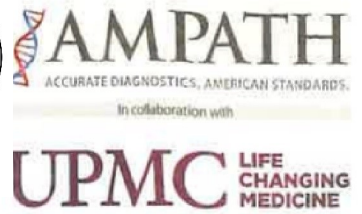


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**LABORATORY REPORT**

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Test Name	Result	Biological Ref. Interval	Unit
<b>Senior Citizens - Female Check</b>			
<b>BIOCHEMISTRY</b>			
<b>Aspartate Aminotransferase (AST/SGOT) (Serum)</b>			
Aspartate Aminotransferase (AST/SGOT) <i>IFCC kinetic</i>	17	<37	U/L
<b>Alanine aminotransferase - (ALT / SGPT) (Serum)</b>			
Alanine aminotransferase - (ALT / SGPT) <i>Kinetic IFCC</i>	15	<41	U/L
<b>Protein Total, Serum (Serum)</b>			
Protein Total, Serum <i>Biuret Method</i>	7.2	6.4-8.3	g/dL

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Test Name	Result	Biological Ref. Interval	Unit
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**Senior Citizens - Female Check**

**HAEMATOLOGY**

**Complete Blood Counts (WB-EDTA)**

**(Automated Hematology Analyzer & Microscopy)**

**(Coulter Principle /Photometric method/VCS/VCSM/Cumulative pulse height/Staining/Calculated and Micr**

Total Leukocyte Count	<b>12.4 H</b>	4.0 - 11.0	10 <sup>3</sup> /μl
RBC Count	4.6	4.5 - 5.5	10 <sup>6</sup> /μL
Hemoglobin	<b>11.9 L</b>	13.0 - 17.0	g/dL
Hematocrit	<b>37.6 L</b>	40 - 50	%
MCV(Mean Corpuscular Volume)	<b>81.1 L</b>	83 - 101	fL
MCH(Mean Corpuscular Hemoglobin)	<b>25.8 L</b>	27 - 32	pg
MCHC(Mean Corpuscular Hemoglobin Concentration)	31.8	31.5 - 34.5	g/dL
RDW	<b>15.4 H</b>	11.6 - 14	%
Platelet Count	299	150 - 410	10 <sup>3</sup> /μl
MPV	8.6	7.5 - 11.5	fL
<b>Differential Counts % (VCSN)</b>			
Neutrophils	70.0	40-80%	%
Lymphocytes	24.0	20-40%	%
Monocytes	5.0	2-10%	%
Eosinophils	1.0	1-6%	%
Basophils	0.0	0-1%	%
<b>Differential Counts, Absolute</b>			
Absolute Neutrophil Count	<b>8.60 H</b>	2.0-7.0	10 <sup>3</sup> /μl
Absolute Lymphocyte Count	3.00	1.0-3.0	10 <sup>3</sup> /μl
Absolute Monocyte Count	0.70	0.2 - 1.0	10 <sup>3</sup> /μl
Absolute Eosinophil Count (AEC)	0.10	0.02-0.5	10 <sup>3</sup> /μl
Absolute Basophil Count	0.00	0.02 - 0.1	10 <sup>3</sup> /μl



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Test Name	Result	Biological Ref. Interval	Unit
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**Senior Citizens - Female Check**

**BIOCHEMISTRY**

**HbA1c - Glycated Hemoglobin (WB-EDTA)**

Glycated Hemoglobin, HbA1c <i>TINIA</i>	<b>7.30 H</b>	Non diabetic range: 4.8-5.6% Prediabetic range: 5.7-6.4% Diabetes range: >=6.5%	%
Estimated Average Glucose	162.8		mg/dL

**Interpretation:**

*Note: HbA1c results may vary in situations of abnormal red cell turnover, such as pregnancy, recent blood loss or transfusion, or some anemias. In such cases only blood glucose criteria should be used to diagnose diabetes (ADA, 2014). Please correlate clinically.*

**Glucose - Fasting (Fluoride Plasma - F)**

Glucose - Fasting <i>Hexokinase</i>	<b>149.0 H</b>	Normal : 70 - 100 Prediabetic: 100 - 125 Diabetic: >=126	mg/dL
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MC-2751

In collaboration with

**LABORATORY REPORT**

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<b>AGE</b>	: ██████████	██████████		<b>COLLECTED ON</b>	██████████
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				<b>REPORT STATUS</b>	: Final Report



Test Name	Result	Biological Ref. Interval	Unit
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**Senior Citizens - Female Check**

**CLINICAL PATHOLOGY**

**Urine Examination - Routine & Microscopy (CUE) (Urine)**

**(Dip Stick , Reflectance Photometer & Microscopy)**

**PHYSICAL EXAMINATION:**

Volume	10.00		mL
Colour	YELLOW	Pale	
<b>Appearance</b>	<b>Hazy</b>	Clear	

**CHEMICAL EXAMINATION:**

pH	5.00	4.8 - 7.4	
<i>Dip stick</i>			
Specific Gravity	<b>1.025 H</b>	1.010 - 1.022	
<i>Dip Stick(Bromothymol blue)</i>			
<b>Protein</b>	<b>POSITIVE (1+)</b>	Negative	
<i>Dip Stick/ Sulfosalicylic acid</i>			
Glucose	NEGATIVE	Negative	
<i>Dip Stick /Benedicts test</i>			
Ketones	NEGATIVE	Negative	
<i>Dip stick</i>			
Urobilinogen	NORMAL	Normal	
<i>Dip Stick / Ehrlich reaction</i>			
Nitrite	NEGATIVE	Negative	
<i>Dip Stick / (Griess test )</i>			
Bilirubin	NEGATIVE	Negative	
Blood	POSITIVE (1+)	Negative	
<i>Dip Stick ( Peroxidase)</i>			

**\*Manual**

**MICROSCOPIC EXAMINATION:**

Pus Cells	8-10 H	0 - 5	/HPF
Epithelial Cells	2-3	< 5	/HPF
Casts	Absent	Absent	/LPF
Crystals	Absent	Absent	/HPF
RBCs	6-7 H	0 - 2	/HPF

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Test Name	Result	Biological Ref. Interval	Unit
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**Senior Citizens - Female Check**

**BIOCHEMISTRY**

**Uric acid (Serum)**

Uric acid	4.2	3.4-7	mg/dL
<i>Uricase</i>			

**Blood Urea Nitrogen, BUN - Serum (Serum)**

Blood Urea Nitrogen, BUN - Serum	15.75	8.4-26	mg/dL
<i>Calculation</i>			

**Creatinine (Serum)**

Creatinine	0.56	0.7-1.4	mg/dL
<i>Modified Jaffe Kinetic</i>			

**Urea (Serum)**

Urea	33.7	18-55	mg/dL
<i>Kinetic, Urease</i>			

**Calcium - Serum (Serum)**

Calcium - Serum	9.30	8.6 - 10.0	mg/dL
<i>NM-BAPTA</i>			

**Electrolytes (Na, K, Cl) - Serum (Serum)**

Sodium	138.0	136 - 145	mmol/L
<i>ISE Indirect</i>			
Potassium - Serum	4.00	3.5-5.1	mmol/L
<i>ISE Indirect</i>			
Chlorides	98.5	98-107	mmol/L
<i>ISE Indirect</i>			

**TSH, Thyroid Stimulating Hormone (Serum)**

TSH, Thyroid Stimulating Hormone	<b>7.040 H</b>	Women (Non pregnant):0.27-4.2	µIU/mL
<i>ECLIA</i>		Pregnant women 1st trimester:0.1-2.5	
		2nd trimester: 0.2-3.0	
		3rd trimester: 0.3-3.0	

**Interpretation:**

The following potential sources of variation should be considered while interpreting thyroid hormone results:

1. Circadian variation in TSH secretion: peak levels are seen between 2-4 am. Minimum levels seen between 6-10 am. This variation may be as much as 50% thus, influence of sampling time needs to be considered for clinical interpretation.
2. Total T3 and T4 levels are seen to have physiological rise during pregnancy and in patients on steroid treatment
3. Circulating forms of T3 and T4 are mostly reversibly bound with Thyroxine binding globulins (TBG), and to a lesser extent with albumin and

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**Senior Citizens - Female Check**

Thyroid binding Pre-Albumin. Thus the conditions in which TBG and protein levels alter such as chronic liver disorders, pregnancy, excess of estrogens, androgens, anabolic steroids and glucocorticoids may cause misleading total T3, total T4 and TSH interpretations.

4. T4 may be normal in the presence of hyperthyroidism under the following conditions : T3 thyrotoxicosis, Hypoproteinemia related reduced binding, in presence of drugs (eg Phenytoin, Salicylates etc)
  5. Neonates and infants have higher levels of T4 due to increased concentration of TBG
  6. TSH levels may be normal in central hypothyroidism, recent rapid correction of hypothyroidism or hyperthyroidism, pregnancy, phenytoin therapy etc.
  7. TSH values of <0.03 uIU/mL must be clinically correlated to evaluate the presence of a rare TSH variant in certain individuals which is undetected by conventional methods.
  8. Presence of Autoimmune disorders may lead to spurious results of thyroid hormones
  9. Various drugs can lead to interference in test results
- It is recommended to evaluate unbound fractions, that is free T3 (fT3) and free T4 (fT4) for clinic-pathologic correlation, as these are the metabolically active forms.

**Vitamin D, 25-Hydroxy (Serum)**

Vitamin D, 25-Hydroxy ECLIA	<b>26.1 L</b>	Deficient: <=20 Insufficiency: 20-29 Desirable: >=30-100 Toxicity: >100	ng/ml
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**Interpretation:**

● **Interpretation:**

- Vitamin D is a fat soluble vitamin produced in the skin by exposure to sun light. Deficiency in children causes rickets and in adults leads to osteomalacia

**Decreased:**

- Impaired cutaneous production (lack of sunlight exposure)
- Dietary absence
- Malabsorption
- Increased metabolism due to drugs like barbiturates, phenytoin.
- Liver disease
- Renal failure
- VIT D receptor mutation

**Increased:**

- Vitamin D intoxication due to increased vit D supplements intake

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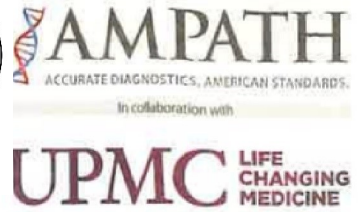
Test Name	Result	Biological Ref. Interval	Unit
<b>Senior Citizens - Female Check</b>			
<b>Serum Iron (Serum)</b>			
Iron <i>FerroZine Colorimetric Assay</i>	62.0	59-158	µg/dL
<b>C-Reactive Protein (CRP) -quantitative (Serum)</b>			
C-Reactive Protein (CRP) Quantitative <i>Immunoturbidimetry</i>	<b>11.3 H</b>	<5.0 (Negative)	mg/L
<b>Rheumatoid Factor (RA) - Quantitative - Serum (Serum)</b>			
Rheumatoid Factor (RA) - Quantitative - Serum <i>Immunoturbidimetry</i>	8.40	<14.0 (Negative)	IU/mL
<b>Lipid profile (Serum)</b>			
Cholesterol Total - Serum <i>Enzymatic colorimetric</i>	127.5	<200 No risk 200-239 Moderate risk >240 High risk	mg/dL
Triglycerides <i>Enzymatic colorimetry</i>	144.2	Normal: <150 Borderline-high: 150–199 High risk 200–499 Very high risk >500	mg/dL
Cholesterol - HDL (Direct) <i>Enzymatic colorimetric</i>	<b>37.5 L</b>	<40 High Risk ; >60 No Risk	mg/dL
VLDL (Very Low Density Lipoprotein) <i>Calculation</i>	28.8		mg/dL
LDL Chol, Calculated	61.10	<100	mg/dL

██████████  
 ██████████  
**Consultant- Biochemist**

██████████  
 ██████████  
**Consultant Pathologist & Hematopathologist**

**Disclaimer:**  
 1. All results released pertain to the specimen as received by the lab for testing and under the assumption that the patient indicated or identified on the bill/test requisition form is the owner of the specimen.  
 2. Clinical details and consent forms, especially in Genetic testing, histopathology, as well as wherever applicable, are mandatory to be accompanied with the test requisition form. The non-availability of such information may lead to delay in reporting as well as misinterpretation of test results. The lab will not be responsible for any such delays or misinterpretations thereof.

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**Senior Citizens - Female Check**

3. Test results are dependent on the quality of the sample received by the lab. In case the samples are preprocessed elsewhere (e.g., paraffin blocks), results may be compromised.

4. Tests are performed as per the schedule given in the test listing and in any unforeseen circumstances, report delivery may be affected.

5. Test results may show inter-laboratory as well as intra-laboratory variations as per the acceptable norms.

6. Genetic reports as well as reports of other tests should be correlated with clinical details and other available test reports by a qualified medical practitioner. Genetic counselling is advised in genetic test reports by a qualified genetic counsellor, medical practitioner or both.

7. Samples will be discarded post processing after a specified period as per the laboratory's retention policy. Kindly get in touch with the lab for more information.

8. If accidental damage, loss, or destruction of the specimen is not attributable to any direct or negligent act or omission on the part of Ampath Labs or its employees, Ampath shall in no event be liable. Ampath lab's liability for a lack of services, or other mistakes and omissions, shall be restricted to the amount of the patient's payment for the pertinent laboratory services.

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