





4.5 - 5.5

40 - 49

83 - 101

7.5 - 10.3

Passport No :	LABORATORY TEST REF	PORT
Patient Information	Sample Information	Client/Location Information
Name : Lyubochka Svetka Sex/Age : Male / 41 Y 01-Feb-1982 Ref. ld : Ref. By :	Lab Id : 02232160XXXX Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : EDTA Blood	Client Name : Sterling Accuris Buddy Location : Approved on : 20-Feb-2023 11:09 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Pald
	Complete Blood Count	
Test Res Hemoglobin Colorimetric 14.5		Biological Ref. Interval 13.0 - 16.5

MCH 30.2 pg 27.1 - 32.5 Calculated **MCHC** 32.5 - 36.7 33.4 g/dL Calculated **RDW CV** 13.60 Total WBC and Differential Count WBC Cou

million/cmm

%

fL

4.79

43.3

90.3

Electrical impedance

Calculated

Derived

Differential Count						Absolut	e Count	
Neutrophils	Microscopic	73	%	40 - 80		7716	/cmm	2000 - 6700
Lymphocytes	Microscopic	19	%	20 - 40	h	2008	/cmm	1100 - 3300
Eosinophils	Microscopic	02	%	1 - 6		211	/cmm	00 - 400
Monocytes	Microscopic	06	%	2 - 10		634	/cmm	200 - 700
Basophils	Microscopic	00	%	0 - 2		0	/cmm	0 - 100
Platelet Count	Electrical impedance	150000	/cmr	n		150000 -	410000	

Peripheral Smear Examination

RBC Count

Hematocrit

MCV

RBC Morphology Normochromic Normocytic

Calculated

WBC Morphology WBCs Series Shows Normal Morphology

H 14.00

Platelets Morphology Platelets are adequate with normal morphology.

fL

Parasites Malarial parasite is not detected.

Erythrocyte Sedimentation Rate

ESR 0 - 14 7 mm/1hr Capillary photometry

MPV

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Passport No :	LABORATORY TEST REI	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. Id : Ref. By :	Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : EDTA Blood, Serum	Location : Approved on : 20-Feb-2023 13:33 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Blood Group

Test Result Unit Biological Ref. Interval

ABO Type "A"

Rh (D) Type Positive

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Passport No :	LABORATORY TE	ST REPORT			
Patient Information	Sample Informati	ion	Client/Location Information		
Name : Mr. Hardik Sompura	Lab ld : 0223216	00126 Client Na	me : Sterling Accuris Buddy		
Sex/Age : Male / 41 Y 01-Feb-19	Registration on : 20-Feb-20 Collected at : non SAWE	Location	:		
Ref. ld : Ref. By :	Collected on : 20-Feb-20 Sample Type : Serum	Approved	n : 28-Feb-2023 10:26		
Lipid Profile					
Test	Result	Unit	Biological Ref. Interval		

Cholesterol Cholesterol oxidase – Peroxidase method	189.0	mg/dL	Desirable : <200 Borderline High : 200-239 High : >240
Triglyceride Ezymatic (Lipase/GK/GPO/POD)	H 168.0	mg/dL	Normal : <150 Borderline : 150-199 High : 200-499 Very High : >500
HDL Cholesterol PTA In a Ci2 Direct LDL Direct measured	60.0 H 100.39	mg/dL mg/dL	Low: <40.0 High: >60.0 Optimal: <100 Near to above Optimal: 100–129 Borderline High: 130-159

Sa_{33.60} Comg/de High: 160–189 Very High: =190 15 - 35 VLDL Calculated 3.1

CHOL/HDL Ratio

Up to 5.0

LDL/HDL Ratio 1.7 Up to 3.5

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Passport No :	LABORATORY TEST REPO			PORT	
Patient Information	Sar	mple	Information		Client/Location Information
Name : Mr. Hardik Sompura	Lab Id	:	022321600126	Client Name	: Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982			20-Feb-2023 09:10	Location	:
Ref. By :	Collected at Collected on Sample Typ	ı :	non SAWPL 20-Feb-2023 08:53 Fluoride plasma	Approved on Printed On Process At	: 20-Feb-2023 11:45 Status : Final : 28-Feb-2023 10:26 : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Biochemistry

Test Result Unit Biological Ref. Interval **Fasting Blood Sugar** H 141.0 mg/dL 74 - 106

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Passport No :	LABORATORY TEST REP	ORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. Id :	Collected at . Hon SAWPL I	Location : Approved on : 20-Feb-2023 11:33 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

HbA1c (Glycosylated Hemoglobin)

Test	Result	Unit	Biological Ref. Interval
HbA1c High Performance Liquid Chromatography	H 7.10	%	For Screening: Diabetes: >6.5% Pre-Diabetes: 5.7% - 6.4% Non-Diabetes: < 5.7%

For Diabetic Patient: Poor Control: > 7.0 % Good Control: 6.0-7.0 %

Mean Blood Glucose 157.07

Explanation:-

- Total haemoglobin A1 c is continuously synthesized in the red blood cell throught its 120 days life span. The concentration of HBA1c in the cell reflects the average blood glucose concentration it encounters.
- The level of HBA1c increases proportionately in patients with uncontrolled diabetes. It reflects the average blood glucose concentration over an extended time period and remains unaffected by short-term fluctua
- The measurement of HbA1c can serve as a convenient test for er aluating the adequacy of diabetic control and in preventing various diabetic complications. Because the average half-life of a red blood cell is sixty days, HbA1c has been accepted as a measurement which reflects the mean daily blood glucose concentration, better than fasting blood glucose determination, and the degree of carbohydrate imbalance over the preceding
- It may also provide a better index of control of the diabetic patient without resorting to glucose loading procedures.

HbA1c assay Interferences:

Erroneous values might be obtained from samples with abnormally elevated quantities of other Haemoglobins as a result of either their simultaneous elution with HbA1c (HbF) or differences in their glycation from that of HbA (HbS).

Reference: ADA Guideline 2023

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Passpor	t No :	LABORATORY TEST REP			PORT	
	Patient Information	Samı	ple	Information		Client/Location Information
Name	: Mr. Hardik Sompura	Lab Id	:	022321600126	Client Name	: Sterling Accuris Buddy
Sex/Age	: Male / 41 Y 01-Feb-1982			20-Feb-2023 09:10 non SAWPL	Location Approved or	: n : 20-Feb-2023 11:38 Status : Final
Ref. Id Ref. By	:	Collected on Sample Type		20-Feb-2023 08:53 Serum	Printed On Process At	: 28-Feb-2023 10:26 : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Thyroid Function Test

Test	Result	Unit	Biological Ref. Interval
T3 - Triiodothyronine	1.01	ng/mL	0.58 - 1.59
T4 - Thyroxine	7.84	mg/mL	4.87 - 11.72
TSH - Thyroid Stimulating Hormone	0.8199	microIU/mL	0.35 - 4.94

TSH	T3/FT3	T4/FT4	Suggested Interpretation for the Thyroid Function Tests Pattern
Within Range	Decreased	Within Range	 Isolated Low T3-offen seen in elderly & associated Non-Thyroidal illness. In elderly the drop in T3 level can be upto 25%
Raised	Within Range	Within Range	Isolated High TSM especially inthe range of 4.7 to 15 mill/limits commonly associated with shyrid layed & Biological TSM Variability. Subdimical Autoimmune Hypothybid in Internimed T4 therapy for the both hold sm Hecovery phase other North fryrold unliness.
Raised	Decreased	Decreased	Chronic autoimmune Thyroiditis Post thyroidectomy, Post radioiodine Hypothyroid phase of transient thyroiditis
Raised or Within Range	Raised	Raised or Within range	Interfering antibodies to thyroid hormones (anti-TPO antibodies) intermittent T4 therapy or T4 overdose Drugin of the ence-time darks. Programme and the ence-time darks. Programme and the ence-time darks.
Decreased	Raised or within Range	Raised or within Raise	Inclated Low TSM - especially in the range of 0.1 to 0.4 eVen seen in elderly & associated with Non-Thyroidal lines - Subclinical Representations - Thyroxine ingestion
Decreased	Decreased	Decreased	Central Hypothyroidism Non-Thyroidal illness Recent treatment for Hyporthyroidism (TSH remains suppressed)
Decreased	Raised	Raised	Primary Hyperthyroidism (Graves disease), Multinodular goitre Toxic nodule Transient thyroiditis:Postpartum, Silent (lymphocytic), Postviral (granulomatous, subacute, DeQuervain'a) Gestational thyrotoxicosis with hyperemesis gravidarum
Decreased or within range	Raised	Within Range	- T3 toxicosis - Non-Thyroidal illness

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Passport No :	LABORATORY TEST REP	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. ld : Ref. By :	Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : Urine	Location : Approved on : 20-Feb-2023 12:40 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Biochemistry

Test	Result	Unit	Biological Ref. Interval
Microalbumin (per urine volume)	10.50	mg/L	< 16.7

In random urine specimens, normal urinary albumin excretion is below 17 mg/g creatinine for males and below 25 mg/g creatinine for females.(3) Microalbuminuria is defined as an albumin:creatinine ratio of 17 to 299 for males and 25 to 299 for females.

A ratio of albumin:creatinine of 300 or higher is indicative of overt proteinuria.

Due to biologic variability, positive results should be confirmed by a second, first-morning random or 24-hour timed urine specimen. If there is discrepancy, a third specimen is recommended. When 2 out of 3 results are in the microalbuminuria range, this is evidence for incipient nephropathy and warrants increased efforts at glucose control, blood pressure control, and institution of therapy with an angiotensin-converting-enzyme (ACE) inhibitor (if the patient can tolerate it).

Reference

- ng <mark>a</mark>nd management of microalbuminuria in patients with diabetes mellitus: recommendations ntific Advisory Board of t National Kidney Foundation from an ad hoc committee or Council on Diabetes Mellitus of the National Kidney Foundation. Am J Kidney Dis 1995;25:107-112
- 2. Krolewski AS, Laffel LM, Krolewski M, et al: Glycosylated hemoglobin and the risk of microalbuminuria in patients with insulin-dependent diabetes mellitus. N Engl J Med 1995;332:1251-1255
- Zelmanovitz T, Gross JL, Olivera JR, al al: The receiver operating characteristics curve in the evaluation of a random urine specimen as a screening test for diabetic nephropathy. Diabetes Care

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Passpor	rt No :	LABORATORY TEST REP		PORT
	Patient Information	Samp	ple Information	Client/Location Information
Name	: Mr. Hardik Sompura	Lab Id	: 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age	: Male / 41 Y 01-Feb-1982		n: 20-Feb-2023 09:10 : non SAWPL	Location
Ref. Id Ref. By	:	Collected on	: 20-Feb-2023 08:53 : Serum	Approved on : 20-Feb-2023 11:41 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Р	r	^	ŀ	iد	n
	п	01	П	-11	п

Test	Result	Unit	Biological Ref. Interval
Total Protein Copper tartrate to colour complex	7.00	g/dL	6.3 - 8.2
Albumin Bromocresol Green Method	4.20	g/dL	3.5 - 5.0
Globulin Calculated	2.80	g/dL	2.3 - 3.5
A/G Ratio	1.50		1.3 - 1.7

Total Bilirubin Azobilirubia curomophores Conjugated Bilirubin Cationic Mordant Binding	en oro	mg/dL mg/dL	0.2 - 1/3
Unconjugated Bilirubin Cationic Mordant Binding	0.20	mg/dL	0.0 - 1.1
Delta Bilirubin Calculated	Sand	OOX	0.0 - 0.2

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Passport No :	LABORATORY TEST REP	ORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. ld : Ref. By :	i Collected at : non SavvPi i	Location : Approved on : 20-Feb-2023 11:29 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Iron Studies

Test	Result	Unit	Biological Ref. Interval
lron Pyridyl azo Dye	103.00	micro g/dL	49 - 181
Total Iron Binding Capacity (TIBC)	352.00		261 - 462
Transferrin Saturation Calculated	29.26	%	Children: >16 Adult: 20 - 50

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Passport No :	LABORATORY TEST RE	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. ld :	Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53	Location : Approved on : 20-Feb-2023 11:38 Status : Final
Ref. By :	Sample Type : Serum	Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Immunoassay

Test	Result	Unit	Biological Ref. Interval
Homocysteine, Serum	H 23.86	micromol/L	6.0 - 14.8

Summary and Uses:

- Total Hcy is a thiol-containing amino acid, produced by the intracellular demethylation of methionine to cysteine.
- Elevated levels of t Hcy may be used to exclude or confirm deficiencies of vitamin B12 or folate.
- It is recommended to test in patients using medications that interfere with folate status (methotrexate, antiepileptics), vegetarians without B12 supplementations, unexplained anemia, peripheral neuropathy or my expathy, recurrent spontaneous abortions or infertility.
- Testing also recommended for patients 40 years of age with coronary artery disease to exclude homocystinuria.
 - Elevations in they levels have also been used as an independent risk factor of coronary or cerebral vascular disease. Treatment of moderate hyperhomocystinemia with folic acid supplementation for primary and secondary cardiovascular protection has met with inconsistent results and at present cannot be routinely recommended.

Limitations:

- The plasma must be seprated immediately on collection to avoid continuous synthesis of Hcy by red cells.
- Samples must be immediately stored on ice and serum centrifuged immediately before a complete clot is formed.
- Certain drugs, such as anticonvulsants, methotrexate, or nitrous oxide, may interfere with the assay.
- Cigarette smoking and coffee consumption increase tHcy levels.
- Intraindividual variability is approximately 8%; it can be as much as 25% in patients with hyperhomocystinemia.
- Generally, a single measurement of tHcy is considered adequate.

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Passport No :	LABORATORY TEST REP	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982	Collected at . Holl SAVVEL	Location : Approved on : 20-Feb-2023 11:10 Status : Final
Ref. Id : Ref. By :	Collected on : 20-Feb-2023 08:53 Sample Type : Serum	Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Biochemistry

	Diocheim	isti y	
Test	Result	Unit	Biological Ref. Interval
Creatinine, Serum Creatinine Amidohydrolase	0.83	mg/dL	0.66 - 1.25
Urea Urease, Colorimetric	L 18.0	mg/dL	19.3 - 43.0
Blood Urea Nitrogen	L 8.41	mg/dL	9.0 - 20.0
Uric Acid Uricase	4.90	mg/dL	3.5 - 8.5
Calcium Arse a 20 III SG PT UV with PSR IFCS	9,10 48.0	mg/dl/ U/L/	8.4 - 10.2 0 - 50
SGOT UV with P5P	27.0	U/L	17 - 59
	Electro	lytes	•
Sodium (Na+) Direct- ISE	143.00	mmol/L	136 - 145
Potassium (K+) Direct- ISE	4.90	mmol/L	3.5 - 5.1
Chloride (CI-) Direct- ISE	105.0	mmol/L	98 - 107

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Passport No :		LABORATORY TEST REP		PORT		
Patient I	nformation	Sam	ple	Information		Client/Location Information
Name : Mr. H	ardik Sompura	Lab Id	:	022321600126	Client Name	e : Sterling Accuris Buddy
Sex/Age : Male	741 Y 01-Feb-1982	_		20-Feb-2023 09:10 non SAWPL	Location	:
Ref. ld :		Collected on	:	20-Feb-2023 08:53 Serum	Approved o Printed On Process At	: 28-Feb-2023 10:26

Immunoassay

Biological Ref. Interval Test Result Unit 8.98 25(OH) Vitamin D ng/mL Deficiency: <10 Insufficiency: 10 - 30 Sufficiency: 30 - 100 Toxicity : >100

Vitamin D is a fat soluble vitamin and exists in two main forms as cholecalciferol(vitamin D3) which is synthesized in skin from 7-dehydrocholesterol in response to sunlight exposure & Ergocalciferol(vitamin D2) present mainly in dietary sources. Both cholecalciferol & Ergocalciferol are converted to

Interpretation:

rient DVS API

Decreased in

- Malabsorption
- Steatorrhea
- Dietary osteomalacia, anticonvulsant osteomalacia
 Biliary and portal cirrhosis
 Thyrotoxicosis

- Pancreatic insufficiency
- Celiac disease
- Rickets
- Alzheimer disease

Limitations:

More recently, it has become clear that receptors for vitamin D are present in a wide variety of cells and that this hormone has biologic effects extending beyond the control of mineral metabolism. Vitamin D deficiency is not clear. Levels needed to prevent rickets and osteomalacia (15 ng/mL) are lower than those that dramatically suppress parathyroid hormone levels (20–30 ng/mL). In turn, those levels are lower than levels needed to optimize intestinal calcium absorption (34 ng/mL). Neuromuscular peak performance is associated with levels approximately 38 ng/mL. A recent study states that increasing mean baseline levels from 29 to 38 ng/mL was associated with a 50% lower risk for colon cancer and levels of 52 ng/mL with a 50% reduction in the incidence of breast cancer. It is recommended to have clinical correlation with serum 25(OH)vitamin D, serum calcium, serum PTH & serum alkaline phosphatase.

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Passport No :	LABORA [*]	TORY TEST REP	ORT	ORT	
Patient Information	Sample Information			Client/Location Information	
Name : Mr. Hardik Sompura	Lab ld :	022321600126	Client Name	: Sterling Accuris Buddy	
Sex/Age : Male / 41 Y 01-Feb-1982		: 20-Feb-2023 09:10 : non SAWPL	Location	:	
Ref. ld : Ref. By :	Collected on :	: 20-Feb-2023 08:53 : Serum	Approved on Printed On Process At	: 20-Feb-2023 12:04 Status : Final: 28-Feb-2023 10:26: 1. NRL SAWPL Gujarat Ahmedabad Paldi	

Immunoassay

Test Result Unit Biological Ref. Interval L < 148 187 - 833 Vitamin B12 pg/mL

Vitamin B12 is essential in DNA synthesis, hematopoiesis, and CNS integrity.

Interpretation:

- Increased In Chronic granulocytic leukemia, COPD and Chronic renal failure, Leukocytosis, Liver cell damage (hepatitis, cirrhosis), Obesity and Severe CHF, Polycythemia vera, Protein malnutrition.
- : Abnormalities of cobalamin transport or metabolism , Bacterial overgrowth , Crohn disease , Dietary deficiency (e.g. in vegetarians) , Diphyllobothrium (fish tapeworm) infestation , Gastric or small intestine urge ry , Hypochlorhydria , Inflammatory bo ntrins malabsorption and

Limitations:

- ase vit<mark>a</mark>min B12 levels. On er hand , alcoh , amin cholestyramine, cimetidine, colchicines, metformin, neomycin, oral contraceptives, ranitidine, and triamterene decrease vitamin B12 levels.
- The evaluation of macrocytic anemia requires measurements of both vitamin B12 and folate levels; ideally they should be measured simultaneously.
- Specimen collection soon after blood transfusion can falsely increase vitamin B12 levels.
- Patients taking vitamin B12 supplementation may have misleading re
- A normal serum concentration of B12 does not rule out tissue deficiency of vitamin B12. ost sensitive test for B12 deficiency at the cellular level is the assay for MMA. If clinical neasure ment of MMA and homocysteine should be considered, even if serum B12 concentrations are normal.

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	Patient Information	Sample Information	Client/Location Information
Name	: Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age Ref. Id Ref. By	:	Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : Serum	Location : Approved on : 20-Feb-2023 11:38 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Immunoassay

Result Biological Ref. Interval **Test** Unit **PSA-Prostate Specific Antigen, Total** 0.573 ng/mL 0 - 4

PSA is a glycoprotein that is expresses by both normal and neoplastic prostate tissue and is prostate tissue specific and not prostate cancer specific. PSA is constantly expressed in nearly all prastate cancers, although its level of expression on a percell basis is lower than in normal prostate epithelium. The absolute value of serum PSA is useful for determining the extent of prostate cancer and assessing the response to prostate cancer treatment; its use as a screening method to detect prostate cancer is also common.

Interpretation

Increased in

Prostate disease (Cancer, Prostatitis, Benign prostatic hyperplasia, Acute urinary retention)

Manipulations (Cystoscopy, Needle biopsy, Radiation therapy, Indwel

Transurethral resection tati isch

Decreased in

- Castration
- Prostatectomy
- Radiation therapy
- Ejaculation withi 24 48 hours
- 5-alpha-reductase inhibitor reduces PS

Limitations

- PSA has been recommended by the American Cancer Society for use in conjunction with a DRE for early detection of prostate cancer starting at the age of 50 years for men with at least 10 year life expectancy
- PSA levels that are measured repeatedly over time may vary because of biologic variability where the true PSA level in a given man is different on different measurements.
- A change in PSA of >30% in man with a PSA initially below 2.0 ng/mL was likely to indicate a true change beyond normal random variation.

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Passport No :	LABORATORY TEST REP	ORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982	Registration on : 20-Feb-2023 09:10	Location :
Ref. ld : Ref. By :	Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : Serum	Approved on : 20-Feb-2023 12:06 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Immunoassay

Test Result Unit Biological Ref. Interval H 492.30 IU/mL 0 - 87IgE

IgE mediates allergic and hypersensitivity reactions. There is a significant overlap in total IgE between allergic and nonallergic individuals. Interpretation:

- Increased In
 - Atopic diseases
 - Exogenous asthama in approximately 60% of patients
 - Hay fever in approximately 30% of patients and Atopic eczema

Influenced by type of allergen, duration of stimulation. Presence of symptoms, and hyposensitization treatment Par<mark>asitic diseases (e.g. ascariasis, visceral arva migra</mark>ns, hoo<mark>kworm dis</mark>eas Echinococcus infestation onoclonal IgE myeloma

- - Hereditary deficiencies
 - Acquired immunodeficiency
 - Ataxia-telangiectasis
 - Non-IgE myeloma

Limitations:

- A normal level of IgE in serum
- Serum total IgE levels for the majority of individuals with IgE-mediated disease can be expected to be elevated compared to the reference range for healthy adults. However, not all allergic patients exhibit elevated serum total IgE levels.
- Since not all atopic reactions are IgE-mediated, a total IgE result in the reference range should always be interpreted in light of other clinical observations.
- Heterophilic antibodies in human serum can react with the immunoglobulins included in the assay components causing interference with in vitro immunoassays.

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Passport No :	LABORATORY TEST REF	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982	Registration on: 20-Feb-2023 09:10 Collected at: non SAWPL	Location : Approved on : 20-Feb-2023 14:35 Status : Final
Ref. ld : Ref. By :	Collected on : 20-Feb-2023 08:53 Sample Type : Serum	Approved on : 20-Feb-2023 14:35 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

Test	Result	Unit	Biological Ref. Interval		
HIV I & II Ab/Ag with P24 Ag	0.070	S/Co	Non Reactive : <1.0 Reactive : >1.0		
Interpretation	Non Reactive				
HBsAg Chemiluminescence	0.290	S/Co	Non Reactive : <1.0 Reactive: >1.0		
Interpretation	Non Reactive				

Additio Information:

- A NON REACTIVE result implies that no Anti HIV-1 or HIV-2 antibodies have been detected in the sample by this the patient has not been exposed to HIV-1 or HIV-2 infection or the sample has been tested during the WINDOW development of detectable levels of antibodies). mple by this method. nat either
- A PROVISIONALITY REACTIVE / BORDERLINE REACTIVE result suggests possibility of HIV-1 or/and HIV-2 infection. However these results must be verified by confirmatory WESTERN BLOT / HIV PCR method before declaring the patient positive for HIV-1 or HIV-2 infection.

3. Very high levels of IgM Antibodies or Anti-HLA ABC and DR Antibodies can give false positive reaction.

**Pre & Post test counselling for HIV testing is responsibility of reffering Physicia

Dr. Siddharth Thummar

M.D. Pathology

Dr. Sanjeev Shah

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MD Path

Dr. Yash Shah

MD Path

Referred Test

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Passport No :	LABORATORY TEST REI	PORT
Patient Information	Sample Information	Client/Location Information
Name : Mr. Hardik Sompura	Lab ld : 022321600126	Client Name : Sterling Accuris Buddy
Sex/Age : Male / 41 Y 01-Feb-1982 Ref. Id : Ref. By :	Registration on : 20-Feb-2023 09:10 Collected at : non SAWPL Collected on : 20-Feb-2023 08:53 Sample Type : EDTA Blood	Location : Approved on : 20-Feb-2023 14:16 Status : Final Printed On : 28-Feb-2023 10:26 Process At : 1. NRL SAWPL Gujarat Ahmedabad Paldi

HB Electrophoresis By HPLC

Instrument Name: BIORAD VARIANT - II Haemoglobin Testing System

Test	Result	Unit	Biological Ref. Interval
Hb A	∟ 84.4	%	96.8 - 97.8
Hb A2	2.8	%	2.2 - 3.2
P2 Peak	5.5	%	
P3 Peak	5.2	%	
Foetal Hb	0.3	%	0.0 - 1.0

Interpretation Negative for typical beta thalassemia trait.

tation: Interpr

- All res of blood tran lood transfusion. usion is recommen
- In case of haemoglobinopathy, parents or family studies and councelling is advised.
- This test detects beta thalassaemia and haemoglobinopathies, DNA analysis is recommended to rule out alpha thalassaemia and silent
- Linearity range of HbF is 1-40%, However, values in excess of the reportable range have been provided for ease of interpretation.
- Mild to moderate increase in fetal haemogloblin can be seen in some acquired condition like pregnancy, megaloblastic anaemia, Throtoxicosis, Hypoxia, Chronic kidney disease, Recovering marrow, MDS, Aplastic anaemia, PVH, Medications (Hydrocyurea, Erythropoietin) ect.
- P3 window-Above 10% is often indicative of eit er denatured froms of hemogl obins or may suggest a possibility of abnormal haemoglobin variant. Hence, repeat analysis with fresh sample or DNA studies is advised.
- P2 Window-Above 10% is indicative of either glycated haemoglonin requring correlation with diabetic staus or may suggest a possibility of abnormal haemoglobin variant further DNA studies for confirmation.

Dr. Hardik Modi

Dr. Sanjeev Shah

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Dr. Yash Shah

Referred Test

Hematopathologist (G-18097)

MD Path

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Patient Data

Sample ID:

022321600126

Patient ID:

Name:

Physician:

Sex:
DOB:

Comments:

Analysis Data

Analysis Performed: 02/20/2023 13:56:59

Injection Number: 2575
Run Number: 95
Rack ID: 0001

Tube Number: 2
Report Generated: 02/20/20

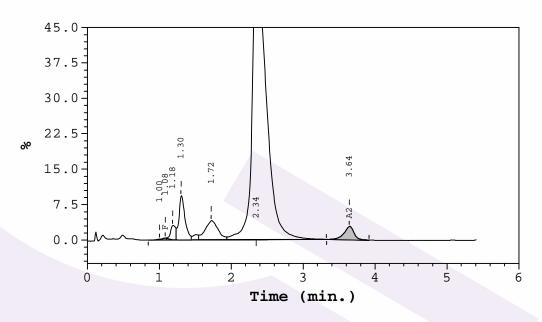
Operator ID:

02/20/2023 14:04:40

	Calibrated		Retention	Peak
Peak Name	Area %	Area %	Time (min)	Area
Unknown		0.1	1.00	2400
F	0.3		1.08	8321
Unknown		1.6	1.18	44109
P2		5.5	1.30	148807
P3		5.2	1.72	139183
Ao		84.4	2.34	2277592
A2	2.8		3.64	76999

F Concentration = 0.3 % A2 Concentration = 2.8 %

Analysis comments: and ox







Client Name : Sterling Accuris Buddy



Client/Location Information

Scan QR code to check report authenticity

LABORATORY TEST REPORT

: 022321600126

Sample Information

Lab Id

Sex/Age : Male / 41 Y 01-Feb-198 Ref. ld : Ref. By :	Registration on : 20-Feb-2023 Collected at : non SAWPL Collected on : 20-Feb-2023 Sample Type : Urine	Approved on	: 20-Feb-2023 11:12 Status: Final : 28-Feb-2023 10:26 : 1. NRL SAWPL Gujarat Ahmedabad Paldi	
Test	Result	Unit	Biological Ref. Interval	
Physical & Chemical (Dip strip) exam	ination_			
Colour	Pale Yellow		Pale Yellow	
Clearity	Clear		Clear	
pH Double indicator	6.0		4.6 - 8.0	
Specific Gravity Polyelectrolyte based reaction	1.030		1.005 - 1.030	
Urine Glucose GOL POD Urine Protein Protein every of indicators Bilirubin Diazo reaction	Present (+) Absent Absent	WS	Absent Absent Absent	
Urobilinogen Modified Ehrlich reaction	Absent		Absent	
Urine Ketone Nitroprusside	Absent		Absent	
Nitrite Nitrite reaction	Absent	UUX	Absent	
Microscopic Examination				
Pus Cells	1-2		Absent	
Red Cells	Nil	/hpf	0 - 2	
Epithelial Cells	1-2	/hpf		
Casts	Absent	/hpf	Absent	
Crystals	Absent	/hpf	Absent	
Amorphous Material	Absent			

Passport No:

Name

Patient Information

: Mr. Hardik Sompura

DR.TEJASWINI DHOTE

M.D. Pathology

Dr. Sanjeev Shah

----- End Of Report --

MD Path

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Dr. Yash Shah

MD Path

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