

**Name** : MR PANKAJ SINGH

**Age** : 34 Yr(s) Sex :Male

**Registration No** : MH011292080

**Lab No** : 202309001986

**Patient Episode** : H18000001080

**Collection Date** : 09 Sep 2023 12:16

**Referred By** : HEALTH CHECK MGD

**Reporting Date** : 09 Sep 2023 18:34

**Receiving Date** : 09 Sep 2023 12:16

## CLINICAL PATHOLOGY

### ROUTINE URINE ANALYSIS (Semi Automated)Specimen-Urine

#### MACROSCOPIC DESCRIPTION

Colour	PALE YELLOW	(Pale Yellow - Yellow)
Appearance	CLEAR	
Reaction[pH]	7.0	(4.6-8.0)
Specific Gravity	1.005	(1.003-1.035)

#### CHEMICAL EXAMINATION

Protein/Albumin	Negative	(NEGATIVE)
Glucose	NIL	(NIL)
Ketone Bodies	Negative	(NEGATIVE)
Urobilinogen	Normal	(NORMAL)

#### MICROSCOPIC EXAMINATION(Automated/Manual)

Pus Cells	1-2 /hpf	(0-5/hpf)
RBC	NIL	(0-2/hpf)
Epithelial Cells	0-1 /hpf	
CASTS	NIL	
Crystals	NIL	
Bacteria	NIL	
OTHERS	NIL	

Name : MR PANKAJ SINGH

Age : 34 Yr(s) Sex :Male

Registration No : MH011292080

Lab No : 202309001986

Patient Episode : H18000001080

Collection Date : 09 Sep 2023 09:46

Referred By : HEALTH CHECK MGD

Reporting Date : 09 Sep 2023 15:44

Receiving Date : 09 Sep 2023 12:16

**BIOCHEMISTRY**

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
<b>Serum LIPID PROFILE</b>			
Serum TOTAL CHOLESTEROL Method:Oxidase,esterase, peroxide	172	mg/dl	[<200] Moderate risk:200-239 High risk:>240
TRIGLYCERIDES (GPO/POD)	131	mg/dl	[<150] Borderline high:151-199 High: 200 - 499 Very high:>500
HDL- CHOLESTEROL Method : Enzymatic Immunoimhibition	56.0	mg/dl	[35.0-65.0]
VLDL- CHOLESTEROL (Calculated)	26	mg/dl	[0-35]
CHOLESTEROL, LDL, CALCULATED	90.0	mg/dl	[<120.0] Near/ Borderline High:130-159 High Risk:160-189
Above optimal-100-129			<4.0 Optimal 4.0-5.0 Borderline >6 High Risk
T.Chol/HDL.Chol ratio(Calculated)	3.1		
LDL.CHOL/HDL.CHOL Ratio(Calculated)	1.6		<3 Optimal 3-4 Borderline >6 High Risk

Note:  
Reference ranges based on ATP III Classifications.

Lipid profile is a panel of blood tests that serves as initial broad medical screening tool for abnormalities in lipids, the results of this tests can identify certain genetic diseases and determine approximate risks for cardiovascular disease, certain forms of pancreatitis and other diseases

Name : MR PANKAJ SINGH

Age : 34 Yr(s) Sex :Male

Registration No : MH011292080

Lab No : 202309001986

Patient Episode : H18000001080

Collection Date : 09 Sep 2023 09:46

Referred By : HEALTH CHECK MGD

Reporting Date : 09 Sep 2023 15:43

Receiving Date : 09 Sep 2023 12:16

### BIOCHEMISTRY

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
<b>KIDNEY PROFILE</b>			
Specimen: Serum			
<b>UREA</b>	<b>9.7 #</b>	<b>mg/dl</b>	<b>[15.0-40.0]</b>
<i>Method: GLDH, Kinatic assay</i>			
<b>BUN, BLOOD UREA NITROGEN</b>	<b>4.5 #</b>	<b>mg/dl</b>	<b>[8.0-20.0]</b>
<i>Method: Calculated</i>			
CREATININE, SERUM	0.76	mg/dl	[0.70-1.20]
<i>Method: Jaffe rate-IDMS Standardization</i>			
URIC ACID	4.5	mg/dl	[4.0-8.5]
<i>Method:uricase PAP</i>			
<b>SODIUM, SERUM</b>	<b>132.90 #</b>	<b>mmol/L</b>	<b>[136.00-144.00]</b>
POTASSIUM, SERUM	4.76	mmol/L	[3.60-5.10]
<b>SERUM CHLORIDE</b>	<b>100.0 #</b>	<b>mmol/L</b>	<b>[101.0-111.0]</b>
<i>Method: ISE Indirect</i>			
eGFR (calculated)	119.0	ml/min/1.73sq.m	[>60.0]

#### Technical Note

eGFR which is primarily based on Serum Creatinine is a derivation of CKD-EPI 2009 equation normalized to 1.73 sq.m BSA and is not applicable to individuals below 18 years. eGFR tends to be less accurate when Serum Creatinine estimation is indeterminate e.g. patients at extremes of muscle mass, on unusual diets etc. and samples with severe Hemolysis Icterus / Lipemia.

**Name** : MR PANKAJ SINGH  
**Registration No** : MH011292080  
**Patient Episode** : H18000001080  
**Referred By** : HEALTH CHECK MGD  
**Receiving Date** : 09 Sep 2023 12:16

**Age** : 34 Yr(s) Sex :Male  
**Lab No** : 202309001986  
**Collection Date** : 09 Sep 2023 09:46  
**Reporting Date** : 09 Sep 2023 15:44

**BIOCHEMISTRY**

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
<b>LIVER FUNCTION TEST</b>			
BILIRUBIN - TOTAL <i>Method: D P D</i>	0.43	mg/dl	[0.30-1.20]
BILIRUBIN - DIRECT <i>Method: DPD</i>	0.08	mg/dl	[0.00-0.30]
INDIRECT BILIRUBIN(SERUM) <i>Method: Calculation</i>	0.35	mg/dl	[0.10-0.90]
TOTAL PROTEINS(SERUM) <i>Method: BIURET</i>	7.60	gm/dl	[6.60-8.70]
ALBUMIN (SERUM) <i>Method: BCG</i>	4.55	g/dl	[3.50-5.20]
GLOBULINS (SERUM) <i>Method: Calculation</i>	3.10	gm/dl	[1.80-3.40]
PROTEIN SERUM (A-G) RATIO <i>Method: Calculation</i>	1.49		[1.00-2.50]
AST(SGOT) (SERUM) <i>Method: IFCC W/O P5P</i>	27.00	U/L	[0.00-40.00]
ALT(SGPT) (SERUM) <i>Method: IFCC W/O P5P</i>	60.20	U/L	[17.00-63.00]
<b>Serum Alkaline Phosphatase</b> <i>Method: AMP BUFFER IFCC)</i>	<b>149.0 #</b>	<b>IU/L</b>	<b>[32.0-91.0]</b>
GGT	46.0	U/L	[7.0-50.0]

Name : MR PANKAJ SINGH

Age : 34 Yr(s) Sex :Male

Registration No : MH011292080

Lab No : 202309001986

Patient Episode : H18000001080

Collection Date : 09 Sep 2023 09:46

Referred By : HEALTH CHECK MGD

Reporting Date : 09 Sep 2023 15:44

Receiving Date : 09 Sep 2023 12:16

**BIOCHEMISTRY**

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
------	--------	------	-------------------------------

Liver function test aids in diagnosis of various pre hepatic, hepatic and post hepatic causes of dysfunction like hemolytic anemia's, viral and alcoholic hepatitis and cholestasis of obstructive causes.

The test encompasses hepatic excretory, synthetic function and also hepatic parenchymal cell damage. LFT helps in evaluating severity, monitoring therapy and assessing prognosis of liver disease and dysfunction.

Page 5 of 7

-----END OF REPORT-----



**Dr. Alka Dixit Vats**  
Consultant Pathologist

Name : MR PANKAJ SINGH

Age : 34 Yr(s) Sex :Male

Registration No : MH011292080

Lab No : 202309001987

Patient Episode : H18000001080

Collection Date : 09 Sep 2023 09:46

Referred By : HEALTH CHECK MGD

Reporting Date : 09 Sep 2023 15:44

Receiving Date : 09 Sep 2023 09:46

**BIOCHEMISTRY**

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
<b>GLUCOSE-Fasting</b> Specimen: Plasma GLUCOSE, FASTING (F) Method: Hexokinase	105.0	mg/dl	[70.0-110.0]

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

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, fibro sarcoma), infant of a diabetic mother enzyme deficiency diseases(e.g.galactosemia),  
Drugs-  
insulin, ethanol, propranolol, sulfonylureas, tobutamide, and other oral hypoglycemic agents.

Page 6 of 7

-----END OF REPORT-----

**Dr. Alka Dixit Vats**  
Consultant Pathologist

Name : MR PANKAJ SINGH

Age : 34 Yr(s) Sex :Male

Registration No : MH011292080

Lab No : 202309001988

Patient Episode : H18000001080

Collection Date : 09 Sep 2023 15:32

Referred By : HEALTH CHECK MGD

Reporting Date : 09 Sep 2023 18:11

Receiving Date : 09 Sep 2023 15:32

**BIOCHEMISTRY**

TEST	RESULT	UNIT	BIOLOGICAL REFERENCE INTERVAL
------	--------	------	-------------------------------

**PLASMA GLUCOSE**

Specimen:Plasma

GLUCOSE, POST PRANDIAL (PP), 2 HOURS	100.0	mg/dl	[80.0-140.0]
--------------------------------------	-------	-------	--------------

Method: Hexokinase

Note:

Conditions which can lead to lower postprandial glucose levels as compared to fasting glucose are excessive insulin release, rapid gastric emptying, brisk glucose absorption , post exercise

Page 7 of 7

-----END OF REPORT-----



**Dr. Alka Dixit Vats**  
Consultant Pathologist