

(Multi Super Speciality 200 Bedded Hospital)

DEPARTMENT OF PATHOLOGY

UHID

Name Age/Gender

Accession Number

Acknowledge Date/Time

Payer Name

CIMS, 7937

Mr Rajendra Singh

35 Y/Male

OPAC-2326

22-01-2024 01:29 PM

Mediwheel Full Body Health

Checkup

Visit Type/No

Order No

Order Date/Time

Collection Date/Time Ordering Doctor

Refer By

OP/EPD-10163 OR-17503

22-01-2024 12:01 PM

Dr Self

22-01-2024

Pathology

		Latitology		
Service Name	Result	Unit	D.f. D	
URINE ANALYSIS/ URINE ROUTINE	E EXAMINATION 1	leine.	Reference Range	Method
Physical Examination		orme		
COLOUR	Pale Yellow			
TRANSPARENCY	Clear			Manual method
SPECIFIC GRAVITY	1.010			Manual
PH URINE	7.0		1.001-1.03	Strip
DEPOSIT	Absent		5-8	Strip
BIOCHEMICAL EXAMINATION	- ADSOIL			Manual
ALBUMIN	Absent			
SUGAR	Absent			Strip
BILE SALTS (BS)	Absent			Strip
BILE PIGMENT (BP)	Absent			Manual
MICROSCOPIC EXAMINATION	rusem			Manual
PUS CELLS	0-1	/ burf		
EPITHELIAL CELLS	0-1	/ hpf		Microscopy
RBC'S	Absent	/ hpf /hpf		Microscopy
CASTS	Absent	, npr		Microscopy
CRYSTALS	Absent			Microscopy
BACTERIA	Absent			Macroscopy
FUNGUS	Absent			Macroscopy
SPERMATOZOA	Absent			Microscopy
OTHERS	Absent			Microscopy
Thyroid Profile -T3, T4, TSH, Blood	Ausen			Microscopy
Triiodothyronine (T3)	1.69			H222400500445
Thyroxine (T4)	88.3	ng/mL	0.69-2.15	CLIA
Thyroid Stimulating Hormone (TSH)	2.10	ng/mL	52-127	CLIA
	2.10	uIU/mL	0.3-4.5	CLIA
Interpretation				

Interpretation

- 1. TSH levels are subject to circadian variation, reaching peak levels between 2 4.a.m. and at a minimum between 6-10 pm . The variation is of the order of 50% . hence time of the day has influence on the measured serum TSH
- 2. Recommended test for T3 and T4 is unbound fraction or free levels as it is metabolically active.
- 3. Physiological rise in Total T3 / T4 levels is seen in pregnancy and in patients on steroid therapy. Clinical Use

Primary Hypothyroidism

Hyperthyroidism Hypothalamic - Pituitary hypothyroidism

Inappropriate TSH secretion

Nonthyroidal illness

Autoimmune thyroid disease

Pregnancy associated thyroid disorders

Thyroid dysfunction in infancy and early childhood

Haematology

All tests have technical limitations Corroborative clinicopathological interpretation is indicated. In case of any disparity in including machine error or typing the test should be repeated immediately.

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(Multi Super Speciality 200 Bedded Hospital)

DEPARTMENT OF RADIOLOGY

NAME: MR. RAJENDRA SINGH	AGE: 35 YRS.	SEX: M
REF. BY: DR. SELF	UHID: 7937	DATE: 22-Jan-2024

ULTRASOUND SCAN OF ABDOMEN

FINDINGS:

Liver is mildly enlarged in size measuring ~ 15.9 cm. Echotexture is slightly echogenic. No focal space occupying lesion is seen within liver parenchyma. Intrahepatic biliary channel are not dilated. Portal vein is normal in caliber.

Gall bladder wall is not thickened. No calculus or mass lesion is seen in gall bladder. Common bile duct is not dilated.

Pancreas is of normal in size and contour. Echo-pattern is normal. No focal lesion is seen within pancreas. (Only head & proximal body is visualized)

Spleen is normal in size (10.4 cm). Echotexture is normal. No focal Lesion is seen.

Right kidney is normally sited and is of normal size (RT \sim 10.4 x 4.9 cm) and shape. Cortico medullary echoes are normal. No focal mass lesion is seen.

Right VUJ shows a calculus measuring approx. 11.5 mm causing mild hydroureteronephrosis.

Left kidney is normally sited and is of normal size (LT \sim 10.6 x 4.6 cm) and shape. Cortico medullary echoes are normal. No focal mass lesion is seen. Collecting system does not show any calculus.

Urinary bladder is normal in distension and wall is not thickened. No calculi seen.

Prostate is normal in size and normal in echotexture.

No free fluid seen in peritoneal cavity.

IMPRESSION-

- > RIGHT VUJ CALCULUS CAUSING MILD PROXIMAL HYDROURETERONEPHROSIS.
- > MILD HEPATOMEGALY WITH GRADE I FATTY CHANGES IN LIVER.

PLEASE CORRELATE CLINICALLY & F/E.

DR. ABHAKRAINA M.B.B.S., D.N.B (RADIO-DIAGNOSIS) CONSULTANT RADIOLOGIST

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Dr Self

Service Name

Result

Unit

Reference Range

Method

BLOOD GROUP (ABO) BLOOD GROUP (ABO)-

RH TYPING

"B"

Positive The upper agglutination test for grouping has some limitations.

HbA1c

GLYCOSYLATED HAEMOGLOBIN (HbA1c)

Method- Immunofluorescence Assay

Glycosylated Hemoglobin (HbA1c)

6.27

<6.5: Non Diabetic

6.5-7 : Good Control 7-8: Weak Control

> 8 : Poor Control 90-120: Excellent Control

Estimated average blood glucose (eAG) 133.2 mg/dl

121-150: Good Control

151-180: Average Control 181-210: Action Suggested

Note:

1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled.

2. Target goals of 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targeting a goal of 7.0 % may not be appropriate. Comments:

HbA1c provides an index of average blood glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.

ESR (Erythrocyte Sedimentation Rate), Blood CBC (Complete Blood Count), Blood	14 H	mm 1st Hr.	0-10	Wintrobe
Hemoglobin (Hb) TLC (Total Leukocyte Count) DIFFERENTIAL LEUCOCYTE COUNT	16.7 7180	gm/dl /cumm	13-17 4000-11000	Spectrophotometry Impedance
Neutrophils Lymphocytes Monocytes Eosinophils Basophils RBC Count PCV / HCt (Hematocrit) MCV	42 48 H 06 04 00 4.95 49.0 H 99.1 H	% % % millions/cumm %	40-80 20-45 1-8 1-6 0-1 4.5-6.0 40-45 76-96	Calculated to NH-10 Manual Calculated to NH-10 M
MCH	33.7 H	pg	27-32	* 100

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Dr Self

Service Name	Result	Unit	Reference Range	Method
MCHC	34.0	g/dL	30-35	Method
Platelet Count	2.99	lakh/eumm	1.5-4.5	Impedance
RDW	12.7	%	1-15	Impedance

Clinical Biochemistry

		Clinical Biocher	mistry	
Service Name	Result	Unit	Reference Range	Method
Glucose (Post Prandial), Plasma Lipid Profile, Serum	119.0	mg/dL	80-150	Method
Cholestrol, serum	193.0	mg%	Optimal: < 200 mg/dl Boder Une High Risk: 150 -240 mg/dl High Risk: > 250 mg/dl	
Triglycerides, serum	125.0	mg%	Optimal: < 150 mg/dl Border Line High Risk: 150 - 199 mg/dl High Risk: 200 - 499 mg /dl Very High Risk: > 500 mg /dl	
IDL Cholesterol	53.2	mg%	Optimal: 70 mg/dl Border Line High Risk: 80 - 100 mg/dl High Risk: > 120 mg/dl	
.DL Cholesterol	114.80	mg%	Optimal: < 100 mg/dl Border Line High Risk: 100 - 129 mg/dl High Risk: > 160 mg/dl	
VLDL Cholestrol	25.00	mg%	Male : 10 - 40 mg/dl Female : 10 - 40 mg/dl	
.DL / HDL Cholesterol ratio	2.16		Child: 10 - 40 mg/dl 0.0-3.5	
ptomust-ti				

Interpretation

- 1. Measurements in the same patient can show physiological & analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycendes, HDL & LDL Cholesterol.
- ATP III recommends a complete lipoprotein profile as the initial test for evaluating cholesterol.
- 3. Friedewald equation to calculate LDL cholesterol is most accurate when Triglyceride fewel is < 400 mg/dL. Measuren LDI, cholestorol is recommended when Triglyceride level is > 400 mg/dL

LFT (Liver Function Test	Profile, Serum
Bilirubin Total, Serum	
Conjugated (Direct), Serum	

Unconjugated (Indirect) SGOT/AST

SGPT/ALT

0.91 0.33 H mg% 0.58 mg% 43.0 H U/L 27.0 U/L

0.1-1.0 0.0-0.3 0.0 - 0.750-40 0 - 48

DMSO DMSO IFCC

Calculated **IFCC**

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Service Name AST/ALT Ratio	Result	Unit	Reference Range	Method
Gamma GT.Serum	1.59 H		0-1	Calculated
Alkaline phosphatase, Serum	46.0 H	U/L	10-45	IFCC
Total Protein, scrum	180.0 H	U/L	53-165	IFCC
Albumin, Serum	7.89	gm/dl	6.0-8.4	Biuret
Globulin	4.27	g/dL	3.5-5.4	BCG
A/G Ratio	3.62 H	g/dL	2.3-3.6	Calculated
KFT (Kidney Profile) -II, Serum	1.18		1.0-2.3	Calculated
Urea, Blood	20.0			
Creatinine, Serum	29.0	mg/dL	15-50	Urease-uv
Blood Urea Nitrogen (BUN)	0.72	mg/dL	0.6-1.2	Enzymatic
BUN-CREATININE RATIO	13.53	mg%	7.5-22.0	Calculated
Sodium, Serum	18.80		10-20	Calculated
Potassium, Serum	136.3	mmol/L	135-150	ISE
Calcium, Serum	4.25	mmol/L	3.5-5.5	ISE
Chloride, Serum	9.76	mg/dL	8.7-11.0	ISE
Jric acid, Scrum	107.2	mmol/L	94-110	ISE
Magnesium, Serum	6.27	mg/dL	3.4-7.0	
Phosphorus, Serum	1.88	ing/dL	1.6-2.8	XYLIDYL BLUE
Alkaline phosphatase, Serum	3.52	mg/dL	2.4-5.0	MOLYBDATE U
Albumin, Serum	180.0 H	U/L	53-165	IFCC
GFT (Kidney Profile) -I, Serum	4.27	g/dL	3.5-5.4	BCG
rea, Blood				BCO
reatinine, Serum	74	mg/dL	15-50	Urease-uv
Blood Urea Nitrogen (BUN)		mg/dL	0.6-1.2	Enzymatic
UN-CREATININE RATIO	- 0	mg%	7.5-22.0	Calculated
odium.Scrum	14.5		10-20	Calculated
otassium, Serum	200	mmol/L	135-150	ISE
alcium, Serum		mmol/L	3.5-5.5	ISE
hloride, Serum		mg/dL	8.7-11.0	ISE
ric acid, Serum	- 12	mmol/L	94-110	ISE
lagnesium, Serum		mg/dL	3.4-7.0	131.
nosphorus, Serum		mg/dL	1.6-2.8	VVI IDVI DI US
lkaline phosphatase, Serum		mg/dL	2.4-5.0	XYLIDYL BLUE
bumin, Serum		U/L	53-165	MOLYBDATE UV
		g/dL	3.5-5.4	IFCC
ucose (Fasting), Plasma	94.0	mg/dL	60-110	BCG



-----End of the Report-----

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NOT VALID FOR MEDICO LEGAL PURPOSE.

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Dr Ambrish Kumar Pathology MD (Pathology)



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