DIAGNOSTICS REPORT

Patient Name : Mr. ROHIT BHAGWAN PARAB

Age/Sex : 32 Year(s)/Male : SHHM 94020

UHID : SHHM.94020 Ref. Doctor : Self

Address : PARAB CHAL, RAMESH WAR SEVA

MANDAL, SANTACRUZ E,Mumbai,

Maharastra, 400055

Facility

Order Date

Report Date

: SEVENHILLS HOSPITAL,

: 08/05/2024 09:28

: 10/05/2024 09:58

MUMBAI

Mobile : 9702628411

X-RAY CHEST PA VIEW

Both lungs are clear.

The frontal cardiac dimensions are normal.

The pleural spaces are clear.

Both hilar shadows are normal in position and density.

No diaphragmatic abnormality is seen.

The soft tissues and bony thorax are normal.

IMPRESSION: No pleuroparenchymal lesion is seen.

Dr.Bhavesh Rajesh Dubey MBBS,MD

RegNo: 2017/03/0656

Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex

Episode

: 32 Year(s) / Male

UHID : SHHM.94020 **Order Date** : 08/05/2024 09:28 : OP

Mobile No Ref. Doctor : Self : 9702628411

2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

DOB : 11/01/1992

Facility : SEVENHILLS HOSPITAL, MUMBAI

Biochemistry

| Test Name | | Resu | ılt | Unit | Bio | logical Reference Interval |
|-------------------------------------|----------------------|-------------|------------|------------------|---------------|----------------------------|
| Sample No: 0033055 | 5C Collection Date : | 08/05/24 09 | Ack Date : | 08/05/2024 10:16 | Report Date : | 08/05/24 10:53 |
| BUN-SERUM | | | | | | |
| BUN - SERUM Method - Urease-GLDH | | | 11.65 | | mg/dl | 4 - 18 |
| References: 1)Pack Insert of Bio | system | | | | | |

End of Report -

Dr.Ritesh Kharche MD, PGD-HM

Consultant Pathologist and Director of Laboratory Services

RegNo: 2006/03/1680



Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

Episode : OP

Ref. Doctor : Self Mobile No : 9702628411

DOB : 11/01/1992

Facility: SEVENHILLS HOSPITAL, MUMBAI

HAEMATOLOGY

| Test Name | | | Result | t | Unit | Bio | logical Reference Interval |
|-------------|-----------------|-------------------|--------------|-------------------|------------------|---------------|----------------------------|
| Sample No : | O0330555A | Collection Date : | 08/05/24 09: | 41 Ack Date : | 08/05/2024 10:15 | Report Date : | 08/05/24 12:13 |
| COMPLETE | E BLOOD COUNT | Γ (CBC) - EDTA | WHOLE BLO | OOD# | | | |
| Total WBC | Count | | | 7.89 | | x10^3/ul | 4 - 10 |
| Neutrophils | | | | 57.2 | | % | 40 - 80 |
| Lymphocyte | es | | | 33.3 | | % | 20 - 40 |
| Eosinophils | | | | 3.7 | | % | 1 - 6 |
| Monocytes | | | | 5.5 | | % | 2 - 10 |
| Basophils | | | | 0.3 ▼ (L) | | % | 1 - 2 |
| Absolute Ne | eutrophil Count | | | 4.52 | | x10^3/ul | 2 - 7 |
| Absolute Ly | mphocyte Count | | | 2.62 | | x10^3/ul | 0.8 - 4 |
| Absolute Eo | sinophil Count | | | 0.29 | | x10^3/ul | 0.02 - 0.5 |
| Absolute Mo | onocyte Count | | | 0.43 | | x10^3/ul | 0.12 - 1.2 |
| Absolute Ba | sophil Count | | | 0.03 | | x10^3/ul | 0 - 0.1 |
| RBCs | | | | 5.74 ▲ (H) | | x10^6/ul | 4.5 - 5.5 |
| Hemoglobin | 1 | | | 17.2 ▲ (H) | | gm/dl | 13 - 17 |
| Hematocrit | | | | 49.5 | | % | 40 - 50 |
| MCV | | | | 86.2 | | fl | 83 - 101 |
| MCH | | | | 30.0 | | pg | 27 - 32 |



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| MCHC | 34.8 ▲ (H) | gm/dl | 31.5 - 34.5 |
|---|--------------------|----------|--------------|
| RED CELL DISTRIBUTION WIDTH-CV (RDW-CV) | 13.0 | % | 11 - 16 |
| RED CELL DISTRIBUTION WIDTH-SD (RDW-SD) | 42.5 | fl | 35 - 56 |
| Platelet | 354 | x10^3/ul | 150 - 410 |
| Mean Platelet Volume (MPV) | 9.2 | fl | 6.78 - 13.46 |
| PLATELET DISTRIBUTION WIDTH (PDW) | 16.3 | % | 9 - 17 |
| PLATELETCRIT (PCT) | 0.324 ▲ (H) | % | 0.11 - 0.28 |

Method:-

HB Colorimetric Method.

RBC/PLT Electrical Impedance Method.

WBC data Flow Cytometry by Laser Method.

MCV,MCH,MCHC,RDW and rest parameters - Calculated.

All Abnormal Haemograms are reviewed confirmed microscopically.

NOTE: Wallach's Interpretation of Diagnostic Tests. 11th Ed, Editors: Rao LV. 2021

NOTE :-

The International Council for Standardization in Haematology (ICSH) recommends reporting of absolute counts of various WBC subsets for clinical decision making. This test has been performed on a fully automated 5 part differential cell counter which counts over 10,000 WBCs to derive differential counts. A complete blood count is a blood panel that gives information about the cells in a patient's blood, such as the cell count for each cell type and the concentrations of Hemoglobin and platelets. The cells that circulate in the bloodstream are generally divided into three types: white blood cells (leukocytes), red blood cells (erythrocytes), and platelets (thrombocytes). Abnormally high or low counts may be physiological or may indicate disease conditions, and hence need to be interpreted clinically.

End of Report

Dr.Ritesh Kharche MD, PGD-HM



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Consultant Pathologist and Director of Laboratory Services

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COMPLETE BLOOD COUNT (CBC) - EDTA WHOLE BLOOD- Report has been amended at May 8 2024 11:29AM by SAKSHI P PALANDE.



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Episode : OP

Ref. Doctor : Self **Mobile No** : 9702628411

DOB : 11/01/1992

Facility: SEVENHILLS HOSPITAL, MUMBAI

Biochemistry

| Test Name | | | Result | | Unit | Biol | ogical Reference Interval |
|-------------|-----------|-------------------|----------------|------------|------------------|---------------|---------------------------|
| Sample No : | O0330555B | Collection Date : | 08/05/24 09:41 | Ack Date : | 08/05/2024 10:16 | Report Date : | 08/05/24 10:53 |
| | | | | | | | |

| GLUCOSE-PLASMA-FASTING | | | |
|------------------------|-------|-------|----------|
| Glucose,Fasting | 99.54 | mg/dl | 70 - 100 |

American Diabetes Association Reference Range:

Normal: < 100 mg/dl

Impaired fasting glucose(Prediabetes): 100 - 126 mg/dl

Diabetes : >= 126 mg/dl

References:

1)Pack Insert of Bio system

2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

Interpretation :-

Conditions that can result in an elevated blood glucose level include: Acromegaly, Acute stress (response to trauma, heart attack, and stroke for instance), Chronic kidney disease, Cushing syndrome, Excessive consumption of food, Hyperthyroidism, Pancreatitis.

A low level of glucose may indicate hypoglycemia, a condition characterized by a drop in blood glucose to a level where first it causes nervous system symptoms (sweating, palpitations, hunger, trembling, and anxiety), then begins to affect the brain (causing confusion, hallucinations, blurred vision, and sometimes even coma and death). A low blood glucose level (hypoglycemia) may be

seen with:Adrenal insufficiency, Drinking excessive alcohol, Severe liver disease, Hypopituitarism, Hypothyroidism, Severe infections, Severe heart failure, Chronic kidney (renal) failure, Insulin overdose, Tumors that produce insulin (insulinomas), Starvation.

| | | | 00/05/04 00 44 | | 00/05/0004 40 46 | | 00/05/04 40 50 |
|------------|-----------|-------------------|----------------|------------|------------------|---------------|----------------|
| Sample No: | O0330555C | Collection Date : | 08/05/24 09:41 | Ack Date : | 08/05/2024 10:16 | Report Date : | 08/05/24 10:53 |

| ALT(SGPT) - SERUM | | | |
|--|--------------------|------|--------|
| SGPT (Alanine Transaminase) - SERUM Method - IFCC | 57.94 ▲ (H) | IU/L | 0 - 45 |
| References : | | | |



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DOB : 11/01/1992

Facility: SEVENHILLS HOSPITAL, MUMBAI

1)Pack Insert of Bio system

2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

| , | · · · · · · · · · · · · · · · · · · · | | |
|--|---------------------------------------|-------|-----------|
| Total Bilirubin - SERUM Method - Diazo | 0.84 | mg/dl | 0 - 2 |
| Direct Bilirubin SERUM Method - Diazotization | 0.33 | mg/dl | 0 - 0.4 |
| Indirect Bilirubin - Calculated Method - Calculated | 0.51 | mg/dl | |
| CREATININE-SERUM | | | |
| Creatinine - SERUM Method - Jaffes Kinetic | 1.02 | mg/dl | 0.5 - 1.3 |

References:

- 1)Pack Insert of Bio system
- 2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

Notes :-

Creatinine is a chemical waste molecule that is generated from muscle metabolism. Creatinine is produced from creatine, a molecule of major importance for energy production in muscles. Approximataly 1-2% of the body's creatine is converted to creatinine every day. Creatinine is transported through the bloodstream to the kidneys. The kidneys filter out host of the creatinine and dispose of it in the urine. The kidneys maintain the blood creatinine in a normal ranges. Creatinine has been found to be a fairly reliable indicator of kidney function.

| GLUCOSE-PLASMA POST PRANDIAL | | | |
|------------------------------|--------|-------|----------|
| Glucose, Post Prandial | 126.28 | mg/dl | 70 - 140 |

American Diabetes Association Reference Range:

Post-Prandial Blood Glucose:

Non- Diabetic: Up to 140mg/dL Pre-Diabetic: 140-199 mg/dL Diabetic: >200 mg/dL

References:

- 1)Pack Insert of Bio system
- 2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018



Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

UHID : SHHM.94020 : 08/05/2024 09:28 **Order Date**

: OP **Episode**

Ref. Doctor Mobile No : Self : 9702628411

DOB : 11/01/1992

: SEVENHILLS HOSPITAL, MUMBAI **Facility**

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Conditions that can result in an elevated blood glucose level include: Acromegaly, Acute stress (response to trauma, heart attack, and stroke for instance), Chronic kidney disease, Cushing syndrome, Excessive consumption of food, Hyperthyroidism, Pancreatitis.

A low level of glucose may indicate hypoglycemia, a condition characterized by a drop in blood glucose to a level where first it causes nervous system symptoms (sweating, palpitations, hunger, trembling, and anxiety), then begins to affect the brain (causing confusion, hallucinations, blurred vision, and sometimes even coma and death). A low blood glucose level (hypoglycemia) may be

seen with:Adrenal insufficiency, Drinking excessive alcohol, Severe liver disease, Hypopituitarism, Hypothyroidism, Severe infections, Severe heart failure, Chronic kidney (renal) failure, Insulin overdose, Tumors that produce insulin (insulinomas), Starvation.

End of Report

MD, PGD-HM

Consultant Pathologist and Director of Laboratory Services

RegNo: 2006/03/1680





Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

Episode : OP

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DOB : 11/01/1992

Facility: SEVENHILLS HOSPITAL, MUMBAI

Urinalysis

| Test Name R | esult Unit | Biological Reference Interval |
|--|----------------------------------|-------------------------------|
| Sample No: O0330555D Collection Date: 08/05/24 | 09:41 Ack Date: 08/05/2024 10:16 | Report Date : 08/05/24 14:39 |
| Physical Examination | | |
| QUANTITY | 20 | ml |
| Colour | Yellow | |
| Appearance | Clear | |
| DEPOSIT | Absent | Absent |
| рН | Acidic | |
| Specific Gravity | 1.015 | |
| Chemical Examination | | |
| Protein | Absent | Absent |
| Sugar | Absent | Absent |
| ketones | Absent | Absent |
| Occult Blood | POSITIVE (+) | Negative |
| Bile Salt | Absent | Absent |
| Bile Pigments | Absent | Absent |
| Urobilinogen | NORMAL | Normal |
| NITRATE | Absent | Absent |
| LEUKOCYTES | Absent | Absent |

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| Microscopic Examination | | | |
|-------------------------|--------|------|--------|
| Pus cells | 2-3 | /HPF | |
| Epithelial Cells | 1-2 | /HPF | |
| RBC | 5-6 | /HPF | Absent |
| Cast | Absent | /LPF | Absent |
| Crystal | Absent | /HPF | Absent |
| Amorphous Materials | Absent | | Absent |
| Yeast | Absent | | Absent |
| Bacteria | Absent | | Absent |

End of Report -

Dr.Pooja Vinod Mishra MD Pathology

Jr Consultant Pathologist, MMC Reg No. 2017052191

RegNo: 2017/05/2191



Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

Episode : OP

Ref. Doctor : Self **Mobile No** : 9702628411

DOB : 11/01/1992

Facility: SEVENHILLS HOSPITAL, MUMBAI

Blood Bank

Test Name Result

Sample No: 00330555A Collection Date: 08/05/24 09:41 Ack Date: 08/05/2024 12:28 Report Date: 08/05/24 13:07

| BLOOD GROUPING/ CROSS-MATCHING BY SEMI AUTOMATION# | | | | |
|--|----------|--|--|--|
| BLOOD GROUP (ABO) | ' AB ' | | | |
| Rh Type Method - Column Agglutination | POSITIVE | | | |

REMARK: THE REPORTED RESULTS PERTAIN TO THE SAMPLE RECEIVED AT THE BLOOD CENTRE.

Interpretation:

Blood typing is used to determine an individual's blood group, to establish whether a person is blood group A, B, AB, or O and whether he or she is Rh positive or Rh negative. Blood typing has the following significance,

- Ensure compatibility between the blood type of a person who requires a transfusion of blood or blood components and the ABO and Rh type of the unit of blood that will be transfused.
- Determine compatibility between a pregnant woman and her developing baby (fetus). Rh typing is especially important during pregnancy because a mother and her fetus could be incompatible.
- Determine the blood group of potential blood donors at a collection facility.
- Determine the blood group of potential donors and recipients of organs, tissues, or bone marrow, as part of a workup for a transplant procedure.

End of Report

Dr.Pooja Vinod Mishra MD Pathology

Jr Consultant Pathologist, MMC Reg No. 2017052191

RegNo: 2017/05/2191



Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

Ref. Doctor: Self **Mobile No**: 9702628411

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ears

20

...... T flat/neg, II III aVF Borderline T abnormalities, inferior leads.....

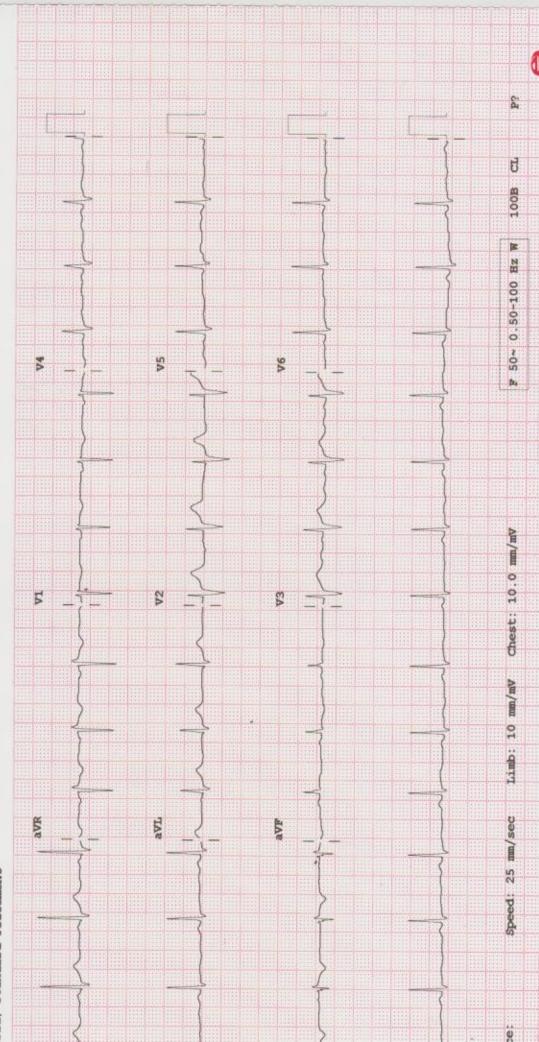
TD I DITTION OF

131 95 354 424 45 23 -26

IS--

- BORDERLINE ECG -

ead; Standard Placement



Patient Name : Mr. ROHIT BHAGWAN PARAB Age/Sex : 32 Year(s) / Male

Episode : OP

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Facility: SEVENHILLS HOSPITAL, MUMBAI

HAEMATOLOGY

| Test Name | | | Result | | Unit Bi | | ological Reference Interval | |
|-------------|------------|-------------------|----------------|------------|------------------|---------------|-----------------------------|--|
| Sample No : | O0330555A | Collection Date : | 08/05/24 09:41 | Ack Date : | 08/05/2024 10:15 | Report Date : | 08/05/24 12:13 | |
| EDVTUDO | CVTE CEDIA | AENTATION DATE /E | (CD) | | | | | |

| ERYTHROCYTE SEDIMENTATION RATE (ESR) | | | |
|--------------------------------------|----|-------|--------|
| ESR | 15 | mm/hr | 0 - 20 |

Method: Westergren Method

INTERPRETATION :-

ESR is a non-specific phenomenon, its measurement is clinically useful in disorders associated with an increased production of acute-phase proteins. It provides an index of progress of the disease in rheumatoid arthritis or tuberculosis, and it is of considerable value in diagnosis of temporal arteritis and polymyalgia rheumatica. It is often used if multiple myeloma is suspected, but when the myeloma is non-secretory or light chain, a normal ESR does not exclude this diagnosis.

An elevated ESR may occur as an early feature in myocardial infarction. Although a normal ESR cannot be taken to exclude the presence of organic disease, the vast majority of acute or chronic infections and most neoplastic and degenerative diseases are associated with changes in the plasma proteins that increased ESR values.

The ESR is influenced by age, stage of the menstrual cycle and medications taken (corticosteroids, contraceptive pills). It is especially low (0–1 mm) in polycythaemia, hypofibrinogenaemia and congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis, or sickle cells. In cases of performance enhancing drug intake by athletes the ESR values are generally lower than the usual value for the individual and as a result of the increase in haemoglobin (i.e. the effect of secondary polycythaemia).

- End of Report -

Dr.Ritesh Kharche MD, PGD-HM

Consultant Pathologist and Director of Laboratory Services

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Page 2 of 2



Arcofemi Healthcare Pvt Ltd

(Formerly known as Arcofemi Healthcare Ltd) F-701A, Lado Sarai, Mehrauli, New Delhi - 110030

MEDICAL FITNESS CERTIFICATE

(To be signed by a registered medical practitioner holding a Medical degree)

This is to certify that <u>Mr.Rohit Bhagwan</u> aged, <u>31yr</u>. Based on the examination, I certify that he is in good mental and physical health and it is free from any physical defects such as deafness, colour blindness, and any chronic or contagious diseases.

Place: Mumbai

Date: 08/05/2024

Mitter

Dr. Nitesh Kumar

Name & Signature of

Medical officer