

AASHIESH GUPTA
Flat No-M-301, Marvel Diva,
Magarpatta Road, Near Seasons Mall,
Hadapsar, Pune-28
Tel No: 918879699138
PID: 11531066

Reference: Dr.--

SID: 121560732

121560732

Collection Date:

18-06-2021 01:21 PM

Sample Date:

18-06-2021 01:21 pm

Report Date:

18-06-2021 06:02 PM

F-----

Age:38.20 Years Sex:MALE

Complete Blood Count

(EDTA Whole Blood)

Hemoglobin (Hb), EDTA whole blood

14.40

14.0 - 17.50 g/dL

Method: Photometry

Total Leucocytes (WBC) count

7,300

4000-10000/ μ L

Method : Coulter Principle / Microscopy

Platelet count

367,000

150000 - 450000 / μ L

Method : Coulter Principle / Microscopy

Red blood cell (RBC) count

5.01

4.52 - 5.90 x 10⁶ / μ L

Method: Coulter Principle

PCV (Packed Cell Volume)

42.70

41.5 - 50.4 %

Method: Calculated

MCV (Mean Corpuscular Volume)

85.20

80.0 - 96.0 fL

Method: Derived from RBC histogram

MCH (Mean Corpuscular Hb)

28.80

27.5 - 33.2 pgms

Method: Calculated

MCHC (Mean Corpuscular Hb Conc.)

33.80

33.4 - 35.5 g/dL

Method: Calculated

RDW (RBC distribution width)

13.30

11.6 - 14.6 %

Method: Derived from RBC Histogram

WBC Differential Count

Method: VCSn / Microscopy / Calculated

Neutrophils

57

40 - 80 %

Absolute Neutrophils

4,161

2000 - 7000 / μ L

Eosinophils

5

1 - 6 %

Absolute Eosinophils

365

20 - 500 / μ L

Basophils

0

0 - 2 %

Absolute Basophils

0

0 - 100 / μ L

Lymphocytes

33

20 - 40 %

Absolute Lymphocytes

2,409

1000 - 3000 / μ L

Monocytes

5

2 - 10 %

Absolute Monocytes

365

200 - 1000 / μ L

-

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"Accreditation as per ISO 15189:2012, Cert.No. MC-3143. Refer scope@ www.nabl-india.org"

Awanti Golwilkar Mehendale
Dr.(Mrs.) Awanti Golwilkar Mehendale
MBBS,MD(Path) Regn.No:2000/02/1052
A.G Diagnostics Pvt. Ltd.

Carrying forward
Dr. Ajit Golwilkar's
legacy of Over
Four Decades

DIAGNOSTICS

BE SURE
BE WELL

ए.जी. डायग्नॉस्टिक्स प्रा. लि. A.G Diagnostics Pvt. Ltd.
a Neuberg associate

Dr. Awanti Golwilkar
MBBS, MD (Pathology)

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Age:38.20 Years Sex:MALE

Complete Blood Count Findings

R.B.C. : Normocytic, Normochromic

W.B.C. : No abnormality detected

Platelets : Adequate

Remark : --

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Age:38.20 Years Sex:MALE

Test Description

Liver Function Test :

Test Description	Observed	Biological Reference Interval
Bilirubin-Total, serum by Diazo method	0.61	0.10 - 1.20 mg/dL Neonates : Upto 15.0 mg/dL
Bilirubin-Conjugated, serum by Diazo method	0.25	Upto 0.5 mg/dL
Bilirubin-Unconjugated, serum by calculation	0.36	0.1 to 1.0 mg/dL
SGOT (AST), serum by Enzymatic method	26	>or= 14 years : 8 - 48 U/Lt
SGPT (ALT), serum by Enzymatic Method	43	7 to 55 U/Lt
Alkaline Phosphatase,serum by pNPP-kinetic	61	Adult Male : (Unit : U/Lt.) 15 - < 17 years : 82 - 331 17 - < 19 years : 55 - 149 > or = 19 years : 40 - 129
Protein (total), serum by Biuret method	7.01	6.4 to 8.2 g/dL
Albumin, serum by Bromocresol purple method	4.16	3.4 to 5.0 g/dL
Globulin, serum by calculation	2.85	2.3 - 3.5 g/dL

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Test Description**Observed Value****Biological Reference Interval****Plasma Glucose :**

Plasma glucose, random by Hexokinase method

231

< 200 mg/dL

American Diabetes Association
Guidelines 2020**Clinical Chemistry**

Urea, serum by GLDH-urease

20

17 to 49 mg/dL

BUN-Blood Urea Nitrogen,serum by calculation

9.35

8 to 23 mg/dL

Creatinine, serum by Jaffe w/o deproteinization

1.00

0.6 to 1.2 mg/dL

Uric Acid, serum by Uricase method

5.30

Male : 3.50 to 7.20 mg/dL

** Uric acid is useful for 1. Diagnosis and follow up of renal failure. 2. Monitoring patients receiving cytotoxic drugs and a variety of other disorders, including gout, leukemia, psoriasis, starvation and other wasting conditions*

*. * Increased uric acid is seen in following conditions :*

*1. Increased purine synthesis 2. Inherited metabolic disorders 3. Excess dietary purine intake
4. Increased nucleic acid turnover 5. Malignancy, cytotoxic drugs 6. Decreased urinary excretion
(due to CRF) 7. Increased renal reabsorption .*

** Uric acid is decreased in : 1. Hepatocellular disease with reduced purine synthesis
2. Defective renal reabsorption 3. Overtreatment of uricemia (allopurinol or cancer
therapies like 6-mercaptopurine, etc).*

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Test Description
Clinical Chemistry :**Observed Value****Biological Reference Interval****Infectious Diseases**

HIV Combo(p24Ag,HIV1Ab,HIV2Ab),Serum by CMIA

Non-reactive (0.06)

Non-reactive : Less than 1.00 S/CO

*HIV Combo(p24Ag,HIV1Ab,HIV2Ab) by CMIA is a 4th generation test.**HIV Antibodies & p24 may be negative in window period (seroconversion period). All positive results need confirmation by another EIA / Western Blot on fresh sample.*

HBsAg, serum by CMIA

Non-reactive(< 0.02)

Non-reactive : < 0.05 IU/mL

Coagulation

Prothrombin Time, Citrated Plasma - Patient value

10.90

10.30 - 13.30 Secs

Prothrombin time - Control value

11.80

Secs

INR Value

0.92

0.85 - 1.15 (ISI : 1.05)

**TEST DONE ON : AUTOMATED BLOOD COAGULATION ANALYZER,
CA- 600 SERIES, SYSMEX CORP., JAPAN.
PHOTO OPTICAL CLOT DETECTION METHOD.**

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Test Description

Coagulation :

Observed Value

Biological Reference Interval



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Urine Routine Examination

(Sample : Urine, Automated / Semiautomated)

Physical

Quantity Examined

Method : Visual

Appearance

Method : Visual / Automated

Colour

Method : Visual / Automated

Chemical (Dipstick)

pH

Method : Indicator Principle

Protein

Method : Sulphosalicylic Acid/ pH Indicator

Glucose

Method : GOD-POD / Benedict's

Acetone

Method : Sodium Nitroprusside reaction

Bile Pigments

Method : Diazo Reaction / Fouchet's test

Urobilinogen

Method : Modified Ehrlich / Watson Schwartz

Microscopy / Flow cytometry

R.B.Cs

Pus cells

Epithelial cells

Casts

Crystals

-

Result

5.0

Clear

Pale yellow

5.5

Absent

Present 1+

Absent

Absent

Not significant

1-2

1-2

Occasional

Not Detected

Not Detected

<-->

Biological Reference Interval

ml

-

-

4.6 - 8.0

Absent

Absent

Absent

Absent

Not Significant

0 - 2 per hpf

0 - 5 per hpf

0 - 5 per hpf

-

-



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Bleeding time & Clotting time

	<u>Result</u>	<u>Biological Reference Interval</u>
Bleeding time	3 Mins 00 Secs	2 - 7 min.s
Clotting time	7 Mins 00 Secs	4 - 9 min.s

Interpretation :

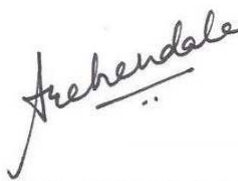
PT, APTT, platelet count and morphology - together can work as a good preoperative panel to assess surgical bleeding.

Bleeding time :

1. Provides useful information during the evaluation of bleeding disorders.
2. Poor predictive value of actual clinical bleeding and hence play limited role as a marker for preoperative hemostatic screening of asymptomatic patients.
3. Bleeding time may be prolonged in :
 - i. Thrombocytopenia (usually Platelet count < 1,00,000 / μ L).
 - ii. von Willebrand disease.
 - iii. Drugs with antiplatelet action (eg: aspirin, clopidogrel, etc.)
 - iv. Acquired and congenital disorders of platelet function.

Clotting time :

Prolonged clotting time may be seen in severe coagulation factor deficiencies (Hemophilia A, Hemophilia B, etc.) and marked thrombocytopenia.


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