40.68 | mg/dL | LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 |
| | | | | 60.0 |
| | | | . / 11 | HIGH HDL: $> OR = 60.0$ |
| DL CHOLESTEROL: S by CALCULATED, SPECTR | | 142.94 ^H | mg/dL | OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129. |
| | | | | BORDERLINE HIGH: 130.0 - |
| | | | | 159.0 HIGH: 160.0 - 189.0 |
| | | | | VERY HIGH: > OR = 190.0 |
| ION HDL CHOLESTER | | 173.27 ^H | mg/dL | OPTIMAL: < 130.0 |
| by CALCULATED, SPECTR | OPHOTOMETRY | | | ABOVE OPTIMAL: 130.0 - 159. BORDERLINE HIGH: 160.0 - |
| | | | | 189.0 |
| | | | | HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0 |
| LDL CHOLESTEROL: | SERUM | 30.33 | mg/dL | 0.00 - 45.00 |
| by CALCULATED, SPECTR | | 579.57 | mg/dL | 350.00 - 700.00 |
| by CALCULATED, SPECTR | | | | |
| HOLESTEROL/HDL R | | 5.26 ^H | RATIO | LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 |
| S, ONLOOLATED, OF LOTA | | | | AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 |
| | | | | HIGH RISK: > 11.0 |
| | | | | |

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| Test Name | | Value | Unit | Biological Reference interval | | |
| LDL/HDL RATIO: S by CALCULATED, SPE | | 3.51 ^H | RATIO | LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0 | | |
| TRIGLYCERIDES/H by CALCULATED, SPE | IDL RATIO: SERUM | 3.73 | RATIO | 3.00 - 5.00 | | |

INTERPRETATION:

1. Measurements in the same patient can show physiological analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.

2. As per NLA-2014 guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.

 Low HDL levels are associated with increased risk for Atherosclerotic Cardiovascular disease (ASCVD) due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.
 NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogeniclipoproteins such as LDL, VLDL, IDL, Lpa, Chylomicron remnants) along with LDL-cholesterol as co- primary target for cholesterol lowering therapy. Note that major risk factors can modify treatment goals for LDL & Non HDL

5. Additional testing for Apolipoprotein B, hsCRP,Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement





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| | | | | |
| Test Name | | Value | Unit | Biological Reference interval |
| BILIRUBIN TOTAI | | FUNCTIO 0.65 | N TEST (COMPLETE) mg/dL | INFANT: 0.20 - 8.00 |
| | SPECTROPHOTOMETRY | 0.03 | liig/ uL | ADULT: 0.00 - 1.20 |
| | T (CONJUGATED): SERUM SPECTROPHOTOMETRY | 0.15 | mg/dL | 0.00 - 0.40 |
| | ECT (UNCONJUGATED): SERUM | 0.5 | mg/dL | 0.10 - 1.00 |
| SGOT/AST: SERUN by IFCC, WITHOUT P | Л YRIDOXAL PHOSPHATE | 20.95 | U/L | 7.00 - 45.00 |
| SGPT/ALT: SERUN by IFCC, WITHOUT P | Л YRIDOXAL PHOSPHATE | 25.54 | U/L | 0.00 - 49.00 |
| AST/ALT RATIO: S | SERUM ectrophotometry | 0.82 | RATIO | 0.00 - 46.00 |
| ALKALINE PHOSP by para nitrophei propanol | HATASE: SERUM NYL PHOSPHATASE BY AMINO METHYL | 103.47 | U/L | 40.0 - 130.0 |
| GAMMA GLUTAMY by SZASZ, SPECTRO | YL TRANSFERASE (GGT): SERUM | 48.92 | U/L | 0.00 - 55.0 |
| TOTAL PROTEINS | | 7.71 | gm/dL | 6.20 - 8.00 |
| ALBUMIN: SERUM | I | 4.26 | gm/dL | 3.50 - 5.50 |
| GLOBULIN: SERUI | | 3.45 | gm/dL | 2.30 - 3.50 |
| A : G RATIO: SERU | | 1.23 | RATIO | 1.00 - 2.00 |

by CALCULATED, SPECTROPHOTOMETRY

INTERPRETATION

NOTE:- To be correlated in individuals having SGOT and SGPT values higher than Normal Referance Range.

USE:- Differential diagnosis of diseases of hepatobiliary system and pancreas.

INCREASED:

| DRUG HEPATOTOXICITY | > 2 |
|--|----------------------------|
| ALCOHOLIC HEPATITIS | > 2 (Highly Suggestive) |
| CIRRHOSIS | 1.4 - 2.0 |
| INTRAHEPATIC CHOLESTATIS | > 1.5 |
| HEPATOCELLULAR CARCINOMA & CHRONIC HEPATITIS | > 1.3 (Slightly Increased) |



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DECREASED:

1. Acute Hepatitis due to virus, drugs, toxins (with AST increased 3 to 10 times upper limit of normal)

2. Extra Hepatic cholestatis: 0.8 (normal or slightly decreased).

| NORMAL | < 0.65 |
|----------------------|-----------|
| GOOD PROGNOSTIC SIGN | 0.3 - 0.6 |
| POOR PROGNOSTIC SIGN | 1.2 - 1.6 |



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| Test Name | | Value | Unit | Biological Reference interval |
| | KIDNE | Y FUNCTION 7 | FEST (COMPLETE) | |
| UREA: SERUM | IATE DEHYDROGENASE (GLDH) | 22.16 | mg/dL | 10.00 - 50.00 |
| CREATININE: SERU | JM | 1.12 | mg/dL | 0.40 - 1.40 |
| - | OGEN (BUN): SERUM | 10.36 | mg/dL | 7.0 - 25.0 |
| BLOOD UREA NITE RATIO: SERUM | OGEN (BUN)/CREATININE | 9.25 ^L | RATIO | 10.0 - 20.0 |
| by CALCULATED, SPE UREA/CREATININ by CALCULATED, SPE | E RATIO: SERUM | 19.79 | RATIO | |
| URIC ACID: SERUM by URICASE - OXIDAS | | 7.58 | mg/dL | 3.60 - 7.70 |
| CALCIUM: SERUM by ARSENAZO III, SPE | | 9.78 | mg/dL | 8.50 - 10.60 |
| PHOSPHOROUS: SE | | 3.86 | mg/dL | 2.30 - 4.70 |
| SODIUM: SERUM by ISE (ION SELECTIV | 'E ELECTRODE) | 139.55 | mmol/L | 135.0 - 150.0 |
| POTASSIUM: SERUE by ISE (ION SELECTIV | M | 4.18 | mmol/L | 3.50 - 5.00 |
| CHLORIDE: SERUM | ſ | 104.66 | mmol/L | 90.0 - 110.0 |
| | IERULAR FILTERATION RATE | | | |
| ESTIMATED GLOM (eGFR): SERUM by CALCULATED INTERPRETATION: | ERULAR FILTERATION RATE | 84.6 | | |

To differentiate between pre- and post renal azotemia.

INCREASED RATIO (>20:1) WITH NORMAL CREATININE:

1. Prerenal azotemia (BUN rises without increase in creatinine) e.g. heart failure, salt depletion, dehydration, blood loss) due to decreased glomerular filtration rate.

2. Catabolic states with increased tissue breakdown.

3. GI haemorrhage.



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|---|---|--|---|---|---------------------------|--|-------------|-------------|
| IAME | : Mr. ABHISHE | K MALIK | | | | | | |
| AGE/ GENDER | : 41 YRS/MALE | | PA | TIENT ID | : 17 | 49590 | | |
| COLLECTED BY | : SURJESH | | RE | G. NO./LAB NO. | :01 | 250208001 | 9 | |
| REFERRED BY | | | RE | GISTRATION DA | ATE • 08 | /Feb/2025 10 | 0·21 AM | |
| BARCODE NO. | : 01525139 | | | LLECTION DAT | | /Feb/2025 10 | | |
| CLIENT CODE. | : KOS DIAGNOS | | | PORTING DATE | | /Feb/2025 1 | | |
| CLIENT ADDRESS | | OLSON ROAD, AMBAL | | I OKING DAII | | / FED/ 2023 1 | 1.55AW | |
| Test Name | | v | alue | Uni | it | Biologi | ical Refere | nce interva |
| INCREASED RĂTIO (>2 1. Postrenal azotemia 2. Prerenal azotemia DECREASED RATIO (< | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises disprosed on superimposed on 10:1) WITH DECRE | TED CREATININE LEVELS oportionately more that renal disease. | | (e.g. obstructive | europathy). | | | |
| NCREASED RATIO (>2 1. Postrenal azotemia 2. Prerenal azotemia DECREASED RATIO (< 1. Acute tubular necr 2. Low protein diet an 3. Severe liver diseas 4. Other causes of de 5. Repeated dialysis (6. Inherited hyperam 7. SIADH (syndrome of 8. Pregnancy. DECREASED RATIO (< 8. Phenacimide thera 2. Rhabdomyolysis (r 8. Muscular patients NAPPROPIATE RATIO 1. Diabetic ketoacido should produce an in 2. Cephalosporin thera 5. CKD STAGE G1 | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises dispro- superimposed on 10:1) WITH DECRE , osis. ad starvation. e. creased urea synt urea rather than monemias (urea in the synthesis of the synthesynthesis of the s | ocorticoids) FED CREATININE LEVELS oportionately more that is renal disease. ASED BUN : thesis. creatinine diffuses out is virtually absent in blutidiuretic harmone) du ASED CREATININE: proversion of creatine to reatinine). al failure. causes false increase i atinine ratio). ith creatinine measure: RATE: DESCRIPTION hal kidney function | in creatinine) c of extracellu ood). le to tubular s o creatinine). n creatinine w ment). GFR (mL/r | lar fluid). secretion of urea with certain meth nin/1.73m2) | hodologies,re ASSOCIAT | ED FINDINGS oteinuria | | hen dehydra |
| NCREASED RATIO (>2 Postrenal azotemia Perenal azotemia DECREASED RATIO (< Acute tubular necr Low protein diet and Severe liver diseas Other causes of de Repeated dialysis Nherited hyperam SIADH (syndrome of Pregnancy. DECREASED RATIO (< Phenacimide thera Rhabdomyolysis (r Diabetic ketoacido hould produce an in Cephalosporin ther STIMATED GLOMERI CKD STAGE | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises dispro- superimposed on 10:1) WITH DECRE , osis. ad starvation. e. creased urea synt urea rather than monemias (urea i finappropiate ar 10:1) WITH INCRE , py (accelerates co eleases muscle cr who develop rena- : sis (acetoacetate creased BUN/crea- apy (interferes w <u>JLAR FILTERATION</u> <u>Norn</u> Kid | ocorticoids) FED CREATININE LEVELS oportionately more that is renal disease. ASED BUN : thesis. creatinine diffuses out is virtually absent in black tidiuretic harmone) du ASED CREATININE: onversion of creatine to reatinine). al failure. causes false increase i atinine ratio). ith creatinine measure: RATE: DESCRIPTION nal kidney function ney damage with | in creatinine) c of extracellu ood). le to tubular s o creatinine). n creatinine w ment). GFR (mL/r | lar fluid). secretion of urea with certain meth | hodologies,re ASSOCIAT | ED FINDINGS oteinuria of Protein , | | hen dehydra |
| NCREASED RATIO (>2 Postrenal azotemia Perenal azotemia DECREASED RATIO (< Acute tubular necr Low protein diet an Severe liver diseas Other causes of de Repeated dialysis Neperated dialysis Pregnancy. DECREASED RATIO (< Phenacimide thera Rhabdomyolysis (r Anuscular patients NAPPROPIATE RATIO Diabetic ketoacido hould produce an in CED STAGE STIMATED GLOMERL CKD STAGE G1 | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises dispro- superimposed on 10:1) WITH DECRE , osis. ad starvation. e. creased urea synt urea rather than monemias (urea in the synthesis of the synthesynthesis of the s | ocorticoids) FED CREATININE LEVELS oportionately more that is renal disease. ASED BUN : thesis. creatinine diffuses out is virtually absent in blutidiuretic harmone) du ASED CREATININE: proversion of creatine to reatinine). al failure. causes false increase i atinine ratio). ith creatinine measure: RATE: DESCRIPTION hal kidney function | in creatinine) c of extracellu ood). le to tubular s o creatinine). ment). GFR (mL/r | lar fluid). secretion of urea with certain meth nin/1.73m2) •90 •90 | hodologies,re ASSOCIAT | ED FINDINGS oteinuria | | hen dehydra |
| INCREASED RATIO (>2 1. Postrenal azotemia 2. Prerenal azotemia DECREASED RATIO (< 1. Acute tubular necr 2. Low protein diet an 3. Severe liver diseas 4. Other causes of de 5. Repeated dialysis (6. Inherited hyperam 7. SIADH (syndrome of 8. Pregnancy. DECREASED RATIO (< 1. Phenacimide thera 2. Rhabdomyolysis (r 3. Muscular patients INAPPROPIATE RATIO 1. Diabetic ketoacido should produce an in 2. Cephalosporin there ESTIMATED GLOMERI G1 G2 G3a G3a G3b | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises dispro- superimposed on 10:1) WITH DECRE osis. ad starvation. e. creased urea synt urea rather than monemias (urea i of inappropiate ar 10:1) WITH INCRE py (accelerates co eleases muscle cr who develop rena- : sis (acetoacetate creased BUN/crea- apy (interferes w <u>UAR FILTERATION</u> Norn Kid noi | ocorticoids) FED CREATININE LEVELS oportionately more that is renal disease. ASED BUN : thesis. creatinine diffuses out is virtually absent in blutidiuretic harmone) du ASED CREATININE: onversion of creatine to reatinine). al failure. causes false increase i atinine ratio). ith creatinine measure: RATE: DESCRIPTION nal kidney function ney damage with rmal or high GFR d decrease in GFR rate decrease in GFR | in creatinine) c of extracellu ood). le to tubular s o creatinine). ment). GFR (mL/r c 60 31 | lar fluid). secretion of urea with certain meth nin/1.73m2) .90 .90 .90 .59 | hodologies,re ASSOCIAT | ED FINDINGS oteinuria of Protein , | | hen dehydra |
| 1. Postrenal azotemia 2. Prerenal azotemia DECREASED RATIO (< 1. Acute tubular necr 2. Low protein diet ar 3. Severe liver diseas 4. Other causes of de 5. Repeated dialysis (6. Inherited hyperam 7. SIADH (syndrome of 8. Pregnancy. DECREASED RATIO (< 1. Phenacimide thera 2. Rhabdomyolysis (r 3. Muscular patients INAPPROPIATE RATIO 1. Diabetic ketoacido should produce an in 2. Cephalosporin ther ESTIMATED GLOMERI G1 G2 G3a | tetracycline, gluc 0:1) WITH ELEVAT (BUN rises dispro- superimposed on 10:1) WITH DECRE osis. ad starvation. e. creased urea synt urea rather than monemias (urea i of inappropiate ar 10:1) WITH INCRE py (accelerates co eleases muscle cr who develop rena- : sis (acetoacetate creased BUN/crea- apy (interferes w <u>UAR FILTERATION</u> Norn Kid noi- Norn Kid Noder | ocorticoids) FED CREATININE LEVELS oportionately more that is renal disease. ASED BUN : thesis. creatinine diffuses out is virtually absent in bl itidiuretic harmone) du ASED CREATININE: onversion of creatine to reatinine). al failure. causes false increase i atinine ratio). ith creatinine measure: RATE: DESCRIPTION nal kidney function ney damage with rmal or high GFR d decrease in GFR | in creatinine) c of extracellu ood). le to tubular s o creatinine). ment). GFR (mL/r 60 30 41 | lar fluid). secretion of urea with certain meth nin/1.73m2) •90 •90 | hodologies,re ASSOCIAT | ED FINDINGS oteinuria of Protein , | | hen dehydra |





DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY)







| | Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Patholog | | (Pathology) |
|---------------------|--|--------------------------|-------------------------------|
| NAME | : Mr. ABHISHEK MALIK | | |
| AGE/ GENDER | : 41 YRS/MALE | PATIENT ID | : 1749590 |
| COLLECTED BY | : SURJESH | REG. NO./LAB NO. | : 012502080019 |
| REFERRED BY | : | REGISTRATION DATE | : 08/Feb/2025 10:21 AM |
| BARCODE NO. | : 01525139 | COLLECTION DATE | : 08/Feb/2025 10:24AM |
| CLIENT CODE. | : KOS DIAGNOSTIC LAB | REPORTING DATE | : 08/Feb/2025 11:55AM |
| CLIENT ADDRESS | : 6349/1, NICHOLSON ROAD, AMBALA CANT | Т | |
| Test Name | Value | Unit | Biological Reference interval |

COMMENTS:

Estimated Glomerular filtration rate (eGFR) is the sum of filtration rates in all functioning nephrons and so an estimation of the GFR provides a measure of functioning nephrons of the kidney.
 eGFR calculated using the 2009 CKD-EPI creatinine equation and GFR category reported as per KDIGO guideline 2012
 In patients, with eGFR creatinine between 45-59 ml/min/1.73 m2 (G3) and without any marker of Kidney damage, It is recommended to measure of CFD with the commended to measure

3. In patients, with eGFR cleaning between 45-59 minimit 1.73 m2 (G3) and without any marker of Kidney damage, it is recommended to measure eGFR with Cystatin C for confirmation of CKD
4. eGFR category G1 OR G2 does not fulfill the criteria for CKD, in the absence of evidence of Kidney Damage
5. In a suspected case of Acute Kidney Injury (AKI), measurement of eGFR should be done after 48-96 hours of any Intervention or procedure
6. eGFR calculated by Serum Creatinine may be less accurate due to certain factors like Race, Muscle Mass, Diet, Certain Drugs. In such cases, eGFR should be calculated using Serum Cystatin C
7. A decrease in eGFR implies either progressive renal disease, or a reversible process causing decreased nephron function (eg, severe dehydration).

ADVICE:

KDIGO guideline, 2012 recommends Chronic Kidney Disease (CKD) should be classified based on cause, eGFR category and Albuminuria (ACR) category. GFR & ACR category combined together reflect risk of progression and helps Clinician to identify the individual who are progressing at more rapid rate than anticipated



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| | | hopra & Microbiology) nsultant Pathologist | Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist | | |
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| BARCODE NO. | :01525139 | | LLECTION DATE | : 08/Feb/2025 10:24AM | |
| CLIENT CODE. CLIENT ADDRESS | : KOS DIAGNOSTIC LAB : 6349/1, NICHOLSON ROAD | | PORTING DATE | : 08/Feb/2025 10:42AM | |
| CLIENT ADDRESS | . 0349/1, MCHOLSON ROAD | , AMDALA CAN I I | | | |
| Test Name | | Value | Unit | Biological Reference interval | |
| | | CLINICAL PA | THOLOGY | | |
| | URINE R | | SCOPIC EXAMINA | ATION | |
| PHYSICAL EXAMI | | | | | |
| QUANTITY RECIEV | | 10 | ml | | |
| | TANCE SPECTROPHOTOMETRY | | | | |
| COLOUR by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | PALE YELLO | W | PALE YELLOW | |
| | TANCE SPECTROPHOTOMETRY | CLEAR | | CLEAR | |
| SPECIFIC GRAVITY | 7 | 1.02 | | 1.002 - 1.030 | |
| | TANCE SPECTROPHOTOMETRY | | | | |
| <u>CHEMICAL EXAM</u> REACTION | INATION | ACIDIC | | | |
| | TANCE SPECTROPHOTOMETRY | | | | |
| PROTEIN by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | Negative | | NEGATIVE (-ve) | |
| SUGAR | | Negative | | NEGATIVE (-ve) | |
| by DIP STICK/REFLEC | CTANCE SPECTROPHOTOMETRY | <=5.0 | | 5.0 - 7.5 | |
| by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | | | | |
| BILIRUBIN by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | Negative | | NEGATIVE (-ve) | |
| NITRITE | TANCE SPECTROPHOTOMETRY. | Negative | | NEGATIVE (-ve) | |
| UROBILINOGEN | | Normal | EU/dL | 0.2 - 1.0 | |
| by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | Negative | | NEGATIVE (-ve) | |
| by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | | | | |
| BLOOD by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY | Negative | | NEGATIVE (-ve) | |
| ASCORBIC ACID | | NEGATIVE (- | ve) | NEGATIVE (-ve) | |
| by DIP STICK/REFLEC | TANCE SPECTROPHOTOMETRY AMINATION | | | | |
| RED BLOOD CELLS | | NEGATIVE (- | ve) /HPF | 0 - 3 | |

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Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist

HEALTHCARE & DIAGNOSTIC EXCELLENCE IN

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

| NAME | : Mr. ABHISHEK MALIK | | | |
|--------------------|------------------------------|-------------|--------------------------|-------------------------------|
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| CLIENT ADDRESS | : 6349/1, NICHOLSON ROAD, | AMBALA CANT | Г | |
| Test Name | | Value | Unit | Biological Reference interval |
| by MICROSCOPY ON | CENTRIFUGED URINARY SEDIMENT | | | |
| PUS CELLS | | 2-3 | /HPF | 0 - 5 |

| by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT | | | | |
|---|----------------|------|----------------|--|
| EPITHELIAL CELLS by MICROSCOPY ON CENTRIFUGED URINARY SEDIMENT | 1-2 | /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 | |

End Of Report



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