



MEDICAL EXAMINATION REPORT (MER)

If the examinee is suffering from an acute life threatening situation, you may be obliged to disclose the result of the medical examination to the examinee.

1. Name of the examinee	: Mr./Mrs./Ms. Smiles - 9
2. Mark of Identification	
Age/Date of Birth	: (Mole/Scar/any other (specify location)): : 434x (287 (1979.) Gender: F/M
4. Photo ID Checked	: (Passport/Election Card/PAN Card/Driving Licence/Company ID)

PHYSICAL DETAILS:

a. Height	b. Weight	c. Girth of Abdomen	
	1 st Reading		
	2 nd Reading		

FAMILY HISTORY:

Relation	Age if Living	Health Status	If deceased, age at the time and cause
Father		/	
Mother		11	PORT OF THE PROPERTY OF THE PR
Brother(s)		/ N S	
Sister(s)		and polipies and Title	Jap Till Y LL Welling a vestel stands time of

HABITS & ADDICTIONS: Does the examinee consume any of the following?

Tobacco in any form	Sedative	Alcohol
Traffer is a series of the finance in the	o namosilnos reliadad inflaid sondi	tall be made your I but imPana

PERSONAL HISTORY

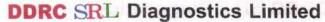
- a. Are you presently in good health and entirely free from any mental or Physical impairment or deformity. If No, please attach details.
- b. Have you undergone/been advised any surgical procedure?
- c. During the last 5 years have you been medically examined, received any advice or treatment or admitted to any hospital?
- d. Have you lost or gained weight in past 12 months?

Have you ever suffered from any of the following?

- Psychological Disorders or any kind of disorders of the Nervous System?
- · Any disorders of Respiratory system?
- · Any Cardiac or Circulatory Disorders?
- · Enlarged glands or any form of Cancer/Tumour?
- · Any Musculoskeletal disorder?

- · Any disorder of Gastrointestinal System?
- Unexplained recurrent or persistent fever, and/or weight loss
- Have you been tested for HIV/HBsAg / HCV before? If yes attach reports
- Are you presently taking medication of any kind?





Any disorders of Urinary System?	Y/N	Any disorder of the Eyes, Ears, Nose, Throat Mouth & Skin	t or Y/N
FOR FEMALE CANDIDATES ONLY			
a. Is there any history of diseases of breast/genital organs?	YN	d. Do you have any history of miscarriage/ abortion or MTP	YN,
b. Is there any history of abnormal PAP Smear/Mammogram/USG of Pelvis or any other tests? (If yes attach reports)	and the same	 e. For Parous Women, were there any complicated during pregnancy such as gestational diabeted hypertension etc 	
c. Do you suspect any disease of Uterus, Cervix or Ovaries?	Y/N	f. Are you now pregnant? If yes, how many mo	onths?
CONFIDENTAIL COMMENTS FROM MEDICA	AL EXA	MINER	
➤ Was the examinee co-operative?			CYN
Is there anything about the examine's health, life his/her job?	estyle tha	t might affect him/her in the near future with reg	gard to Y/N
Are there any points on which you suggest further	er inform	nation be obtained?	Y/N
Based on your clinical impression, please provid	le your si	aggestions and recommendations below;	
8.10	- 1	A second of	
NS -	Post	2010	
	.0.3		
➤ Do you think he/she is MEDICALLY FIT or UN	IFIT for	employment	
		• •	
	FIT		
MEDICAL EXAMINER'S DECLARATION			
I hereby confirm that I have examined the above indi- above are true and correct to the best of my knowledge		ter verification of his/her identity and the finding	gs stated
Name & Signature of the Medical Examiner :	360		
Seal of Medical Examiner :	r. GEOI	RGE THOMAS	

MEDICAL EXAMINER Reg: 86614

Name & Seal of DDRC SRL Branch

Date & Time

DDRC SRL Diagnostics Limited

DIAGNOSTIC SERVICES

Trom

Smithé. G. Bauk og Baroda

DORCSRL

Sub: Medical Check Up - Regarding

Du thé abour Course lión i novomld like & inform you that I don't require the following chek up/test

(Denlas

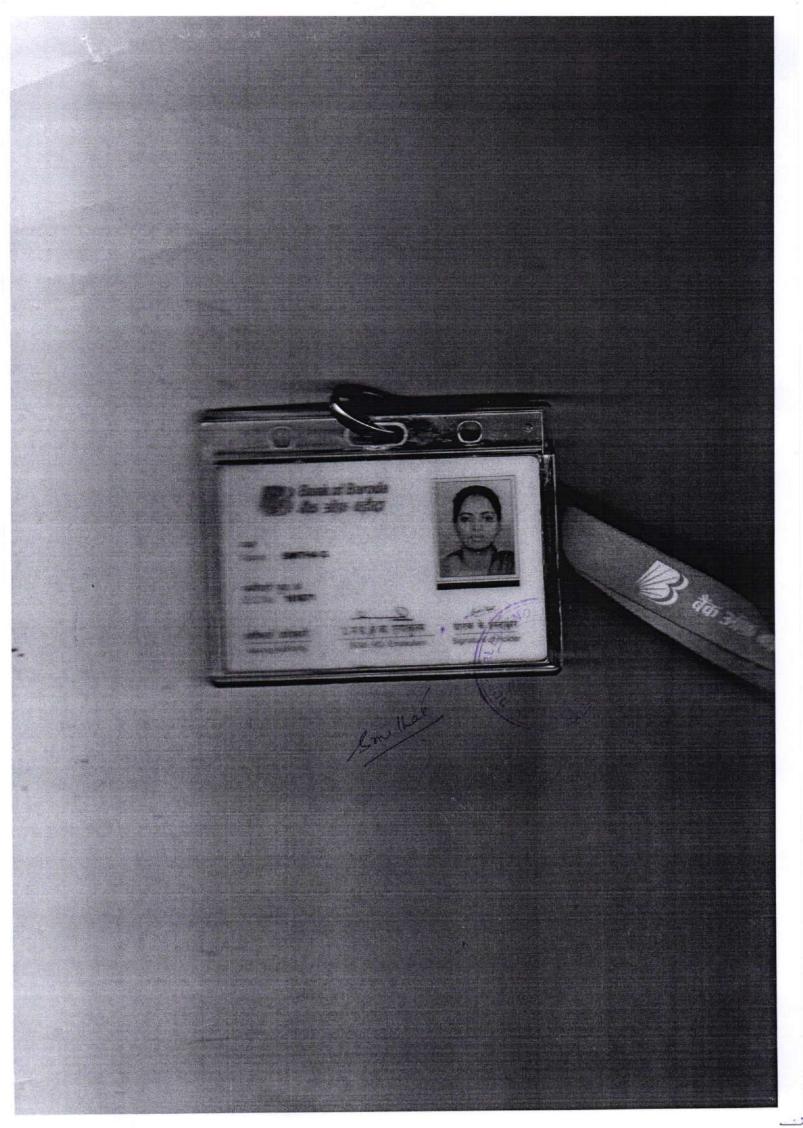
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DDRC SRL DIAGNOSTICS DDRC SRL Tower, G-131,Panampilly Nagar, PANAMPALLY NAGAR, 682036 KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: G.SMITHA

PATIENT ID : GSMIF2502804126

ACCESSION NO: 4126WB009076 AGE: 43 Years

ABHA NO :

DRAWN :

RECEIVED: 25/02/2023 08:54

REPORTED: 26/02/2023 12:47

REFERRING DOCTOR: DR. BOB

CLIENT PATIENT ID :

Test Report Status

Preliminary

Results

SEX: Female

Biological Reference Interval Units

MEDIWHEEL HEALTH CHECKUP ABOVE 40(F)TMT

OPTHAL

OPTHAL

TEST COMPLETED

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CIN: U85190MH2006PTC161480







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MEDIWHEEL HEALTH CHECKUP ABOVE 40(F	TMT(
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN METHOD: UREASE - UV	9	Adult(<60 yrs): 6 to	20 mg/dL
BUN/CREAT RATIO			
BUN/CREAT RATIO CREATININE, SERUM	6.92		
CREATININE METHOD: JAFFE KINETIC METHOD	0.66	18 - 60 yrs : 0.6 - 1.1	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA			
GLUCOSE, POST-PRANDIAL, PLASMA	87	Diabetes Mellitus : > c Impaired Glucose toler Prediabetes : 140 - 19 Hypoglycemia : < 55.	rance/
METHOD : HEXOKINASE		nypoglycemia : < 55.	
LIPID PROFILE, SERUM			
CHOLESTEROL	187	Desirable : < 200 Borderline : 200-239 High : > or = 240	mg/dL
METHOD : CHOD-POD		11igii . 701 = 240	
TRIGLYCERIDES	59	Normal : < 150 High : 150-199 Hypertriglyceridemia : Very High : > 499	mg/dL 200-499
HDL CHOLESTEROL METHOD: DIRECT ENZYME CLEARANCE	44	General range: 40-60	mg/dL
DIRECT LDL CHOLESTEROL	128	Optimum : < 10 Above Optimum : 100 Borderline High : 130 High : 160 Very High : >or=	139 -159 -189
NON HDL CHOLESTEROL	143	High Desirable: Less than 1: Above Desirable: 130 - Borderline High: 160 - High: 190 - 219 Very high: > or = 220	30 mg/dL 159
VERY LOW DENSITY LIPOPROTEIN	11.8	Desirable value : 10 - 35	mg/dL











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CHOL/HDL RATIO		4.3	3.3-4.4 Low Risk 4.5-7.0 Average Risk 7.1-11.0 Moderate Risk > 11.0 High Risk
LDL/HDL RATIO		2.9	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk











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Preliminary

Results

Units

Interpretation(s)

I) Cholesterol levels help assess the patient risk status and to follow the progress of patient under treatment to lower serum cholesterol concentrations.

2) Serum Triglyceride (TG) are a type of fat and a major source of energy for the body. Both quantity and composition of the diet impact on plasma triglyceride concentrations. Elevations in TG levels are the result of overproduction and impaired clearance. High TG are associated with increased risk for CAD (Coronary artery disease) in patients with other risk factors, such as low HDL-C, some patient groups with elevated apolipoprotein B concentrations, and patients with forms of LDL that may be particularly atherogenic.

3)HDL-C plays a crucial role in the initial step of reverse cholesterol transport, this considered to be the primary atheroprotective function of HDL

4) LDL -C plays a key role in causing and influencing the progression of atherosclerosis and, in particular, coronary sclerosis. The majority of cholesterol stored in atherosclerotic plaques originates from LDL, thus LDL-C value is the most powerful clinical predictor.

5)Non HDL cholesterol: Non-HDL-C measures the cholesterol content of all atherogenic lipoproteins, including LDL hence it is a better marker of risk in both primary and secondary prevention studies. Non-HDL-C also covers, to some extent, the excess ASCVD risk imparted by the sdLDL, which is significantly more atherogenic than the normal large buoyant particles, an elevated non-HDL-C indirectly suggests greater proportion of the small, dense variety of LDL particles

Serum lipid profile is measured for cardiovascular risk prediction. Lipid Association of India recommends LDL-C as primary target and Non HDL-C as co-primary treatment target.

Risk Stratification for ASCVD (Atherosclerotic cardiovascular disease) by Lipid Association of India

Extreme risk group	A.CAD with > 1 feature of high risk group	1	
		group or recurrent ACS (within 1 year) despite LDL-C	
Very High Risk	Established ASCVD 2. Diabetes with 2 major risk factors or evidence of end organ damage. Familial Homozygous Hypercholesterolemia.		
High Risk	organ damage. 3, CKD stage 3B or 4, 4,	hiabetes with 1 major risk factor or no evidence of end LDL>190 mg/dl 5. Extreme of a single risk factor. 6. 7. Lipoprotein a >/= 50mg/dl 8. Non stenotic carotid	
Moderate Risk	2 major ASCVD risk factors		
Moderate Risk Low Risk	2 major ASCVD risk factors 0-1 major ASCVD risk factors		
Low Risk	0-1 major ASCVD risk factors	eactors .	
Low Risk Major ASCVD (Ath	0-1 major ASCVD risk factors erosclerotic cardiovascular disease) Risk I		
Low Risk Major ASCVD (Ath	0-1 major ASCVD risk factors erosclerotic cardiovascular disease) Risk I s in males and > or = 55 years in females	3. Current Cigarette smoking or tobacco use 4. High blood pressure	

Newer treatment goals and statin initiation thresholds based on the risk categories proposed by LAI in 2020.

Risk Group	oup Treatment Goals Consider Drug Therapy		erapy	
	LDL-C (mg/dl)	Non-HDL (mg/dl)	LDL-C (mg/dl)	Non-HDL (mg/dl)



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GSMIF2502804126

Units

ACCESSION NO: 4126WB009076 AGE: 43 Years

Test Report Status

SEX: Female

Results

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MEAN PLASMA GLUCOSE

ALBUMIN/GLOBULIN RATIO

CLIENT PATIENT ID :

Extreme Risk Group Category A	<50 (Optional goal < OR = 30)	< 80 (Optional goal <or 60)<="" =="" th=""><th>>OR = 50</th><th>>OR = 80</th></or>	>OR = 50	>OR = 80
Extreme Risk Group Category B	<or 30<="" =="" td=""><td><or 60<="" =="" td=""><td>> 30</td><td>>60</td></or></td></or>	<or 60<="" =="" td=""><td>> 30</td><td>>60</td></or>	> 30	>60
Very High Risk	<50	<80	>OR= 50	>OR= 80
High Risk	<70	<100	>OR= 70	>OR= 100
Moderate Risk	<100	<130	>OR= 100	>OR= 130
Low Risk	<100	<130	>OR= 130*	>OR 150

^{*}After an adequate non-pharmacological intervention for at least 3 months.

Preliminary

References: Management of Dyslipidaemia for the Prevention of Stroke: Clinical Practice Recommendations from the Lipid Association of India. Current Vascular Pharmacology, 2022, 20, 134-155.

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE

GLYCOSYLATED HEMOGLOBIN (HBA1C)

: 4.0 - 5.6%. %

Non-diabetic level : < 5.7%. Diabetic

: >6.5%

Glycemic control goal

More stringent goal : < 6.5 %. : < 7%. General goal Less stringent goal : < 8%.

Glycemic targets in CKD :-If eGFR > 60 : < 7%.

If eGFR < 60: 7 - 8.5%. < 116.0

LIVER FUNCTION TEST WITH GGT			
BILIRUBIN, TOTAL METHOD: DIAZO METHOD	0.43	General Range : < 1.1	mg/dL
BILIRUBIN, DIRECT	0.18	General Range : < 0.3	mg/dL

99.7

METHOD : DIAZO METHOD BILIRUBIN, INDIRECT 0.25 0.00 - 0.60mg/dL TOTAL PROTEIN 7.1 Ambulatory: 6.4 - 8.3 g/dL Recumbant: 6 - 7.8

ALBUMIN 4.1 20-60yrs: 3.5 - 5.2 g/dL GLOBULIN 3.0 2.0 - 4.0 g/dL

1.4

Neonates -Pre Mature:

0.29 - 1.041.00 - 2.00

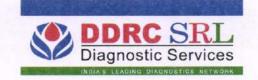
RATIO





mg/dL





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Test Report Status Preliminary	Results		Units
ASPARTATE AMINOTRANSFERASI (AST/SGOT)	E 20	Adults : < 33	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT) METHOD: IFCC WITHOUT PDP	16	Adults : < 34	U/L
ALKALINE PHOSPHATASE	89	Adult (<60yrs): 35 - 105	U/L
GAMMA GLUTAMYL TRANSFERAS TOTAL PROTEIN, SERUM	E (GGT) 9	Adult (female) : < 40	U/L
FOTAL PROTEIN METHOD: BIURET JRIC ACID, SERUM	7.1	Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
JRIC ACID METHOD: SPECTROPHOTOMETRY	4.1	Adults: 2.4-5.7	mg/dL
BO GROUP & RH TYPE, EDTA WHOLI	E BLOOD		
ABO GROUP METHOD : GEL CARD METHOD	0		
RH TYPE	POSITIVE		
LOOD COUNTS, EDTA WHOLE BLOOD			
HEMOGLOBIN METHOD : NON CYANMETHEMOGLOBIN	12.3	12.0 - 15.0	g/dL
ED BLOOD CELL COUNT METHOD: IMPEDANCE	4.29	3.8 - 4.8	mil/µL
VHITE BLOOD CELL COUNT METHOD: IMPEDANCE	6.14	4.0 - 10.0	thou/µL
LATELET COUNT METHOD: IMPEDANCE	316	150 - 410	thou/µL
BC AND PLATELET INDICES			
EMATOCRIT METHOD : CALCULATED	36.5	36 - 46	%
IEAN CORPUSCULAR VOL METHOD: DERIVED FROM IMPEDANCE MEASURE	85.0	83 - 101	fL
IEAN CORPUSCULAR HGB.	28.6	27.0 - 32.0	pg









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MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION METHOD: CALCULATED	33.6		31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH MENTZER INDEX	15.1 19.8		12.0 - 18.0	%
MEAN PLATELET VOLUME METHOD: DERIVED FROM IMPEDANCE MEASURE	7.1		6.8 - 10.9	fL
WBC DIFFERENTIAL COUNT				
SEGMENTED NEUTROPHILS METHOD: DHSS FLOWCYTOMETRY	59		40 - 80	%
LYMPHOCYTES METHOD: DHSS FLOWCYTOMETRY	33		20 - 40	%
MONOCYTES METHOD: DHSS FLOWCYTOMETRY	6		2 - 10	%
EOSINOPHILS METHOD: DHSS FLOWCYTOMETRY	2		1 - 6	%
BASOPHILS METHOD: IMPEDANCE	0		0 - 2	%
ABSOLUTE NEUTROPHIL COUNT	3.62		2.0 - 7.0	thou/µL
ABSOLUTE LYMPHOCYTE COUNT	2.03		1 - 3	thou/µL
ABSOLUTE MONOCYTE COUNT	0.37		0.20 - 1.00	thou/µL
ABSOLUTE EOSINOPHIL COUNT	0.12		0.02 - 0.50	thou/µL
ABSOLUTE BASOPHIL COUNT	0.00		0.00 - 0.10	thou/µL
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.8			
ERYTHROCYTE SEDIMENTATION RATE (ESR),WBLOOD	HOLE			
SEDIMENTATION RATE (ESR) METHOD: WESTERGREN METHOD	24	High	0 - 20	mm at 1 hr
* SUGAR URINE - POST PRANDIAL				
SUGAR URINE - POST PRANDIAL CYTOLOGY - CS (PAP SMEAR)	NOT DETECTED RESULT PENDING		NOT DETECTED	
THYROID PANEL, SERUM				
T3	105.10		80 - 200	ng/dL
Τ4	8.18		5.1 - 14.1	µg/dl



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TSH 3RD GENERATION

2.170

Non-Pregnant: 0.4-4.2

µIU/mL

Pregnant Trimester-wise :

1st : 0.1 - 2.5 2nd: 0.2 - 3 3rd: 0.3 - 3

Interpretation(s)

Triiodothyronine T3, Thyroxine T4, and Thyroid Stimulating Hormone TSH are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate. Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hyperthyroidism, TSH levels are low. Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3. Measurement of the serum TT3 level is a more sensitive test for the diagnosis of hyperthyroidism, and measurement of TT4 is more useful in the diagnosis of hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active. It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.

Sr. No.	TSH	Total T4	FT4	Total T3	Possible Conditions
1	High	Low	Low	Low	(1) Primary Hypothyroidism (2) Chronic autoimmune Thyroiditis (3) Post Thyroidectomy (4) Post Radio-Iodine treatment
2	High	Normal	Normal	Normal	(1)Subclinical Hypothyroidism (2) Patient with insufficient thyroid hormone replacement therapy (3) In cases of Autoimmune/Hashimoto thyroiditis (4). Isolated increase in TSH levels can be due to Subclinical inflammation, drugs like amphetamines, Iodine containing drug and dopamine antagonist e.g. domperidone and other physiological reasons.
3	Normal/Low	Low	Low	Low	(1) Secondary and Tertiary Hypothyroidism
4	Low	High	High	High	(1) Primary Hyperthyroidism (Graves Disease) (2) Multinodular Goitre (3)Toxic Nodular Goitre (4) Thyroiditis (5) Over treatment of thyroid hormone (6) Drug effect e.g. Glucocorticoids, dopamine, T4 replacement therapy (7) First trimester of Pregnancy
5	Low	Normal	Normal	Normal	(1) Subclinical Hyperthyroidism
6	High	High	High	High	(1) TSH secreting pituitary adenoma (2) TRH secreting tumor
7	Low	Low	Low	Low	(1) Central Hypothyroidism (2) Euthyroid sick syndrome (3) Recent treatment for Hypothyroidism
8	Normal/Low	Normal	Normal	High	(1) T3 thyrotoxicosis (2) Non-Thyroidal illness
9	Low	High	High	Normal	(1) T4 Ingestion (2) Thyroiditis (3) Interfering Anti TPO antibodies

REF: 1. TIETZ Fundamentals of Clinical chemistry 2. Guidlines of the American Thyroid association during pregnancy and Postpartum, 2011. NOTE: It is advisable to detect Free T3, Free T4 along with TSH, instead of testing for albumin bound Total T3, Total T4. TSH is not affected by variation in thyroid - binding protein. TSH has a diurnal rhythm, with peaks at 2:00 - 4:00 a.m. And troughs at 5:00 - 6:00 p.m. With ultradian variations.



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Results

REPORTED: 26/02/2023 12:47

REFERRING DOCTOR: DR. BOB

Test Report Status Preliminary

CLIENT PATIENT ID :

	- Todato		Units
* SUGAR URINE - FASTING			
SUGAR URINE - FASTING PHYSICAL EXAMINATION, URINE	NOT DETECTED	NOT DETECTED	
COLOR	AMBER		
APPEARANCE	CLEAR		
CHEMICAL EXAMINATION, URINE			
PH	5.0	4.8 - 7.4	
SPECIFIC GRAVITY	1.020	1.015 - 1.030	
PROTEIN	NOT DETECTED	NOT DETECTED	
GLUCOSE	NOT DETECTED	NOT DETECTED	
KETONES	NOT DETECTED	NOT DETECTED	
BLOOD	NOT DETECTED	NOT DETECTED	
BILIRUBIN	NOT DETECTED	NOT DETECTED	
UROBILINOGEN	NORMAL	NORMAL	
NITRITE	NOT DETECTED	NOT DETECTED	
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS	0 - 1	NOT DETECTED	/HPF
WBC	2-3	0-5	/HPF
EPITHELIAL CELLS	5-7	0-5	/HPF
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		
BACTERIA	NOT DETECTED	NOT DETECTED	
YEAST	NOT DETECTED	NOT DETECTED	









CLIENT CODE: CA00008282 - BANK OF BARODA CLIENT'S NAME AND ADDRESST:

BANK OF BARODA - MEDI ASSIST MEDI ASSIST HEALTHCARE SERVICE TOWER D, 4TH FLOOR, IBC KNOWLEDGE PARK, 4/1 BANNERGHATTA ROAD, BHAVANI NAGA BENGALURU BENGALURU 560029 KARNATAKA INDIA

Cert. No. MC-2354

DDRC SRL DIAGNOSTICS DDRC SRL Tower, G-131,Panampilly Nagar, PANAMPALLY NAGAR, 682036 KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: G.SMITHA

PATIENT ID: GSMIF2502804126

ACCESSION NO: 4126WB009076 AGE: 43 Years

SEX: Female

ABHA NO:

DRAWN:

9742187722

RECEIVED: 25/02/2023 08:54

26/02/2023 12:47 REPORTED:

REFERRING DOCTOR: DR. BOB

CLIENT PATIENT ID :

Test Report Status

Preliminary

Results

Units

Interpretation(s)

The following table describes the probable conditions, in which the analytes are present in urine

Presence of	Conditions
Proteins	Inflammation or immune illnesses
Pus (White Blood Cells)	Urinary tract infection, urinary tract or kidney stone, tumors or any kind of kidney impairment
Glucose	Diabetes or kidney disease
Ketones	Diabetic ketoacidosis (DKA), starvation or thirst
Urobilinogen	Liver disease such as hepatitis or cirrhosis
Blood	Renal or genital disorders/trauma
Bilirubin	Liver disease
Erythrocytes	Urological diseases (e.g. kidney and bladder cancer, urolithiasis), urinary tract infection and glomerular diseases
Leukocytes	Urinary tract infection, glomerulonephritis, interstitial nephritis either acute or chronic, polycystic kidney disease, urolithiasis, contamination by genital secretions
Epithelial cells	Urolithiasis, bladder carcinoma or hydronephrosis, ureteric stents or bladder catheters for prolonged periods of time
Granular Casts	Low intratubular pH, high urine osmolality and sodium concentration, interaction with Bence-Jones protein
Hyaline casts	Physical stress, fever, dehydration, acute congestive heart failure, renal diseases
Calcium oxalate	Metabolic stone disease, primary or secondary hyperoxaluria, intravenous infusion of large doses of vitamin C, the use of vasodilator naftidrofuryl oxalate or the gastrointestinal lipase inhibitor orlistat, ingestion of ethylene glycol or of star fruit (Averrhoa carambola) or its juice
Uric acid	arthritis
Bacteria	Urinary infectionwhen present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis

GLUCOSE FASTING, FLUORIDE PLASMA

GLUCOSE, FASTING, PLASMA

98

Diabetes Mellitus : > or = 126.

Impaired fasting Glucose/ Prediabetes: 101 - 125. Hypoglycemia : < 55.

* PHYSICAL EXAMINATION, STOOL

RESULT PENDING

* CHEMICAL EXAMINATION, STOOL

RESULT PENDING









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Test Report Status

Preliminary

Results

Units

* MICROSCOPIC EXAMINATION, STOOL

RESULT PENDING



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CIN: U85190MH2006PTC161480







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Test Report Status

Preliminary

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Units

Interpretation(s)

Stool routine analysis is only a screening test for disorders of gastrointentestinal tract like infection, malabsorption, etc. The following table describes the probable conditions, in which the analytes are present in stool.

PRESENCE OF	CONDITION
Pus cells	Pus in the stool is an indication of infection
Red Blood cells	Parasitic or bacterial infection or an inflammatory bowel condition such as ulcerative colitis
Parasites	Infection of the digestive system. Stool examination for ova and parasite detects presence of parasitic infestation of gastrointestinal tract. Various forms of parasite that can be detected include cyst, trophozoite and larvae. One negative result does not rule out the possibility of parasitic infestation. Intermittent shedding of parasites warrants examinations of multiple specimens tested on consecutive days. Stool specimens for parasitic examination should be collected before initiation of antidiarrheal therapy or antiparasitic therapy. This test does not detect presence of opportunistic parasites like Cyclospora, Cryptosporidia and Isospora species. Examination of Ova and Parasite has been carried out by direct and concentration techniques.
Mucus	Mucus is a protective layer that lubricates, protects& reduces damage due to bacteria or viruses.
Charcot-Leyden crystal	Parasitic diseases.
Ova & cyst	Ova & cyst indicate parasitic infestation of intestine.
Frank blood	Bleeding in the rectum or colon.
Occult blood	Occult blood indicates upper GI bleeding.
Macrophages	Macrophages in stool are an indication of infection as they are protective cells.
Epithelial cells	Epithelial cells that normally line the body surface and internal organs show up in stool when there is inflammation or infection.
Fat	Increased fat in stool maybe seen in conditions like diarrhoea or malabsorption.
pH	Normal stool pH is slightly acidic to neutral. Breast-fed babies generally have an acidic stool.

ADDITIONAL STOOL TESTS:

- Stool Culture: This test is done to find cause of GI infection, make decision about best treatment for GI infection & to find out if treatment for GI infection worked.
- Fecal Calprotectin: It is a marker of intestinal inflammation. This test is done to differentiate Inflammatory Bowel Disease (IBD) from Irritable Bowel Syndrome (IBS).
- 3. Fecal Occult Blood Test(FOBT): This test is done to screen for colon cancer & to evaluate possible cause of unexplained anaemia.
- Clostridium Difficile Toxin Assay: This test is strongly recommended in healthcare associated bloody or waterydiarrhoea, due to
 overuse of broad spectrum antibiotics which alter the normal GI flora.
- Biofire (Film Array) GI PANEL: In patients of Diarrhoea, Dysentry, Rice watery Stool, FDA approved, Biofire Film Array
 Test, (Real Time Multiplex PCR) is strongly recommended as it identifies organisms, bacteria, fungi, virus parasite and other
 opportunistic pathogens, Vibrio cholera infections only in 3 hours. Sensitivity 96% & Specificity 99%.

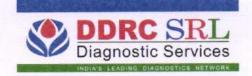


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CIN: U85190MH2006PTC161480







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Rota Virus Immunoassay: This test is recommended in severe gastroenteritis in infants & children associated with watery diarrhoea, vomitting& abdominal cramps. Adults are also affected. It is highly contagious in nature.

ELOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Mailgnancy, Nephrolithiasis, Prostatism)

Causes of decreased level include Liver disease, SIADH.

CREATININE, SERUM-Higher than normal level may be due to:

Blockage in the urinary tract

Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
 Loss of body fluid (dehydration)

Muscle problems, such as breakdown of muscle fibers
 Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

Lower tran formal level may be due to.
 Myasthenia Gravis
 Muscular dystrophy
 GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.Additional test HbA1c GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-Used For:

1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.

1.Evaluating the long-term control of blood glucose concentrations in diabetic patients.

2.Diagnosing diabetes.

3.Identifying patients at increased risk for diabetes (prediabetes).

The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range.

1.eAG (Estimated average glucose) converts percentage HbA1c to md/dl, to compare blood glucose levels.

2. eAG gives an evaluation of blood glucose levels for the last couple of months.

3. eAG is calculated as eAG (mg/dl) = 28.7 * HbA1c - 46.7

HbA1c Estimation can get affected due to:

I.Shortened Erythrocyte survival: Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss, hemolytic anemia) will falsely lower HbA1c test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.

II.Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin.

III.Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results.

IV.Interference of hemoglobinopathies in HbA1c estimation is seen in a. Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.

b.Heterozygous state detected (D10 is corrected for HbS & HbC trait.)

Antimotygous state detected (D10 is corrected for HbS & HbC trait.)

C.HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

TOTAL PROTEIN, SERUM-Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum...Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom'''''''s disease Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc.

URIC ACID, SERUM-Causes of Increased levels:-Dietary(High Protein Intake, Prolonged Fasting, Rapid weight loss), Gout, Lesch nyhan syndrome, Type 2 DM, Metabolic

syndrome

syndrome

Causes of decreased levels-Low Zinc intake,OCP,Multiple Sclerosis

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD
Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.



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Preliminary

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Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.

BLOOD COUNTS,EDTA WHOLE BLOOD-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13)

RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen book to differentiate to the displacement of the several patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.

(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION:

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic; it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition.CRP is superior to ESR because it is more sensitive and reflects a more rapid change.

TEST INTERPRETATION

Increase in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.

Finding a very accelerated ESR(>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).

In pregnancy BRI in first trimester is 0-48 mm/hr(62 if anemic) and in second trimester (0-70 mm/hr(95 if anemic). ESR returns to normal 4th week post partum.

Decreased in: Polycythermia vera, Sickle cell anemia

False elevated ESR: Increased fibrinogen, Drugs(Vitamin A, Dextran etc.), Hypercholesterolemia
False Decreased: Poikilocytosis, (SickleCells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs(Quinine, salicylates)

I. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin; 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition.

SUGAR URINE - POST PRANDIAL-METHOD: DIPSTICK/BENEDICT'S TEST
SUGAR URINE - FASTING-METHOD: DIPSTICK/BENEDICT'S TEST
GLUCOSE FASTING, FLUORIDE PLASMA-TEST DESCRIPTION

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and sothat no glucose is excreted in the

Increased in

Diabetes mellitus, Cushing's syndrome (10 - 15%), chronic pancreatitis (30%). Drugs:corticosteroids, phenytoin, estrogen, thiazides.

Decreased in Pancreatic islet cell disease with increased insulin, insulinoma, adrenocortical insufficiency, hypopituitarism, diffuse liver disease, malignancy (adrenocortical, stomach, fibrosarcoma), infant of a diabetic mother, enzyme deficiency diseases(e.g., galactosemia), Drugs- insulin, ethanol, propranolol; sulfonylureas, tolbutamide, and other oral hypoglycemic agents.

NOTE: While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values), there is wide fluctuation within individuals. Thus, glycosylated hemoglobin(HbALc) levels are favored to monitor glycemic control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

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Units

MEDIWHEEL HEALTH CHECKUP ABOVE 40(F)TMT

* ECG WITH REPORT

REPORT

TEST COMPLETEDY

* MAMMOGRAPHY -BOTH

REPORT

TEST COMPLETED

* USG ABDOMEN AND PELVIS

TEST COMPLETED

* CHEST X-RAY WITH REPORT

REPORT

TEST COMPLETED

End Of Report

Please visit www.srlworld.com for related Test Information for this accession TEST MARKED WITH '*' ARE OUTSIDE THE NABL ACCREDITED SCOPE OF THE LABORATORY.

DR.HARI SHANKAR, MBBS MD (Reg No - TCMC:62092) **HEAD - Biochemistry &** Immunology

DR.VIJAY K N,MBBS MD(PATH) (Reg No - KMC:91816) **HEAD-HAEMATOLOGY &** CLINICAL PATHOLOGY

DR.SMITHA PAULSON, MD (PATH), DPB (Reg No - TCMC:35960) LAB DIRECTOR & HEAD-HISTOPATHOLOGY & CYTOLOGY

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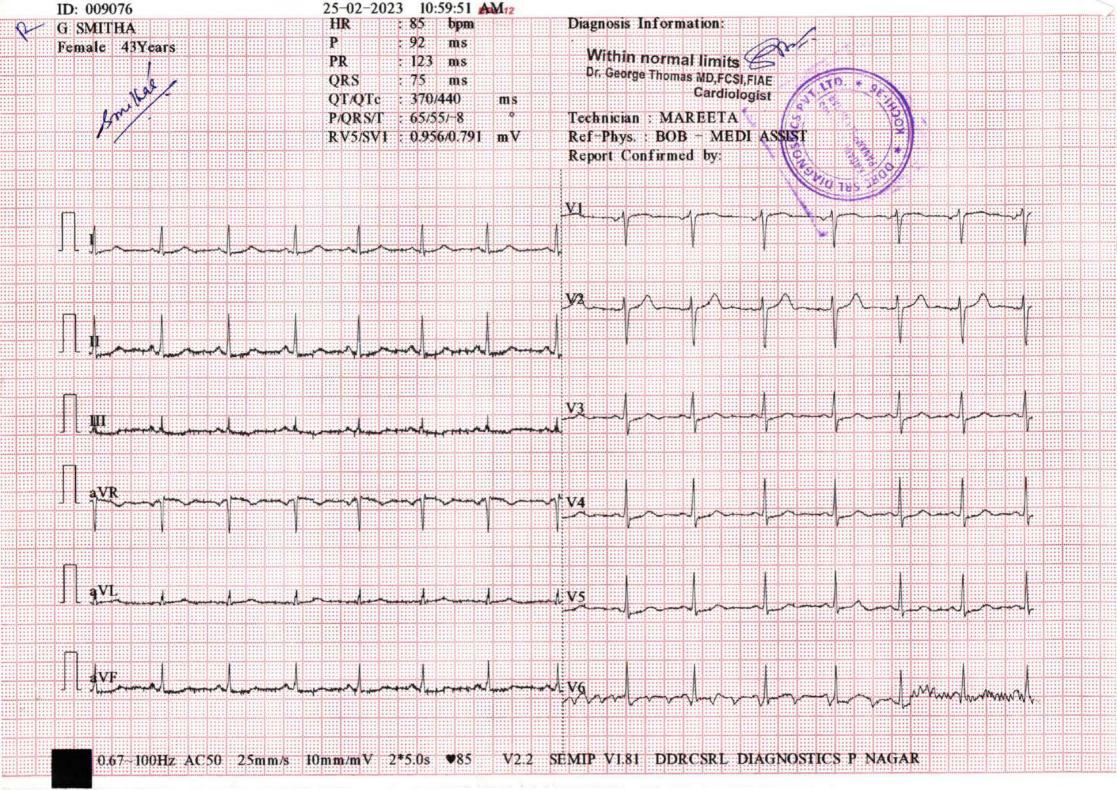
CIN: U85190MH2006PTC161480



Date. 25.02. 2023

OPHTHALMOLOGY REPORT

This is to certif	y that I have examined
Mr / Ms :	miMaAged
visual standard	s is as follows:
Visual Acuity:	
	R:
For far vision	ւ։616
For near vision	R: No
roi near vision	L:
Color Vision :	Normal
	Nannu Elizabeth (Optometrist)





NAME: MRS G SMITHA	STUDY DATE 25/02/2023
AGE / SEX : 43 YRS / F	REPORTING DATE 25/02/2023
REFERRED BY :BOB	ACC NO: 4126WB009076

X - RAY - CHEST PA VIEW

- Both the lung fields are clear.
- B/L hila and mediastinal shadows are normal.
- Cardiac silhouette appears normal.
- Cardio thoracic ratio is normal.
- Bilateral CP angles and domes of diaphragm appear normal.

IMPRESSION: NORMAL STUDY

Kindly correlate clinically

Dr. NAVNEET KAUR, MBBS,MD Consultant Radiologist.

CIN: U85190MH2006PTC161480
(Refer to "CONDITIONS OF REPORTING" Overleaf)





NAME	MRS G SMITHA	AGE	43 YRS
SEX	FEMALE	DATE	February 25, 2023
REFERRAL	BANK OF BARODA	ACC NO	4126WB009076

USG BOTH BREASTS

Clinical details: Screening

Real Time gray scale B mode imaging of breasts reveals:

RIGHT BREAST

- Normal in Echotexture.
- Retroareolar region normal.
- · Nipple and areola appears normal.
- Axillary tail normal.
- No evidence of any regional lymphadenopathy

LEFT BREAST

- · Normal in Echotexture.
- · Retroareolar region normal.
- · Nipple and areola appears normal.
- · Axillary tail normal.
- · No evidence of any regional lymphadenopathy

IMPRESSION:

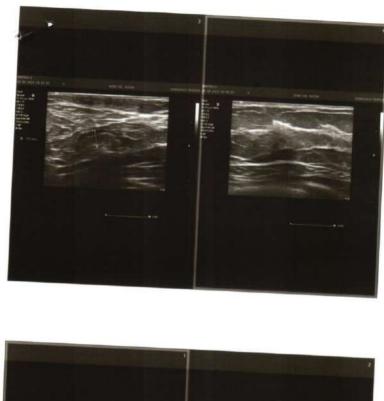
No significant abnormality detected (BIRADS I).

Dr. NAVNEET KAUR MBBS . MD Consultant Radiologist

Thank you for referral. Your feedback will be appreciated.

NOTE: This report is only a professional opinion based on the real time image finding and not a diagnosis by itself. It has to be correlated and interpreted with clinical and other investigation findings. Review scan is advised, If this ultrasound opinion and other clinical findings / reports don't correlate.

CIN: U85190MH2006PTC161480







NAME	MRS G SMITHA	AGE	43 YRS
SEX	FEMALE	DATE	February 25, 2023
REFERRAL	BANK OF BARODA	ACC NO	4126WB009076

USG ABDOMEN AND PELVIS

LIVER

Measures ~ 11.5 cm. Bright echotexture.

Smooth margins and no obvious focal lesion within. No IHBR dilatation. Portal vein normal in caliber.

GB

Contracted.

SPLEEN

Measures ~ 8.5 cm, normal to visualized extent. Splenic vein normal.

PANCREAS

Normal to visualized extent, PD is not dilated.

KIDNEYS

RK: 8.7 x 3.6 cm, appears normal in size and echotexture. LK: 8.8 x 5.1 cm, appears normal in size and echotexture.

No focal lesion / calculus within.

Maintained corticomedullary differentiation and normal parenchymal thickness.

No hydroureteronephrosis.

BLADDER

Normal wall caliber, no internal echoes/calculus within.

UTERUS

Anteverted, normal in size [6.1 x 3.6 x 5.7 cm] and echopattern.

No focal lesion seen.

ET - 8.7 mm.

OVARIES

RT OV: $2.6 \times 2 \times 2.1 \text{ cm}$ [volume $\sim 6 \text{ cc}$]. LT OV: $2.7 \times 1.3 \times 2.6 \text{ cm}$ [volume $\sim 5 \text{ cc}$].

NODES/FLUID

Nil to visualized extent.

BOWEL

Visualized bowel loops appear normal.

IMPRESSION

♣ Grade I fatty liver.

Kindly correlate clinically.

Dr. NAVNEET KAUR MBBS . MD Consultant Radiologist

Thank you for referral. Your feedback will be appreciated.

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