



(Govt. Recognised Advanced 3D-4D Ultrasound and State-of-the-art Digital X-Ray Centre) Regn. No.: RMEE1905483

80 (Near Nagar Nigam Office), Vasundhara, Ghaziabad, Phone: 0120-4127778, 9899004884, E-mail : premdharamhospital@ymail.com Name : Mr. Karan Kakkar Patient ID : 24/230300015

Received On

Patient ID

Visit No. : SR242303015

: 23/03/2024 14:24 Age/Gender : 43 Y/Male Collected On : 23/03/2024 14:24 Referred by : PREM-DHARAM HOSPITAL

Reported On : 23/03/2024 19:35 : PDH42291A Barcode

HAEMATOLOGY COMPLETE BLOOD COUNT WITH ESR **Test Name Obtained Value** Units Bio. Ref. Intervals(Age/Gender specific)

			00	Bio. Nei. Intervals(Age/Gender specific)
	HAEMOGLOBIN Methodology: Colorimetric	14.7	g/dl	13.0- 17.5
	RED BLOOD CELL COUNT (RBC)	4.85	millions/mm ³	4.5 - 6
	Methodology : Electrical Impedence			a reserve of the second of
	PACKED CELL VOLUME/HEMATOCRIT (PCV)	42.8	% Vol	40 - 50
	Methodology : Calculated			
	MEAN CORPUSCULAR VOLUME (MCV) Methodology : Calculated	88.3	fL	80 - 96
	MEAN CORPUSCULAR HAEMOGLOBIN (MCH)	30.4	pg	27 - 33
	Methodology: Calculated			
	MEAN CORPUSCULAR HAEMOGLOBIN CONCENTRATION (MCHC)	34.4	g/dl	31 - 36
	Methodology: Calculated			
	RED CELL DISTRIBUTION WIDTH (RDW-CV)	15.9	%	11 - 16
	Methodology : Automated-Cell Counter			
	RED CELL DISTRIBUTION WIDTH (RDW-SD)	46.8	fL	35 - 56
	Methodology : Automated-Cell Counter			$\tilde{\epsilon}_{\odot}$
	TOTAL LEUCOCYTE COUNT	7.41	10^3/µL	Ta and 1 = 1 = 1
	Methodology : Flow Cytometry	F 10 10 10 10 10 10 10 10 10 10 10 10 10	10 3/μι	4 - 11
	DIFFERENTIAL COUNT (DC)			
1	NEUTROPHILS .	61	%	40. 75
l	LYMPHOCYTES	32	%	40 - 75
E	EOSINOPHILS	03	%	20 - 45
١	MONOCYTES	04		0 - 6
	BASOPHILS		%	0 - 10
		00	%	0 - 1
٨	ABSOLUTE NEUTROPHIL COUNT (ANC) Methodology: Calculated	4.53	10^3/μL	2 - 8
	ABSOLUTE LYMPHOCYTE COUNT (ALC)	2.38	10^3/μL	0.8 - 7
IV	lethodology · Calculated			

ABSOLUTE EOSINOPHIL COUNT (AEC) Methodology : Calculated

Methodology: Calculated

Note: If the test results are alarming or unexpected, Client is advised to contact the laboratory



0.02 - 0.8 3D/4D Ultrasound

Whole Body Color Doppler

Digital X-Ray (24 Hours)

immediately for possible remedial action graphy

ECG-3 Channel

FOR TELEPHONIC APPOINTMENT RADIOLOGY: 8287900395 (9 AM - 2:30 PM, 5:30 PM - 8:45 PM)





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			Barcode	: PDH42291A
ABSOLUTE MONOCYTE COUNT (AMC) Methodology: Calculated	0.28	10^3/μL	0.12 - 1.2	
ABSOLUTE BASOPHIL COUNT (ABC) Methodology: Calculated	00	10^3/µL	0 - 0.1	
PLATELET COUNT Methodology : Electrical Impedence	260	10^3/µL	150 - 450	
MEAN PLATELET VOLUME (MPV) Methodology: Electrical Impedence	8.7	fL	7 - 12	
PLATELET DISTRIBUTION WIDTH (PDW) Methodology: Calculated	16.4	 fL	9 - 17	
PCT(PLATELET CRIT) Methodology: Calculated	0.22	%	0.108 - 0.282	
P-LCR Methodology: Calculated	20.3	%	11 - 45	
P-LCC Methodology : Calculated	53	10?9/L	30 - 90	
ERYTHROCYTE SEDIMENTATION RATE (ESR)	04	mm in 1st hr	0 - 10	
Methodology: Westergreen				

Sample Type: Whole Blood-EDTA

Processed by : Nisha Sharma Verified by : Tabassum

*** End Of Report ***







- 3D/4D Ultrasound Whole Body Color Doppler
- Digital X-Ray (24 Hours) d, Client is advised to contact the laboratory immediately for possible remediataction og raphy
 - ECG-3 Channel

FOR OPD APPOINTMENT: 8287900395



PREM-DHARAM



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CLINICAL BIOCHEMISTRY

LIVER FUNCTION TEST

Test Name	Obtained Value	Units	Bio Ref Intervals	Age/Gender specific)
TOTAL BILIRUBIN	0.72	mg/dL	0.2 - 1.2	Age/Gender specific)
Methodology: Diazo Method		9,	0.2 - 1.2	
DIRECT BILIRUBIN	0.27	· mg/dL	0 - 0.3	
Methodology : Diazo Method		g, az	0 - 0.5	
INDIRECT BILIRUBIN	0.45	mg/dL		
Methodology: Calculated	W.	9, a2		
SGOT/AST	19.1	U/L	0 - 40	
Methodology : IFCC	V-45000000 10.	2,2	0 - 40	
SGPT/ALT .	23.7	U/L	0 - 35	
Methodology : IFCC			0-33	
ALKALINE PHOSPHATASE	112.6	U/L	.40 - 130	
Methodology: IFCC		-/-	,40 - 130	
TOTAL PROTEIN	7.98	g/dl	6 - 8.3	
Methodology : Biuret		g,	0 - 0.3	
SERUM ALBUMIN	4.24	g/dl	3.2 - 5.2	
Methodology : BCG		3,	. J.L - J.L	
GLOBULIN SERUM .	3.74	g/dl	2.3 - 4.5	
Methodology : Calculated		9, 01	2.3 - 4.5	
A/G RATIO	1.13	Ratio	1 - 2.5	
Methodology : Calculated		1,10,10	1 - 2.5	
Sample Type : serum				

Processed by : Nisha Sharma

Verified by : Tabassum

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CLINICAL BIOCHEMISTRY RENAL FUNCTION TEST

RENAL FUNCTION TEST					
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific)		
BLOOD UREA	26.1	mg/dL	10 - 45		
Methodology: Urease		3,			
BLOOD UREA NITROGEN (BUN)	12	mg/dL	5 - 21		
Methodology : Calculated		mg/ ac	3-21		
SERUM CREATININE	0.72	mg/dL	0.7 - 1.4		
Methodology : Jaffe Kinetic	6.5.	mg/ac	0.7 - 1.4		
SODIUM - SERUM	142.6	meg/L	135 - 155		
Methodology: ISE		meq/L	133 - 133		
POTASSIUM - SERUM	3.98	meq/L	3.5 - 5.5		
Methodology: ISE	S-00M-M/10	meq/L	3.3 - 3.3		
CHLORIDE - SERUM	104.5	mmol/L	98 - 106		
Methodology : ISE		oy L	30 - 100		
CALCIUM - SERUM	9.0	mg/dL	8.6 - 11		
Methodology : Arsenazo		mg/ at			
EGFR	153	mL/min/1.73 m2	00 400		
		111L/11111/1.73 M2	90 - 180 > = 90 : Normal		
			60 - 89 : Mild Decreas		
			45 - 59 : Mild to Moderate Decrease		
			30 - 44 : Moderate to Severe Decrease		
IDIC ACID CEDINA			15 - 29 : Severe Decrease		
URIC ACID - SERUM	4.28	mg/dL	3.5 - 7.2		
Methodology : URICASE					
Sample Type : serum					

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Consultant Pathologist









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CLINICAL BIOCHEMISTRY

LIPID	PROFILE	
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-		PID PROFILE		
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific)	
TOTAL CHOLESTEROL	266.3	mg/dL	1-200 Desirable < 200 Borderline high risk 200 - 240 High risk > 240	
Methodology.: CHO-POD				
Comments: KINDLY CORRELATE CLINIC	CALLY			
HDL CHOLESTEROL Methodology: Direct	60.	mg/dL	NO RISK : - > 60.0 MODERATE RISK :- 35 - 55 HIGH RISK : - < 35.0	
LDL CHOLESTEROL	203.90	mg/dL	0 - 130 Desirable < 130 Borderline high risk 130 -160	
Methodology : Calculated			'High risk > 160	
VLDL . Methodology: Calculated	2.40	mg/dL	0 - 45	
TRIGLYCERIDES (TG) - SERUM	121	mg/dL	0 - 200 Desirable: < 200 (fasting) Borderline high: 200 - 400	
Methodology : GPO-POD			Elevated > 400	
CHOL/HDL Ratio Methodology : Calculated	4.44	Ratio	3.5 - 5.5	
LDL/HDL Ratio Methodology : Calculated	3.40	mg/dL	2.5 - 3.5	
Sample Type : serum				

*** End Of Report ***

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Dr. Vivek Kapoor Consultant Pathologist







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UL	-IIN	ICAL	BIU	CHEN	/ISTRY

Test Name	Obtained Value Units		Bio. Ref. Intervals(Age/Gender specific)
PLASMA GLUCOSE FASTING (FBS)	95.9	mg/dL	70 - 110
24 14 14 14 15 15 15 15 15 15 15 15 15 15 15 15 15		-	

Methodology : Hexokinase

Interpretation Notes:

Interpretation (In accordance with the American diabetes association guidelines):

A fasting plasma glucose level below 100 mg/dL is considered normal.

· A fasting plasma glucose level between 100-126 mg/dL is considered as glucose intolerant or pre diabetic. A fasting and postprandial blood sugar test (after consumption of 75 gm of glucose) is recommended for all such patient.

A fasting plasma glucose level of above 126 mg/dL is highly suggestive of a diabetic state. A repeat fasting test is strongly recommended for all such patients. A fasting plasma glucose level in excess of 126 mg/dL on both the occasions is confirmatory of a diabetic state.

PLASMA GLUCOSE POST PRANDIAL

99.8

mg/dL

80 - 140

(PPBS)

Methodology: GPO-POD Interpretation Notes:

Fasting Glucose Plasma 02 hr Plasma Glucose

Diagnosis

</=99

</=139

Normal

100 to 125

1'40 to 199

Pre Diabetes

>126

>200

Diabetes

Confirm by repeating the test on a different day

Impaired glucose tolerance (IGT) fasting, means a person has an increased risk of developing type 2 diabetes but does not have it yet. A level of 126 mg/dL or above, confirmed by repeating the test on another day, means a person has diabetes. IGT (2 hrs Post meal), means a person has an increased risk of developing type 2 diabetes but does not have it yet. A 2-hour glucose level of 200 mg/dL or above, confirmed by repeating the test on another day, means a person has diabetes

Ref : American Diabetes association standards of medical care.

Sample Type: Plasma.

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HAEMATOLOGY

Test Name Obtained Value Units Bio. Ref. Intervals(Age/Gender specific)

BLOOD GROUP, RH FACTOR

Methodology : Forward & Reverse

Blood Grouping

Visit No.

"B"

RH Typing

POSITIVE .

Sample Type: Whole Blood-EDTA

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PREM-DHARAM **HOSPITAL & DIAGNOSTICS**



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HAEMATOLOGY

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+1		HbA1c	
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific)
GLYCOSYLATED HAEMOGLOBIN(HbA1c)	4.85	%	4.5 - 6.0
· · · · · · · · ·	24 21		Good Control : 6.1-7.0 Fare Control : 7.1-9.0
Methodology : HPLC			Poor Control : >9.0
ESTIMATED AVERAGE GLUCOSE(eAG)	92.50	mg/dL	90 – 120 Excellent Control 121 – 150 Good Control
Methodology : Calarias d			151 – 180 Average Control 181 – 210 Action Suggested > 211 Panic Value

Methodology: Calculated

Sample Type : Whole Blood-EDTA

Interpretation Notes:

In vitro quantitative determination of HbA1c in whole blood is utilized in long term monitoring of glycemia. The HbA1c level correlates with the mean glucose concentration prevailing in the course of the patient's recent history (approx - 6-8 weeks) and therefore provides much more reliable information for glycemia monitoring than do determinations of blood glucose or urinary glucose. It is recommended that the determination of HbA1c be performed at intervals of 4-6 weeks during Diabetes Mellitus therapy. Results of HbA1c should be assessed in conjunction with the patient's medical history, clinical examinations and other findings.

Note: If variant hemoglobin is observed in HbA1c HPLC screen, HbA1c levels may not truly represent in vivo condition. In such condition HbA1c analysis by HPLC may not be the method of choice. You are advised to consult your referring physician and discuss the alternative tests as

Advised:

1.To follow patient for glycemic control test like fructosamine or glycated albumin may be performed instead. 2.Hemoglobin HPLC screen to analyze abnormal hemoglobin variant.

estimated Average Glucose (eAG):

estimated Average Glucose (eAG) based on value calculated according to National Glycohemoglobin Standardization Program (NGSP) criteria.

*** End Of Report ***

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Dr. Vivek Kapoor

onsultant Pathologist

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Reported On : 26/03/2024 18:18

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THYROID PROFILE				
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific)	
TOTAL TRIIODOTHYRONINE (T3)	1.12	ng/dL	0.70-2.04	
Methodology : Chemiluminescence Immunoass	say(CLIA)	(c) 3 (
TOTAL THYROXINE (T4)	7.62	ug/dl	4.5 - 12	
Methodology : Chemiluminescence Immunoass	say(CLIA)	-9/	- I - I - I - I - I - I - I - I - I - I	
THYROID STIMULATING HORMONE (TSH)	3.66	uIU/mL	0.35 –5.50	
			Newborns: 0.70 - 15.2 Peadiatric:	
			2weeks-4 months :1.7-9.1	
8			<12 months : 1.36 - 8.8 1- 6 years : 0.85 - 6.5	
			7-12 years : 0.28 – 4.3	
			Pregnancy:	
· · · · · · · · · · · · · · · · · · ·			1st Trimester: 0.1-2.5	
Anthodol Cl V			2nd&3rd Trimester:0.2-3.0	

Methodology: Chemiluminescence Immunoassay(CLIA)

Sample Type : serum

Interpretation Notes:

- 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 concentrations.
- 2. Recommended test for T3 and T4 is unbound fraction or free levels as it is metabolically
- 3. Physiological rise in Total T3/T4 levels is seen in pregnancy and in patients on steroid therapy.

*** End Of Report ***

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Verified by : Nisha Sharma MSC MB

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	IMMUNOLOGY				
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific)		
PROSTATE SPECIFIC ANTIGEN -TOTAL	0.29	ng/mL	0.0-4.0		

Methodology: Chemiluminiscence

Interpretation Notes:

The PSA test and digital rectal exam (DRE) may be used to screen both asymptomatic and symptomatic men for prostate cancer. PSA is a protein produced primarily by cells in the prostate and most of the PSA is released into semen, but small amounts of it are also released into the blood. PSA exists in two forms in the blood: free (not bound) and complexed (cPSA, bound to other proteins). Lab tests can measure free PSA or total PSA (bound plus unbound). Some organizations, such as the U.S. Preventive Services Task Force, feel that the harms associated with over-diagnosis and over-treatment outweigh the potential benefits and advise against using PSA to screen for prostate cancer in healthy men of any age. The American Cancer Society and the American Urological Association recommend that men discuss the advantages and disadvantages of PSA-based screening for prostate cancer with their healthcare provider before making an informed decision about whether to be screened or not. While elevated PSA levels are associated with cancer, they may be caused by other conditions, such as benign prostatic hyperplasia (BPH) and inflammation of the prostate. An elevated PSA may be followed by a biopsy, which has risk of complications such as pain, fever, blood in the urine, or urinary tract infection. (Read the article on Anatomic Pathology for more information about biopsies.)

Sample Type: serum

Processed by : Nisha Sharma Verified by : Tabassum

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	CLINICA	L PATHOLOG	Y
Test Name	Obtained Value	Units	Bio. Ref. Intervals(Age/Gender specific
URINE ROUTINE			A
PHYSICAL EXAMINATION	a, _ 11 as 11		and the
Quantity	20	ml	
colour	PALE YELLOW		-
Appearance	CLEAR		
рН	6.0		4.5 - 8
Specific Gravity	1.005		1.005 - 1.025
MICROSCOPIC EXAMINATION			
Pus Cells	0-1	/HPF	1 - 3
RBC CELLS	NIL	/HPF	
pithelial Cells	0-1	/HPF	1 - 2
Casts	ABSENT	/Hpf	
Crystals .	ABSENT	/Hpf	1
CHEMICAL EXAMINATION	•		
Albumin/Protein	ABSENT		-
Glucose	ABSENT .		
Jrobilinogen	ABSENT		
lood	ABSENT		
litrite	ABSENT		
eucocyte	ABSENT		
nterpretation Notes:	0.000	200	
ample Type : URINE			

*** End Of Report ***

Processed by : Tabassum Verified by : Tabassum



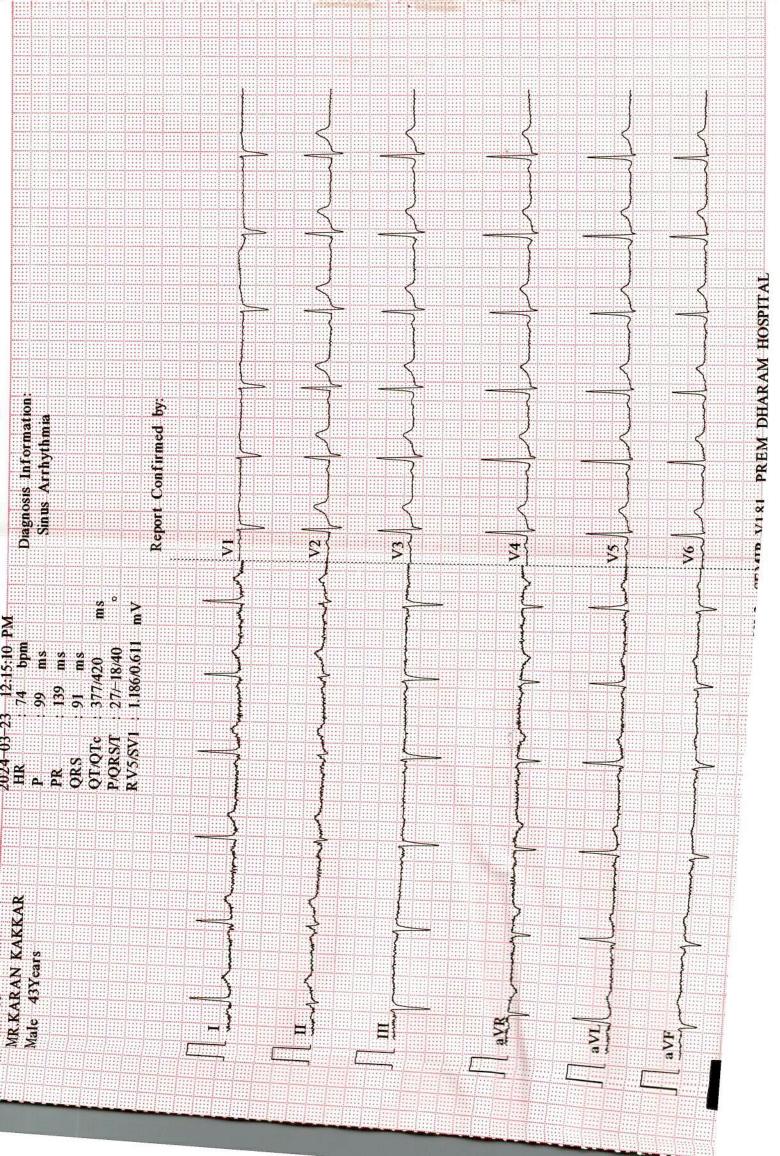






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Date

: 23/03/2024

Receipt No. :

Pt.ID No. : PAT/22_23/2203:

Patient Name : Mr. KARAN KAKKAR

Age

43 Y 0 M

9539

Sex

: Male

Ref.By

: PREM DHARAM HOSPITAL

XRAY CHEST PA

Bilateral lung fields show no obvious parenchymal lesion.

Trachea is central.

Hila and mediastinum are unremarkable.

Both domes of diaphragm are normal.

Both cardiophrenic and costophrenic angles are normal.

Cardiac size is normal.

Bony thoracic cage appears normal.

ADVISED: CLINICOPATHOLOGICAL CORRELATION.

(1) This report is a professional opinion based on imaging findings. NOT valid for medico-legal pu



Dr.Nidhi Tyagi Consultant Radiologist



- 3D/4D Ultrasound
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 Whole Body Color Doppler
- Digital X-Ray (24 Hours)
- 2D Echocardiography
- ECG-3 Channel







FOR TELEPHONIC APPOINTMENT RADIOLOGY: 8287900395 (9 AM - 2:30 PM, 5:30 PM - 8:45 PM) FOR ICU & AMBULANCE ENQUIRY: 8287900395 FOR OPD APPOINTMENT: 8287900395





(Govt. Recognised Advanced 3D-4D Ultrasound and State-of-the-art Digital X-Ray Centre) Regn. No.: RMEE1905483

0D-180 (Near Nagar Nigam Office), Vasundhara, Ghaziabad, Phone: 0120-4127778, 9899004884, E-mail : premdharamhospital@ymail.com

Date

23/03/2024

Srl No. 13

Pt. Id.

Name

MR. KARAN KAKKAR

Age 43 Yrs.

Sex

PD/2303202411

Ref. By

PREM DHARAM HOSPITAL

WHOLE ABDOMEN MALE

Liver is normal in size, outline and echopattern. No focal SOL noted. Vascular channels are clear. No evidence of I.H.B.D. Portal vein is normal in course and caliber.

Gall Bladder is partially distended with normal wall thickness. A tiny isoechoic focus measuring approximately 2.9X1.81 mm seen along gall bladder wall at present scan? prominent mucosal fold. Needs-Follow up USG for better evaluation.CBD is normal.

Spleen is normal in size, shape and shows homogenous echotexture. No focal SOL noted.

Pancreas shows normal outline and echopattern. No focal SOL noted. Pancreatic duct is not dilated.

Retroperitoneum No evidence of significant retroperitoneal lymphadenopathy noted.

Right Kidney is normal in size, (measures approximately mm) outline and parenchymal echopattern.

Cortico-medullary differentiation is well preserved. Pelvi-calyceal system is compact.

A calculus measuring approximately 4.3 mm seen at upper pole of right kidney.

Left Kidney is normal in size, (measures approximately 107.02X55.26 mm) outline and normal parenchymal echopattern. Cortico-medullary differentiation is well preserved. Pelvi-calyceal system is compact. No intra renal calculus seen.

Tiny echogenic non-shadowing foci measuring approximately 2.3 mm, 1.78 mm and 2.15 mm seen at mid pole of left kidney? concretions. Adv-Follow up USG for better evaluation.

Urinary Bladder is partially distended and appears anechoic. The outline is smooth with mildly echogenic walls. The bladder wall thickness appears normal. There is no evidence of any debris or echogenic calculus in

Prostate is mildly enlarged in size(measuring approximately 36.3x44.0x38.1mm vol -(30.15-31.86 cc), shows normal outline and echopattern. No focal SOL noted.

Bowel loops show no abnormal dilatation or bowel wall thickening.

No free fluid in visualized peritoneal or pleural cavity.

IMPRESSION:

USG WHOLE ABDOMEN REVEALS -RIGHT RENAL CALCULUS WITH BODERLINE PROSTATOMEGALY AND ? TINY LEFT RENAL CONCRETIONS

Clinical and lab correlation /Follow up USG is recommended for further evaluation.

Note: (1) This report is a professional opinion based on imaging findings. NOT valid for medico legal p (2) in case of any discrepancy due to machine error or typing error, please get it rectified immediately.

> Dr. NIDHI TYAGI CONSULTANT RADIOLOGIST



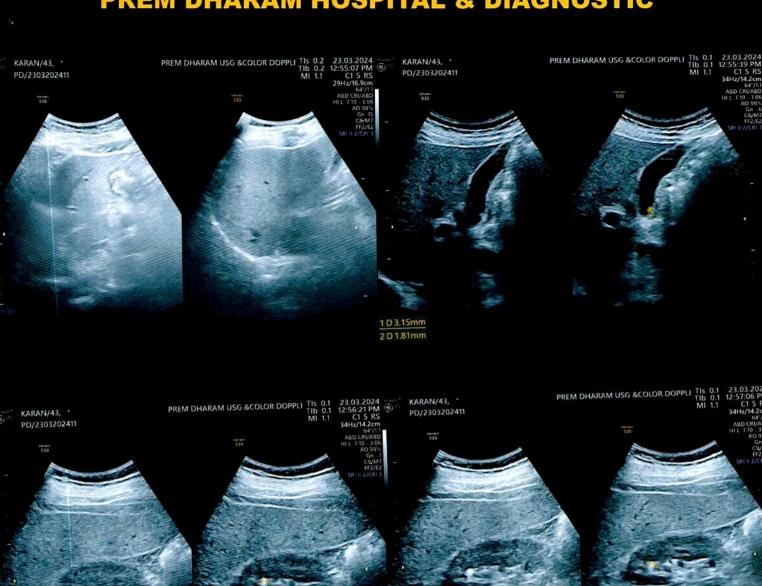






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1 D 3.75mm

1 D 4.30mm



1 D 2.37mm

1 D 1.78mm