Sample Collection Date	:17-09-2021		
Lab Ref. No.	:CP/78		
Name	:PRABHAT KUMAR	Age	:40Y
Ref. by	: DR.PARASAR HOSPITAL	Sex	: F

PATHOLOGY REPORT

/////

TESTS	RESULTS	EXPECTED VALUES	
THYROID			
TSH	0.96	Euthyroidism: 0.25-5µIU/ml	
Т3	1.12	Euthyroidism: 0.61-1.63ng/ml	
T4	8.99	Euthyroidism: 4.68-9.36 µg/dl	

Wallach's reference range for Thyroid for neonates and children

1-4 daya	1-39	11.08-21.61	0.97-7.42	
1-4 webs	1.7-9.1	8.29-17.24	1.04-3.45	
1-12 mon	0.8-8.2	5.93-16.38	1.04-2.47	
1-6 yrs	0.7-5.7	7.33-15.04	1.04-2.66	
6-10 yrs	0.7-5.7	6.40-13.33	0.91-2.40	
11-15 yrs	0.7-5.7	5.54-11.78	0.84-2.14	
15-18 yrs	0.7-5.7	4.21-11.86	0.78-2.0	
Pregnancy	T	SH	T3	T4
let Trimenter	0.3 t	to 4.5 0.1	81 -1.90	7.8-14.77
2 nd Trimester	0.5 t	to 4.6	00-2.60	7.14-19.58
3 rd Trimester	0.8 1	to 5.2 1.	00-2.60	8.32-17.02

Note:

- TSH stimulates the thyroid gland to produce the main thyroid hormones T3 and T4.
- In cases of hyperthyroidism TSH level is severely inhibited and may even be undetectable.
- In rare forms of high-origin hyperthyroidism, the TSH level is not reduced, since the negative-feedback control of the thyroid hormones has no effect.
- In cases of primary hypothyroidism, TSH levels are always much higher than normal and thyroid hormone levels are low.
- The TSH assay aids in diagnosing thyroid or hypophysial disorders.
- The T4 assay aids in assessing thyroid function, which is characterized by a decrease in thyroxine levels in patients with hypothyroidism and an increase in patients with hyperthyroidism.
- The T3 plays an important part in maintaining euthyroidism.
- TSH, T4 & T3 determination may be associated with other tests such as FT4 & FT3
- assay, as well as with the clinical examination of the patient

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Indra Diagnostics WE ASSURE QUALITY

Add. : G-127, PC Colony, Kankarbagh, Patna-20 Mob. : 7870077439, Ph. : 0612-2351609 email : indradiagnostics439@gmail.com

TUMOR MARKER					
Test Description Result Reference Range Unit(s)					
Referred By	: PRASHAR HOSPITAL	Accession N			
Organization	MUZAFFARPUR-CC		ate & Time: 18/09/2021, 12:49 AM		
Sex	: Male	Received Da	ate & Time: 17/09/2021, 10:30 PM	回当处于在产品。	
Age	: 49 years	Collected O	n : 17/09/2021, 12:40 PM		
Patient Name	PRABHAT KUMAR	Primary San	nple : Serum		
Pt. No	MUZ3096	Location		· · · · · · · · · · · · · · · · · · ·	

PROSTATE SPECIFIC ANTIGEN

PSA (Prostate Specific Antigen)-Total

Method : Serum, CLIA

0 - 4.0

Interpretation:

- Increased levels are noted in Prostate cancer, Bengin prostatic hypertrophy. Prostatitis. ٠
- Values obtained with different assay methods or kits may be different and cannot be used interchangeably. ٠

1.30

Specimens drawn from patients undergoing prostate manipulation, especially needle biopsy and transurethral resection, may ٠ show erroneously high prostatic-specific antigen (PSA) results. Care should be taken that specimens are drawn before these procedures are performed.

END OF REPORT



PATIENT NAME	:PRABHAT KUMAR		REFF.NO	:18/09/21CP/93
AGE/SEX	:49Y/Adult-M		TEST.DATE	:18/09/2021
REF.BY DR.	:DR.PARASAR HOSPITAL		RPT.DATE	:18/09/2021
INVESTIGATIONS		VALUE	REF.R	ANGE UNIT

URINE ROUTINE

Physical Examination	
Colour	Pale Straw
Appearance	Clear
Chemical Examination	
PH	Neutral
Specific Gravity	1.025
Sugar	Negative
Albumin	Negative
Microscopic Examination	
Crystal	Nil
RBC	Nil
WBC	0-1/HPF
Epithelial Cells	A Few
Granular Cast	Absent
Bacteria	Absent
Cast	Absent
Other	NIL

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PATIENT NAME	:PRABHAT KUMAR		REFF.NO	:18/09/2	1CP/93
AGE/SEX	:49Y/Adult-M		TEST.DATE	:18/09/2	021
REF.BY DR.	:DR.PARASAR HOSPITAL		RPT.DATE	:18/09/20)21
INVESTIGATIONS		VALUE	RE	EF. RANGE	UNIT

ROUTINE

Stool Examination.

Colour	Yellow
Consistency	Semi Solid
Reducing Sugar	Nil
Mucus	Nil
Epethelial Cells	Nil
* Occult Blood	Negative
Parasites	E.Histolytica(+)
RBC	Nil
Undigested Food Particles	Absent
WBC	Nil
CYSTS	Nil
AVO	Nil

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PATIENT NAME : PRABHAT KUMAR		REFF.NO	:17/09/21CP/78
AGE/SEX :49Y/Adult-M		TEST.DATE	:17/09/2021
REF.BY DR. :DR.PARASAR HOSPI	TAL	RPT.DATE	:18/09/2021
INVESTIGATIONS	VALUE	REF.RANGE	UNIT
	IAEMATOLOG	Ĩ	
CBC (COMPLETE BLOOD COUNT)			
RBC (Red Blood Cells)	4.90	3.8-6.5	M/uL
Hb (Heamoglobin)	14.2	12-16	gm/dl
HCT (Haematocrit)	45.8	37-54	<u>_</u>
MCV	93.4	76-96	fl
MCH	33.6	27-32	рq
MCHC	36.0	31-36	gm%
TC of WBC	8,800	4500-10500	/cumm
DC of WBC			
Neutrophil (seg)	60	50 - 70	8
Lymphocyte	35	20 - 40	8
Eosinophil	04	1 - 6	<u>e</u>
Monocyte	01	2 - 8	<u>_</u>
Basophil	00	0 - 1	8
Platelets	1.25	1.5-3.5	lakh/cumm
Red Cell Distribution Width (RDW)	13.5	11.5-14.5	fL
ESR (DONE BY AUTO ANALYZER)	04	0-20	mm/hr
В	IOCHEMISTR	RY	
Plasma Glucose (Fasting)	73	70-110	mg/dl
Plasma Glucose(Post Prandial)	106	100-140	mg/dl
LIVER PROFILE			-
Serum Bilirubin			
Total	0.9	0.1-1.2	mg⊹
Direct	0.5	0-0.3	mg%
Indirect	0.40	0-0.6	mg%
Serum SGPT(ALAT)	26	9-38	IU/L

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PATIENT NAME :PRABHAT KUMAR		REFF.NO	:17/09/21CP/78
AGE/SEX :49Y/Adult-M		TEST.DATE	:17/09/2021
REF.BY DR. :DR.PARASAR HOSPI	TAL	RPT.DATE	:18/09/2021
INVESTIGATIONS	VALUE	REF . RANGE	UNIT
Serum SGOT(ASAT)	32	10-40	IU/L
Alkaline Phosphatase	236	80-306	U/L
Total Protien			
Protein	7.4	6.0-8.0	g/dl
Albumin	4.6	3.7-5.3	g/dl
Globulin	2.80	2.3-3.6	g/dl
A:G Ratio	1.64:1	1.0-2.3	Ratio
Gamma Glutamyl transferase(GGT)	26	9-58	U/L
RENAL FUNCTION TEST (RFT)			
Serum Creatinine	1.1	0.7-1.4	mg/dl
Blood Urea Nitrogen(Bun)	13.08	7-21	mg/dl
Serum Uric Acid	5.1	3.6-7.7	mg/dl
Serum Calcium(Total)	10.0	8.7-11.0	mg/dl
Blood Urea	28	13-45	mg/dl
SERUM ELECTROLYTE			
Sodium (Na)+	144.7	135-150	m mol/L
Potassium (K)+	4.1	3.5-5.1	m mol/L
Chloride (Cl)-	106.2	97-111	m mol/L
LIPID PROFILE			
Serum Total Cholesterol	245	50-230	mg/dl
Serum HDL Cholesterol Direct	35	30-65	mg/dl
Serum Triglycerides	298	25-200	mg/dl
Serum LDL Cholesterol	150.40	85-130	mg/dl
Serum VLDL Cholesterol	59.60	7-32	mg/dl
TC/HDL Cholesterol Ratio	7.00	3.0-5.0	Ratio
LDL/HDL Ratio	4.30:1	1.5-3.5	Ratio

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INVESTIGATIONS		VALUE	REF.]	RANGE	UNIT
REF.BY DR.	:DR.PARASAR HOSPITAL		RPT.DATE	:18/09/	2021
AGE/SEX	:49Y/Adult-M		TEST.DATE	:17/09	/2021
PATIENT NAME	:PRABHAT KUMAR		REFF.NO	:17/09	/21CP/78

IMMUNOLOGY

Blood Group

ABORH

"O"POSITIVE

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CURE PATHOLOGY (A Unit of Anand arya Healthcare)	 Level And Constraints of the end of the en
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Patient Name	PRABHAT KUMAR	CP	:78	Age/Sex	:49Y/M
Refered Doctor	DR.PARASAR HOSPITAL	OPD/IPD	: OPD		
		Date	17-09-2021		

BLOOD EXAMINATION REPORT

Investigation	Result	R.Range	Unit
GLYCOSYLATED NEMOGLOBIN, BLOOD			
GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.3	High Non-diabetic: < 5.7 Pre-diabetics: 5.7 - Diabetics: > or = 6. ADA Target: 7.0	6.4 5
MEAN PLASMA GLUCOSE	91	Action suggested: > High < 116.0	8.0 mg/dL

Interpretation(s)

GLYCOSYLATED HEMOGLOBIN, BLOOD-

Glycation is nonenzymatic addition of sugar residue to amino groups of proteins. HbA1C is formed by the condensation of glucose with n-terminal valine residue of each beta chain of hb a to form an unstable schif base. It is the major fraction, constituting approximately 80% of HbA1. Formation of glycated hemoglobin (GHb) is essentially irreversible and the concentration in the blood depends on both the lifespan of the red blood (120 days) and the blood glucose concentration. The GHB concentration represents the integrated values for glucose aver cells (RBC) the period of 6 to 8 weeks. GHb values are free of day to day glucose fluctuations and are unaffected by recent exercise or food ingestion. Concentration of plasma glucose concentration in GHb depends on the time interval, with more recent values providing a larger contribution than earlier values.

The interpretation of GHb depends on RBC having a normal life span. Patients with hemolytic disease or other conditions with shortened RBC survival exhibit a substantial reduction of GHb. High GHb have been reported in iron deficiency anemia .

GHb has been firmly established as an index of long term blood glucose concentrations and as a measure of the risk for the development of patients with diabetes mellitus. The absolute risk of retinopathy and nephropathy are directly proportional to the complications in mean of HbAIC.

"Targets should be individualized More or less stringent glycemic goals may be appropriate for individual patients. Goals should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycemia unawareness, and individual patient considerations."

End Of Report

End of Report

Dr Sima kumari



