

Name

TEST

: Mrs KALPNABEN B VASAVA





REMARKS

LABORATORY REPORT

Sex/Age : Female / 35 Years Case ID: 40308001493

BIOLOGICAL REF. INTERVAL

0

Pt. ID Ref. By : Mediwheel Full Body Health Checkup Dis. At

Bill. Loc. : Health packages Pt. Loc

Reg Date and Time : 30-Mar-2024 07:57 Sample Type : Whole Blood EDTA Mobile No. :

Ref Id1 Sample Date and Time : 30-Mar-2024 07:57 Sample Coll. By : non Report Date and Time : 30-Mar-2024 09:57 Acc. Remarks Ref Id2

UNIT

HAEMOGRAM REPORT HB AND INDICES Haemoglobin 11.4 G% 12.0 - 15.0 **RBC** (Electrical Impedance) 5.60 millions/cumm 3.80 - 4.80

PCV(Calc) 37.41 % 36.00 - 46.00 MCV (RBC histogram) L 66.8 fL 83.00 - 101.00 20.3 27.00 - 32.00 MCH (Calc) pg MCHC (Calc) 30.4 gm/dL 31.50 - 34.50

RESULTS

RDW (RBC histogram) 15.40 % 11.00 - 16.00

TOTAL AND DIFFERENTIAL WBC COUNT **Total WBC Count** 6390 /µL 4000.00 - 10000.00

EXPECTED VALUES 40.00 - 70.00 **EXPECTED VALUES** /μL 2000.00 - 7000.00 Neutrophil /µL 1000.00 - 3000.00 Lymphocyte 19 20.00 - 40.00 1214 **Eosinophil** 02 % 1.00 - 6.00 128 /µL 20.00 - 500.00 2.00 - 10.00 383 /µL 200.00 - 1000.00 Monocytes 06 0.00 - 2.00/µL 0.00 - 100.00

PLATELET COUNT

Platelet Count 258000 /µL 150000.00 - 410000.00 **MPV** fL 6.5 - 12 9.20

PDW 8 - 13 15.5

00

Method:

Basophil

TLC-SF cube technology(Flow Cytometry+ fluorescence),

DC by microscopy,

Platelet count by electrical impedance+/-SF cube technology

Note:(LL-VeryLow,L-Low,H-High,HH-VeryHigh ,A-Abnormal)

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Consultant Pathologist

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SKATOKT REPORT

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TEST RESULTS UNIT BIOLOGICAL REF RANGE REMARKS

HAEMATOLOGY INVESTIGATIONS

ESR 12 mm after 1hr 3 - 20 Westergren Method

BLOOD GROUP AND RH TYPING (Erythrocyte Magnetized Technology) (Both Forward and Reverse Group)

ABO Type AB

Rh Type POSITIVE

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RY REPORT

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Reg Date and Time : 30-Mar-2024 07:57 | Sample Type : Plasma Fluoride F,Plasma | Mobile No. :

Fluoride PP,Whole Blood

EDTA

Sample Date and Time : 30-Mar-2024 07:57 | Sample Coll. By : non | Ref Id1

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TEST RESULTS UNIT BIOLOGICAL REF RANGE REMARKS

Plasma Glucose - F
Photometric, Hexokinase

H 104.27 mg/dL 70 - 100 FUS: NIL

Plasma Glucose - PP
Photometric, Hexokinase

133.98 mg/dL 70 - 140 PPUS: NIL

Glycated Haemoglobin Estimation

HbA1C 5.3 % of total Hb <5.7: Normal Immunoturbidimetric 5.4 6.4: Prod

5.7-6.4: Prediabetes

>=6.5: Diabetes

Estimated Avg Glucose (3 Mths) 105.41 mg/dL Not available

Please Note change in reference range as per ADA 2021 guidelines.

Interpretation:

HbA1C level reflects the mean glucose concentration over previous 8-12 weeks and provides better indication of long term glycemic control. Levels of HbA1C may be low as result of shortened RBC life span in case of hemolytic anemia.

Increased HbA1C values may be found in patients with polycythemia or post splenectomy patients.

Patients with Homozygous forms of rare variant Hb(CC,SS,EE,SC) HbA1c can not be quantitated as there is no HbA.

In such circumstances glycemic control can be monitored using plasma glucose levels or serum Fructosamine.

The A1c target should be individualized based on numerous factors, such as age, life expectancy, comorbid conditions, duration of diabetes, risk of hypoglycemia or adverse consequences from hypoglycemia, patient motivation and adherence.

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Pt. ID Ref. By : Mediwheel Full Body Health Checkup Dis. At Bill. Loc. : Health packages Pt. Loc :

Reg Date and Time : 30-Mar-2024 07:57 Sample Type : Serum Mobile No. :

Sample Date and Time : 30-Mar-2024 07:57 Sample Coll. By : non Ref Id1 : 30-Mar-2024 09:57 Ref Id2 Report Date and Time Acc. Remarks

TEST	RESULTS	UNIT	BIOLOGICAL REF RANGE	REMARKS
BUN (Blood Urea Nitrogen)	10.1	mg/dL	7.00 - 18.70	
Uric Acid Uricase-Peroxidase method	5.53	mg/dL	2.6 - 6.2	
Creatinine Jaffe compensated	0.71	mg/dL	0.55 - 1.02	

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: Mrs KALPNABEN B VASAVA

Name





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TEST RESULTS UNITBIOLOGICAL REF RANGE **REMARKS**

BIOCHEMICAL INVESTIGATIONS

Lipid Profile

Cholesterol Colorimetric, CHOD-POD	162.73	mg/dL	110 - 200	
HDL Cholesterol	46.0	mg/dL	40 - 60	
Triglyceride GPO-POD	94.55	mg/dL	40 - 200	
VLDL Calculated	18.91	mg/dL	10 - 40	
Chol/HDL Calculated	3.54		0 - 4.1	
LDL Cholesterol Calculated	97.82	mg/dL	0.00 - 100.00	

NEW ATP III GUIDELINES (MAY 2001), MODIFICATION OF NCEP

LDL CHOLESTEROL	CHOLESTEROL	HDL CHOLESTEROL	TRIGLYCERIDES
Optimal<100	Desirable<200	Low<40	Normal<150
Near Optimal 100-129	Border Line 200-239	High >60	Border High 150-199
Borderline 130-159	High >240		High 200-499
High 160-189	-		-

- LDL Cholesterol level is primary goal for treatment and varies with risk category and assesment
- For LDL Cholesterol level Please consider direct LDL value
 - Risk assessment from HDL and Triglyceride has been revised. Also LDL goals have changed.
- Detail test interpreation available from the lab
- All tests are done according to NCEP guidelines and with FDA approved kits.
- · LDL Cholesterol level is primary goal for treatment and varies with risk category and assesment

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RESULTS TEST UNITBIOLOGICAL REF RANGE **REMARKS**

BIOCHEMICAL INVESTIGATIONS

Liver Function Test

Liver Function Fest					
S.G.P.T.	19.28	U/L	0 - 59		
S.G.O.T.	15.91	U/L	15 - 37		
Alkaline Phosphatase Modified IFCC method	98.43	U/L	40 - 150		
Proteins (Total) Biuret	7.05	g/dL	6.4 - 8.2		
Albumin Bromo Cresol Green	4.33	g/dL	3.4 - 5.0		
Globulin Calculated	2.72	gm/dL	2 - 4.1		
A/G Ratio Calculated	1.6		1.0 - 2.1		
Bilirubin Total Diazotized Sulfanilic Acid Method	0.26	mg/dL	0.2 - 1.0		
Bilirubin Conjugated Diazotized Sulfanilic Acid Method	0.12	mg/dL			
Bilirubin Unconjugated Calculated	0.14	mg/dL	0 - 0.8		

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: Mrs KALPNABEN B VASAVA





LABORATORY REPORT

Cay/Aga . Famala / 25 Vagra

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TEST RESULTS UNIT BIOLOGICAL REF RANGE REMARKS

BIOCHEMICAL INVESTIGATIONS

Thyroid Function Test

Triiodothyronine (T3)	1.43	ng/mL	0.70 - 2.04
Thyroxine (T4) ECLIA	8.25	μg/dL	5.5 - 11.0
TSH ECLIA	3.960	μIU/mL	0.40 - 4.20

INTERPRETATIONS

Name

Useful for Monitoring patients on thyroid replacement therapy, Confirmation of thyroid-stimulating hormone (TSH) suppression in thyroid cancer patients on thyroxine therapy, for Prediction of thyrotropin-releasing hormone-stimulated TSH response, as An aid in the diagnosis of primary hyperthyroidism, for Differential diagnosis of hypothyroidism.

The ability to quantitate circulating levels of thyroid-stimulating hormone (TSH) is important in evaluating thyroid function. It is especially useful in the differential diagnosis of primary (thyroid) from secondary (pituitary) and tertiary (hypothalamus) hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low or normal. Concentrations of 5.1 mIU/ml to 7.0 mIU/ml are considered borderline hypothyroid

CAUTIONS

Sick, hospitalized patients may have falsely low or transiently elevated thyroid stimulating hormone.

Some patients who have been exposed to animal antigens, either in the environment or as part of treatment or imaging procedure, may have circulating antianimal antibodies present. These antibodies may interfere with the assay reagents to produce unreliable results.

TSH ref range in Pregnacy Reference range (microlU/ml)

First trimester 0.24 - 2.00 Second trimester 0.43-2.2 Third trimester 0.8-2.5

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TEST RESULTS UNIT BIOLOGICAL REF RANGE REMARKS

URINE EXAMINATION (STRIP METHOD AND FLOWCYTOMETRY)

Physical examination

Name

Colour Pale yellow

Transparency Clear

Chemical Examination By Sysmex UC-3500

Sp.Gravity 1.025 1.003 - 1.035

рΗ 6.0 4.6 - 8 Leucocytes (ESTERASE) **Negative** Negative **Protein** Negative Negative Glucose **Negative** Negative **Ketone Bodies Urine** Negative Negative Urobilinogen **Negative** Negative **Bilirubin** Negative Negative **Blood** Negative Negative **Nitrite** Negative Negative

Flowcytometric Examination By Sysmex UF-5000

 Leucocyte
 Nil
 /HPF
 Nil

 Red Blood Cell
 Nil
 /HPF
 Nil

 Epithelial Cell
 3-5
 /HPF
 Present(+)

 Bacteria
 Nil
 /μL
 Nil

 Bacteria
 Nil
 /μL
 Nil

 Yeast
 Nil
 /μL
 Nil

 Cast
 Nil
 /HPF
 Nil

 Crystals
 Nil
 /HPF
 Nil

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Parameter Unit	Unit	Unit Expected value		Result/Notations				
		Trace	+	++	+++	++++		
pН	-	4.6-8.0	1 11 1 1 1 1 2	- 23	1-4-		2 1111	
SG	-	1.003-1.035	100		7		2) 	
Protein	mg/dL	Negative (<10)	10	25	75	150	500	
Glucose	mg/dL	Negative (<30)	30	50	100	300	1000	
Bilirubin	mg/dL	Negative (0.2)	0.2	1	3	6		
Ketone	mg/dL	Negative (<5)	5	15	50	150	1 2	
Urobilinogen	mg/dL	Negative (<1)	1	4	8	12	-	

Parameter	Unit	Expected value	e Result/Notifications			ons	
			Trace	+	++	+++	++++
Leukocytes (Strip)	/micro L	Negative (<10)	10	25	100	500	-
Nitrite(Strip)	-	Negative	-	-	-	-	-0
Erythrocytes(Strip)	/micro L	Negative (<5)	10	25	50	150	250
Pus cells (Microscopic)	/hpf	<5	71	5	87.8	-	ħ
Red blood cells (Microscopic)	/hpf	<2	27	-	-	-	-
Cast (Microscopic)	/lpf	<2	-		070	-	-

Pending Services ------ End Of Report ------ Stool Examination

For test performed on specimens received or collected from non-NSRL locations, it is presumed that the specimen belongs to the patient named or identified as labeled on the container/test request and such verification has been carried out at the point generation of the said specimen by the sender. NSRL will be responsible Only for the analytical part of test carried out. All other responsibility will be of referring Laboratory.

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