

Patient Name : Mr. SUNNY
Age / Gender : 36 Years / Male
Referred By : Dr. KARAN CHHABRA
Req.No : 2195772
Patient Type : OPD

UHID : 24009
IPNO :
Requisitions : 28/03/2022 / 9.28 AM
Sample collection : 28/03/2022 / 09.34 AM
Sample Receiving : null /
Reported on : 28/03/2022 / 1.09 PM

BIOCHEMISTRY

BLOOD SUGAR RANDOM

Specimen Type

TEST NAME

Glucose Random

RESULT

83.3

UNITS

mg/dL

BIOLOGICAL

REFERENCE INTERVAL

70 - 140

METHOD

GOD-POD Hexokinase

Note: Random Glucose refers to levels between meals with a minimum time lapse of 2 hours.

-**** End of Report ****-

Please Correlate With Clinical Findings

Gaurvi

Lab Technician Dr. GAURVI PIPLANI
MD (Pathology)

Dr. KANIKA GUPTA
MD (Pathology)

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Sample Receiving : null /
Reported on : 28/03/2022 / 1:12 PM

HAEMATOLOGY

COMPLETE HAEMOGRAM (CBC ESR)

Specimen Type	Whole Blood		BIOLOGICAL	
TEST NAME	RESULT	UNITS	REFERENCE INTERVAL	METHOD
Haemoglobin	15.9	gm/dl	13.0 - 17.5	Cyanide-Free Colorimetry
Total Leucocyte Count	7500	/ μ l	4000 - 11000	Impedance Variation
DIFFERENTIAL COUNT				
Neutrophils.	57	%	40.0 - 75.0	Flow Cytometry
Lymphocytes.	36	%	20.0 - 45.0	Flow Cytometry
Monocytes	06	%	2.0 - 10.0	Flow Cytometry
Eosinophils.	01	%	0.0 - 4.0	Flow Cytometry
Basophils	00	%	0.0 - 1.0	Flow Cytometry
Platelet Count	3.02	1000/cumm	1.50 - 4.50	Electrical Impedance
RED BLOOD CELL COUNT	6.05	millions/cum m	3.5 - 5.5	Electrical Impedance
PACKED CELL VOLUME	47.6	%	36 - 46	Calculated
MEAN CORPUSCULAR VOLUME	78.7	fL	76 - 96	Measured
MEAN CORPUSCULAR HAEMOGLOBIN	26.3	pg	27 - 32	Calculated
MEAN CORPUSCULAR Hb CONC	33.4	gm/dl	33 - 37	Calculated

----- End of Report -----
Please Correlate With Clinical Findings

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Sample Receiving : null / 09:34:01
Reported on : 28/03/2022 / 5.39 PM

CLINICAL PATHOLOGY
URINE ROUTINE MICROSCOPY

Specimen Type	BIOLOGICAL			
TEST NAME	RESULT	UNITS	REFERENCE INTERVAL	METHOD
PHYSICAL EXAMINATION				
volume	20	ml		
colour	Pale Yellow		Pale Yellow	
Appearance	Clear		Clear	
Specific Gravity	1.020			Polyelectrolytes Ionic
reaction	Acidic		Acidic	
pH -Urine	6.0			PH paper
Albumin	NIL		NIL	Protein-error-of-Indicator Sulphosalicylic Acid
Glucose	NIL		NIL	GODPOD/Benedicts
Bile Salt	NIL		NIL	
Bile Pigment	NIL		NIL	Diazo/Fouchets Test
Urobilinogen	NIL		NIL	Elrich Aldehyde

-**** End of Report ****
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Sample Receiving : null / 09.34:04
Reported on : 28/03/2022 / 5.39 PM

CLINICAL PATHOLOGY

MICROSCOPIC EXAMINATION

PUS CELLS - URINE	2-4	/HPF		
Red blood cells	Nil		NIL	
Epithelial Cells - Urine	1-2		4---5/HPF	
Casts	NIL		NIL	Microscopic
Crystals.	NIL		NIL	Microscopic

Albumin test positive by Multistrip Method is confirmed by Sulphosalicylic acid method.

-**** End of Report ****

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Sample Receiving : 28/03/2022 / 09:34.04
Reported on : 28/03/2022 / 5.33 PM

HAEMATOLOGY

HBA1C

Specimen Type

TEST NAME	RESULT	UNITS	REFERENCE INTERVAL	METHOD
Glycosylated Haemoglobin (Hb A1c)	6.1	%	NON DIABETIC:<5.7 PRE DIABETIC:5.7-6.4 DIABETICS: >OR 6.5 ADA TRAGET:7.0	Latex immunoagglutination inhibition methodology

*Done on DCA Vantage

*Results of these tests should always be interpreted in conjunction with patients medical history, clinical presentation and other findings.

*The results of HbA1c are not influenced by recent meals, physical activity or emotional stress.

-**** End of Report ****-

Please Correlate With Clinical Findings

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BIOCHEMISTRY

LIPID PROFILE.(TOTAL CHOLESTEROL,LDL,HDL,TRIGLYCERIDES)

Specimen Type

BIOLOGICAL

TEST NAME	RESULT	UNITS	REFERENCE INTERVAL	METHOD
<u>LIPID PROFILE</u>				
SERUM CHOLESTROL	156.1	mg/dl	0 - 200	Cholestrol Oxidase
Serum Triglycerides	140.7	mg/dl	Up to 150	GPO -Trinder
HDL Cholesterol	45.2	mg/dl	0 - >50	Direct Method
LDL Cholesterol	82.76	mg/dl	Optimal <100,Above Opt. 100-129 -high 160-189	Direct Measure
VLDL Cholesterol	28.14	mg/dL	*Less than 30	Calculated

*Automated Direct HDL And LDL Estimation.

*Results of these tests should always be interpreted in conjunction with patients medical history, clinical presentation and other findings

***** End of Report *****

Please Correlate With Clinical Findings

Gaurvi

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 MD (Pathology)

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BIOCHEMISTRY

KFT(KIDNEY FUNCTION TEST)/RFT/Renal Profile

Serum

TEST NAME	RESULT	UNITS	REFERENCE INTERVAL	METHOD
<u>Urea Creatinine</u>				
Serum Urea	14.0	mg/dl	13 - 45	UreaseGLDH
Serum Creatinine	0.97	mg/dl	Male: 0.6 - 1.3	Modified JAFFES
Serum Uric Acid	4.3	mg/dl	Adult Male: 3.5 - 7.2	Uricase Trinder, End Point (Toos)

*Results of these tests should always be interpreted in conjunction with patients medical history, clinical presentation and other findings.
 *Performed on fully Automated Dimension X-Pand plus BioChemistry Analyser.
 *External Quality Control by Biorad Laboratory.

******* End of Report *******
Please Correlate With Clinical Findings

Lab Technician **Dr. GAURVI PIPLANI** **Dr. KANIKA GUPTA**
 MD (Pathology) MD (Pathology)

Patient Name : Mr. SUNNY
Age / Gender : 36 Years / Male
Referred By : Dr. KARAN CHHABRA
Req.No : 2195772
Patient Type : OPD

UHID : 24609
IPNO :
Requisitions : 28/03/2022 / 09:34 AM
Sample collection : 28/03/2022 / 09:34 AM
Sample Receiving : null /
Reported on : 28/03/2022 / 1:10 PM

BIOCHEMISTRY

LFT(LIVER FUNCTION TEST)

Specimen Type	Serum	RESULT	UNITS	REFERENCE INTERVAL	BIOLOGICAL METHOD
TEST NAME					
TOTAL BILIRUBIN		0.83	mg/dL	0.1 - 1.2	Diazotized Sulphanilic Acid
DIRECT BILIRUBIN		0.40	mg/dL	0.00 - 0.20	Diazotized Sulphanilic Acid
INDIRECT BILIRUBIN		0.43	mg/dL	0.0 - 0.9	Diazotized Sulphanilic Acid
SGOT (AST)		20.8	IU/L	0 - 35	IFCC WPP AMP
SGPT (ALT)		12.7	IU/L	5 - 40	IFCC WPP AMP
Alkaline Phosphatase		95.6	IU/L	Adult: 50 - 136	Modified IFCC
Total Protein		7.06	g/dl	6.4-8.2	Biuret Endpoint
Albumin - Serum		4.44	g/DL	3.2 - 5.0	Photometric Column test BCG Dve
Globulin		2.62	gms%	2.3 - 4.5	

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-**** End of Report ****
 Please Correlate With Clinical Findings

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Dr. KANIKA GUPTA
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Reported on : 28/03/2022 / 1:38 PM

BIOCHEMISTRY

BSPP (BLOOD SUGAR PP)

Specimen Type

TEST NAME

BIOLOGICAL

RESULT UNITS REFERENCE INTERVAL. METHOD

FASTING PP

Plasma glucose(fasting.)

83.3 mg/dl

70 - 110

GOD-POD Hexokinase

Plasma Glucose(POST Prandial)

94.0 mg/dl

90 - 140

GOD-POD Hexokinase

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*Performed on fully Automated Dimension X-Pand plus BioChemistry Analyser

*External Quality Control by Biorad Laboratory.

**** End of Report ****
Please Correlate With Clinical Findings

Gaurvi

Lab Technician Dr. GAURVI PIPLANI
MD (Pathology)

Dr. KANIKA GUPTA
MD (Pathology)

Client Code: HR46
Client Name And Address:
PARK HOSPITAL,
AMBALA CHANDIGARH ROAD, AMBALA CITY



NAME : Mr. SUNNY 24009
AGE/GENDER : 36 Y/Male
TEST REQUEST ID : 012203280133
REFERRED BY : Dr. PARK HOSPITAL,
SAMPLE ID : 10229236

Patient_ID : 230821
SPECIMEN DATE : 28/Mar/2022 07:24PM
SPECIMEN RECEIVED : 28/Mar/2022 07:46PM
REPORT DATE : 28/Mar/2022 10:11PM
PRINT DATE : 07/Apr/2022 12:00PM

Investigation Name	Result	Unit	Biological Ref.Interval
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Thyroid Function Test(T3,T4,TSH)

Primary Sample Type:Serum

Triiodothyronine total [T3] Chemiluminescence Microparticle Immuno Assay	93	ng/dL	70-200
Thyroxine total [t4] Chemiluminescence Microparticle Immuno Assay	7.10	ug/dL	4.87-11.72
TSH (4th Generation) Chemiluminescence Microparticle Immuno Assay	3.162	uIU/mL	0.35-4.94

INTERPRETATION

Link with age for Males > 20 years

REFERENCE GROUP	REFERENCE RANGE IN uIU/mL
Males > 20 years	0.5 - 4.8

Below mentioned Table to appear only for female patients > 20 years.No value in reference range

REFERENCE GROUP	REFERENCE RANGE in uIU/mL (As per American Thyroid Association)
Adult Females (> 20 years)	0.5 - 4.8
Pregnancy	Reference Range
First Trimester	0.10- 2.50
Second Trimester	0.20 - 3.00
Third Trimester	0.30 - 3.00

Note: 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.

Clinical Use

V.K. Dogra

Dr. V.K. Dogra
MD Path.
Director
Sr. Consultant Pathologist



Client Code: HR46
Client Name And Address:
PARK HOSPITAL (P)
AMBALA CHANDIGARH ROAD,AMBALA CITY



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Investigation Name	Result	Unit	Biological Ref.Interval
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- Diagnose Hypothyroidism and Hyperthyroidism
- Monitor T4 replacement or T4 suppressive therapy
- Quantify TSH levels in the subnormal

Range Increased Levels:

- Primary hypothyroidism
- Subclinical hypothyroidism
- TSH dependent Hyperthyroidism
- Thyroid hormone resistance

Decreased Levels:

- Graves disease
- Autonomous thyroid hormone secretion
- TSH deficiency

Comment

T₃ or 3,5,3 triiodothyronine is a hormone synthesized and secreted from the thyroid gland, and formed by peripheral deiodination of thyroxine (T₄). The determination of T₃ levels in serum is essential in assessing thyroid functions. T₃ is secreted by thyroid glands and circulates in the blood stream; mostly (99.7%) bound to the plasma protein, thyroxin binding globulin (TBG) and prealbumin (TBPA) and albumin. The remaining (0.3%) is free, unbound and its metabolic potency is much greater. T₃ hormone regulates cell metabolism and body growth and its level is a good indicator of thyroid disease state and body metabolism. Further the concentrations of the carrier protein are altered in many conditions such as pregnancy in normal thyroid function, as the concentrations of the carrier proteins alters, the total T₃ level changes so that free T₃ concentration remains constant. Thus, measurements of the free T₃ concentrations correlate excellently with clinical status than total T₃ levels.

T₄ or Thyroxine or 3,5,3,5-tetraiodothyronine is a hormone synthesized and secreted by the thyroid gland and plays an important role in regulating metabolism. In the peripheral tissues it act as a prohormone which is further metabolized to another most active thyroid hormone, triiodothyronine (T₃) and other inactive metabolites such as reverse T₃.

TSH or Thyroid-stimulating hormone is a hormone synthesized and secreted by Pituitary gland. TSH is glycoprotein with two non-covalently bound alpha and beta subunits. The beta subunit of TSH is unique, which results in the specific biochemical and immunological properties of this hormone. The ability to quantitate circulating levels of 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. The measurement of serum TSH has proven to be one of the most sensitive methods for the detection of primary hypothyroidism. In primary hypothyroidism the production of thyroid hormones is impaired and the TSH levels are observed to be higher. However in secondary and tertiary hypothyroidism the TSH levels are low because of pituitary or hypothalamic lesions. In hyperthyroidism the circulating levels of TSH is usually subnormal. In some instance however this condition may result from hyperstimulation of thyroid.

Vasu Dogra

Dr. V.K. Dogra

MD Path.

Director

Sr. Consultant Pathologist

