

A unit of Aarthi Scans and Labs

Patient Name: MR. MALLIKARJUNA GOWDA

Age / Gender: 40 years / Male

Patient ID: 124127

Referral : MediWheel

Collection Time : Mar 10, 2023, 09:56 a.m. **Reporting Time :** Mar 10, 2023, 02:53 p.m.

Sample ID:

001306923P

,	001300923F			
Test Description	Value(s)	Unit(s)	Reference Range	
COMPLETE BLOOD COUNT (CBC)				
Hemoglobin (Hb)	12.3	gm/dL	13.5 - 18.0	
Erythrocyte (RBC) Count	4.7	mil/cu.mm	4.7 - 6.0	
Packed Cell Volume (PCV)	36.6	%	42 - 52	
Mean Cell Volume (MCV)	77.87	fL	78 - 100	
Mean Cell Haemoglobin (MCH)	26.17	pg	27 - 31	
Mean Corpuscular Hb Concn. (MCHC)	33.61	g/dL	32 - 36	
Red Cell Distribution Width (RDW)	13.0	%	11.5 - 14.0	
Total Leucocytes (WBC) Count	6640	cell/cu.mm	4000-10000	
Neutrophils	56	%	40 - 80	
Lymphocytes	33	%	20 - 40	
Monocytes	5	%	2 - 10	
Eosinophils	5	%	1 - 6	
Basophils	1	%	1-2	
Absolute Neutrophil Count	3718.40	/c.mm	2000 - 7000	
Absolute Lymphocyte Count	2191.20	/c.mm	1000 - 3000	
Absolute Monocyte Count	332	/c.mm	200 - 1000	
Absolute Eosinophil Count	332	/c.mm	20 - 500	
Absolute Basophils Count	66.40	/c.mm	20 - 100	
Platelet Count	330	10^3/ul	150 - 450	
Mean Platelet Volume (MPV)	10.1	fL	7.2 - 11.7	
PCT	0.33	%	0.2 - 0.5	
PDW	11.8	%	9.0 - 17.0	
ESR	35.0	mm/hr	13.5 - 18.0	

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URINE COMPLETE ANALYSIS,				
	Physical Ex	<u>amination</u>		
Quantity	25	ml	-	
Colour	Pale Yellow		Pale yellow/Yellow	
Appearance	Clear		Clear	
Specific Gravity	1.020		1.005-1.025	
рН	6.5		5.0 - 8.0	
Deposit	Absent		Absent	
	Chemical Ex	<u>kamination</u>		
Protein	Absent		Absent	
Sugar	Absent		Absent	
Ketones	Absent		Absent	
Bile Salt	Absent		Absent	
Bile Pigment	Absent		Absent	
Urobilinogen	Normal		Normal	
	Microscopic Exa	mination (/hpf)		
Pus Cell	2-4		Upto 5	
Epithelial Cells	0 - 1		Upto 5	
Red Blood Cells	Absent		Absent	
Casts	Absent		Absent	
Crystals	Absent		Absent	
Amorphous Deposit	Absent		Absent	
Yeast Cells	Absent		Absent	
Bacteria	Absent		Absent	

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STOOL ROUTINE ANALYSIS			
Color	Brownish		Brown
Consistency	Solid		Solid - Semi solid
Reaction (pH) Method : Methyl Red & Bromothymol Blue	Alkaline		Acidic - Alkaline
Mucous	Absent		Absent
Blood	Absent		Absent
Pus cells	1-2/hpf	/hpf	Few
Epithelial cells	1-2/hpf	/hpf	
RBC	Absent	/hpf	Absent
Ova	Not found	/hpf	Absent
Cyst	Not found	/hpf	Absent
Starch granules	Absent	/hpf	None to small amount
Vegetable cells	Absent	/hpf	
Fat globules	Absent	/hpf	Absent
Others Method : Microscopy (Concentration technique)	Nil	/hpf	







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Test Description	Value(s)	Unit(s)	Reference Range	
BLOOD GROUP & RH TYPING				
Blood Group (ABO typing) Method : Manual-Hemagglutination	"O"			
RhD Factor (Rh Typing) Method : Manual hemagglutination	Positive			

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Test Description Value(s) Unit(s) Reference Range

urine glucose

urine glucose (R)

Negative

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Test Description	Value(s)	Unit(s)	Reference Range	
Glycosylated HbA1c				
HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD	5.7	%		
Method : (HPLC, NGSP certified)				
Estimated Average Glucose :	116.89	mg/dL	-	
Interpretation				
As per American Diabetes Association (ADA)				
Reference Group	HbA1c in %			
Non diabetic adults >=18 years	<5.7			
At risk (Prediabetes)	5.7 - 6.4			
Diagnosing Diabetes	>= 6.5			
	Age > 19 yea	rs		
	Goal of therap	oy: < 7.0		
Therapeutic goals for glycemic control	Action sugges	sted: > 8.0		
	Age < 19 yea	rs		
	Goal of therap	oy: <7.5		







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			0010003201	
Test Description	Value(s)	Unit(s)	Reference Range	
THYROID PROFILE TEST - TOTAL				
T3-Total	119.0	ng/dL	60 - 200	
T4-Total	13.5	ug/dL	4.52 - 12.8	
TSH-Ultrasensitive Method : CLIA	0.35	uIU/mL	0.32 - 5.5	

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Test Description	Value(s)	Unit(s)	Reference Range
LIPID PROFILE			
Cholesterol-Total	234	mg/dL	Desirable level < 200
Method : Spectrophotometry			Borderline High 200-239
			High >or = 240
Triglycerides	246	mg/dL	Normal: < 150
Method : Serum, Enzymatic, endpoint			Borderline High: 150-199
			High: 200-499
			Very High: >= 500
HDL Cholesterol	36	mg/dL	Normal: > 40
Method : Serum, Direct measure-PEG			Major Risk for Heart: < 40
LDL Cholesterol	148.80	mg/dL	Optimal < 100
Method : Enzymatic selective protection			Near / Above Optimal 100-129
			Borderline High 130-159
			High 160-189
			Very High >or = 190
VLDL Cholesterol	49.20	mg/dL	6 - 38
Method : Serum, Enzymatic			
CHOL/HDL Ratio	6.50		3.5 - 5.0
Method : Serum, Enzymatic			
LDL/HDL Ratio	4.13		2.5 - 3.5
Method : Serum, Enzymatic			
Note:			
8-10 hours fasting sample is required.			







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RENAL PROFILE				
Urea	13.6	mg/dL	19-42	
Method : Uricase				
Blood Urea Nitrogen-BUN	6.35	mg/dL	9-20	
Method : Serum, Urease				
Creatinine	0.85	mg/dL	0.66-1.25	
Method : Serum, Jaffe				
Uric Acid	6.7	mg/dL	3.5-8.5	
Method : Serum, Uricase				
Remark:				

In blood, Urea is usually reported as BUN and expressed in mg/dl. BUN mass units can be converted to urea mass units by multiplying by 2.14.

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Test Description	Value(s)	Unit(s)	Reference Range	
LIVER FUNCTION TEST				
Total Protein	7.2	g/dL	6.3-8.2	
Method : Serum, Biuret, reagent blank end point				
Albumin	3.8	g/dL	3.5-5.0	
Method : Serum, Bromocresol green				
Globulin	3.40	g/dL	1.8 - 3.6	
Method : Serum, EIA				
A/G Ratio	1.12		1.2 - 2.2	
Method : Serum, EIA				
Bilirubin - Total	0.2	mg/dL	0.3-1.2	
Method : Serum, Jendrassik Grof				
Bilirubin - Direct	0.1	mg/dL	< 0.2	
Method : Serum, Diazotization				
Bilirubin - Indirect	0.10	mg/dL	0.1 - 1.0	
Method : Serum, Calculated				
SGOT	27	U/L	17-59	
Method : Serum, UV with P5P, IFCC 37 degree				
SGPT	30	U/L	21-72	
Method : Serum, UV with P5P, IFCC 37 degree				
Alkaline Phosphatase	73	U/L	30 - 120	
Method : PNPP-AMP Buffer/Kinetic				
GGT-Gamma Glutamyl Transpeptidae	28	U/L	< 55	
Method : Serum, G-glutamyl-carboxy-nitoanilide				

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Test Description	Value(s)	Unit(s)	Reference Range	
PSA-Total (Prostate-specific antigen-Total	<u>D</u>			
PSA Profile *				
PSA (Prostate Specific Antigen)-Total	0.45	ng/mL	0 - 4.0	
Method : Serum, CLIA				
Interpretation:				
1. Increased levels are noted in Prostate cancer	r, Bengin prostatio	hypertrophy, Prost	atitis	
PSA (Prostate-Specificantigen)-Free *	-	ng/mL	0.0 - 0.5	

Interpretation & Remarks:

Method: Serum, CLIA

- Normal results do not eliminate the possibility of prostate cancer.
- · Values obtained with different assay methods or kits may be different and cannot be used interchangeably.
- Tumor markers are not specific for malignancy. Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease
- Specimens drawn from patients undergoing prostate manipulation, especially needle biopsy and transurethral resection, may show erroneously high prostatic-specific antigen (PSA) results. Care should be taken that specimens are drawn before these procedures are performed.
- The percentage of free PSA can be used to estimate how likely it is that a biopsy will show cancer.
- If the percentage of free PSA is higher than 25%, the likelihood of prostate cancer is about 8%.
- If the percentage of free PSA is less than 10%, then the likelihood of prostate cancer rises to 56%.

Free PSA / Total PSA % - -

Method : Serum

Interpretation

- When total prostate-specific antigen (PSA) concentration is <2.0 ng/mL, the probability of prostate cancer in asymptomatic men is low, further testing and free PSA may provide little additional information. When total PSA concentration is >10.0 ng/mL, the probability of cancer is high and prostate biopsy is generally recommended.
- The total PSA range of 4.0 to 10.0 ng/mL has been described as a diagnostic "gray zone," in which the free:total PSA ratio helps to determine the relative risk of prostate cancer (see table below). Therefore, some urologists recommend using the free:total ratio to help select which men should undergo biopsy. However even a negative result of prostate biopsy does not

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Test Description Value(s) Unit(s) Reference Range

rule-out prostate cancer. Up to 20% of men with negative biopsy results have subsequently been found to have cancer. Based on free:total PSA ratio: the percent probability of finding prostate cancer on a needle biopsy by age in years:

Free:total PSA ratio	50-59 years	60-69 years	> or =70 years
< or =0.10	49.2%	57.5%	64.5%
0.11-0.18	26.9%	33.9%	40.8%
0.19-0.25	18.3%	23.9%	29.7%
>0.25	9.1%	12.2%	15.8%

Cautions

- · Normal results do not eliminate the possibility of prostate cancer.
- · Values obtained with different assay methods or kits may be different and cannot be used interchangeably
- Tumor markers are not specific for malignancy. Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease.

Interfering factors:

- Prostatic massage
- Proctoscopy
- Prostatic biopsy
- Prostate cancer patients receiving treatment with antiandrogens and luteinizing hormone-releasing factor agonists may exhibit
 markedly decreased levels of PSA. Also, men treated for benign prostatic hyperplasia with inhibitors of 5-alpha-reductase
 (finasteride) may demonstrate a significant reduction in PSA levels compared to values before treatment. Care should be taken
 in interpreting values for these individuals.
- In patients receiving therapy with high biotin doses (ie, >5 mg/day), no sample should be taken until at least 8 hours after the last biotin administration.

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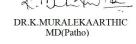
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			0010000201	
Test Description	Value(s)	Unit(s)	Reference Range	
GLUCOSE (F)				
Glucose fasting Method : GOD-POD	94	mg/dL	Normal: 70 - 120	



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GLUCOSE (PP)			
Blood Glucose-Post Prandial Method: GOD-POD	128.0	mg/dL	80 - 140

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