

REPORT

Name	: Miss. SRIVIDYA	Sample ID	: A0934376, A0934350
Age/Gender	: 23 Years/Female	Reg. No	: 0312409160071
Referred by	: Dr. VENKATA RAMANA Y	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 16-Sep-2024 06:29 PM
Primary Sample	: Whole Blood	Received On	: 16-Sep-2024 11:25 PM
Sample Tested In	: Citrated Plasma, Capillary Tub	Reported On	: 17-Sep-2024 12:03 AM
Client Address	: Kimtee colony ,Gokul Nagar, Tarnaka	Report Status	: Final Report

HAEMATOLOGY

Test Name	Results	Units	Ref. Range	Method
Activated Partial Thromboplastin Time (APTT/PTTK)				
Patient Value	30.50	sec	26-40	Photo Optical Clot Detection
Control Value	33.00	Sec		Agglutination

Comments:APTT measures intrinsic and common pathways of the coagulation cascade. Prolonged APTT may be caused by heparin and other anticoagulants, factor deficiencies or inhibitors such as lupus anticoagulants

Bleeding Time & Clotting Time

Bleeding Time (BT)	2:30 Sec	Minutes	2 - 5	Capillary Method
Clotting Time (CT)	5:50 Sec	Minutes	3 - 7	Capillary Method

PROTHROMBIN TIME (P TIME)

PT-Patient Value	13.7	Secs	10-15	Photo Optical Clot Detection
PT-Mean Control Value	13.00	Seconds		
PT Ratio	1.05			
PT INR	1.00		0.9-1.2	

Interpretation :

Prothrombin time measures the extrinsic coagulation pathway which consists of activated Factor VII (VIIa), Tissue factor and Proteins of the common pathway (Factors X, V, II & Fibrinogen). This assay is used to control long term oral anticoagulant therapy, evaluation of liver function & to evaluate coagulation disorders specially factors involved in the extrinsic pathway like Factors V, VII, X, Prothrombin & Fibrinogen.

Note

1. INR is the parameter of choice in monitoring adequacy of oral anticoagulant therapy. Appropriate therapeutic range varies with the disease and treatment intensity
2. Prolonged INR suggests potential bleeding disorder / bleeding complications
3. Results should be clinically correlated
4. Test conducted on Citrated plasma



Swarnabala - M
DR.SWARNA BALA
MD PATHOLOGY

REPORT

Name	: Miss. SRIVIDYA	Sample ID	: A0933742
Age/Gender	: 23 Years/Female	Reg. No	: 0312409160071
Referred by	: Dr. VENKATA RAMANA Y	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 16-Sep-2024 06:29 PM
Primary Sample	: Whole Blood	Received On	: 16-Sep-2024 11:25 PM
Sample Tested In	: Whole Blood EDTA	Reported On	: 17-Sep-2024 12:36 AM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

HAEMATOLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
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Blood Grouping (A B O)

B

Tube Agglutination

Rh Typing

Positive

Tube Agglutination

*** End Of Report ***

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Swannabala - M
DR.SWARNA BALA
MD PATHOLOGY

REPORT

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Referred by	: Dr. VENKATA RAMANA Y	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 16-Sep-2024 06:29 PM
Primary Sample	: Whole Blood	Received On	: 16-Sep-2024 11:25 PM
Sample Tested In	: Whole Blood EDTA	Reported On	: 16-Sep-2024 11:55 PM
Client Address	: Kimtee colony ,Gokul Nagar ,Tarnaka	Report Status	: Final Report

HAEMATOLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
Complete Blood Picture(CBP)				
Haemoglobin (Hb)	11.9	g/dL	12-15	Cynmeth Method
Haematocrit (HCT)	40.0	%	40-50	Calculated
RBC Count	4.32	10 ¹² /L	3.8-4.8	Cell Impedance
MCV	91	fl	81-101	Calculated
MCH	27.4	pg	27-32	Calculated
MCHC	33.6	g/dL	32.5-34.5	Calculated
RDW-CV	12.8	%	11.6-14.0	Calculated
Platelet Count (PLT)	353	10 ⁹ /L	150-410	Cell Impedance
Total WBC Count	8.8	10 ⁹ /L	4.0-10.0	Impedance
Differential Leucocyte Count (DC)				
Neutrophils	70	%	40-70	Cell Impedance
Lymphocytes	25	%	20-40	Cell Impedance
Monocytes	03	%	2-10	Microscopy
Eosinophils	02	%	1-6	Microscopy
Basophils	00	%	1-2	Microscopy
Absolute Neutrophils Count	6.16	10 ⁹ /L	2.0-7.0	Impedance
Absolute Lymphocyte Count	2.2	10 ⁹ /L	1.0-3.0	Impedance
Absolute Monocyte Count	0.26	10 ⁹ /L	0.2-1.0	Calculated
Absolute Eosinophils Count	0.18	10 ⁹ /L	0.02-0.5	Calculated
Absolute Basophil ICount	0.00	10 ⁹ /L	0.0-0.3	Calculated
Morphology	Normocytic Normochromic			PAPs Staining



Swannabala - M
DR.SWARNA BALA
MD PATHOLOGY

REPORT

Name	: Miss. SRIVIDYA	Sample ID	: A0934375, A0934373
Age/Gender	: 23 Years/Female	Reg. No	: 0312409160071
Referred by	: Dr. VENKATA RAMANA Y	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 16-Sep-2024 06:29 PM
Primary Sample	: Whole Blood	Received On	: 16-Sep-2024 11:36 PM
Sample Tested In	: Plasma-NaF(R), Serum	Reported On	: 17-Sep-2024 12:38 AM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

CLINICAL BIOCHEMISTRY

Test Name	Results	Units	Ref. Range	Method
25 - Hydroxy Vitamin D	42.71	ng/mL	<20.0-Deficiency 20.0-30.0-Insufficiency 30.0-100.0-Sufficiency >100.0-Potential Intoxication	CLIA

Interpretation:

- Vitamin D helps your body absorb calcium and maintain strong bones throughout your entire life. Your body produces vitamin D when the sun's UV rays contact your skin. Other good sources of the vitamin include fish, eggs, and fortified dairy products. It's also available as a dietary supplement.
- Vitamin D must go through several processes in your body before your body can use it. The first transformation occurs in the liver. Here, your body converts vitamin D to a chemical known as 25-hydroxyvitamin D, also called calcidiol.
- The 25-hydroxy vitamin D test is the best way to monitor vitamin D levels. The amount of 25-hydroxyvitamin D in your blood is a good indication of how much vitamin D your body has. The test can determine if your vitamin D levels are too high or too low.
- The test is also known as the 25-OH vitamin D test and the calcidiol 25-hydroxycholecalciferol test. It can be an important indicator of osteoporosis (bone weakness) and rickets (bone malformation).

Those who are at high risk of having low levels of vitamin D include:

- people who don't get much exposure to the sun
- older adults
- people with obesity.
- dietary deficiency

Increased Levels: Vitamin D Intoxication

Method : CLIA

Vitamin- B12 (cyanocobalamin)	339	pg/mL	200-911	CLIA
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Interpretation:

This test is most often done when other blood tests suggest a condition called megaloblastic anemia. Pernicious anemia is a form of megaloblastic anemia caused by poor vitamin B12 absorption. This can occur when the stomach makes less of the substance the body needs to properly absorb vitamin B12.

Causes of vitamin B12 deficiency include:Diseases that cause malabsorption

- Lack of intrinsic factor, a protein that helps the intestine absorb vitamin B12
- Above normal heat production (for example, with hyperthyroidism)

An increased vitamin B12 level is uncommon in:

- Liver disease (such as cirrhosis or hepatitis)
- Myeloproliferative disorders (for example, polycythemia vera and chronic myelogenous leukemia)
- .



Dr. Vaishnavi
DR. VAISHNAVI
MD BIOCHEMISTRY

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Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 16-Sep-2024 06:29 PM
Primary Sample	: Whole Blood	Received On	: 16-Sep-2024 11:36 PM
Sample Tested In	: Serum	Reported On	: 17-Sep-2024 12:46 AM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

IMMUNOLOGY & SEROLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
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VDRL- Syphilis Antibodies	Non Reactive		Non Reactive	Slide Flocculation
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The serological diagnosis of syphilis is classified into two groups: Nontreponemal tests (RPR/VDRL) and Treponemal tests (TPHA/CLIA). Syphilis serology is a treponemal assay for the qualitative determination of antibodies to T. pallidum in human serum or plasma as an aid in the diagnosis of syphilis. Treponemal tests may remain reactive for life, even following adequate therapy thus a positive result suggests infection with Treponema pallidum but does not distinguish between treated and untreated infections. Therefore, the results of a nontreponemal assay, such as rapid plasma reagin, are needed to provide information on a patient's disease state and history of therapy. Nontreponemal tests lack sensitivity in late stage of infection and screening with these tests alone may yield false positive reactions in various acute and chronic conditions in the absence of syphilis (biological false positive reactions).

Result rechecked and verified for abnormal cases

*** End Of Report ***

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DR. RUTURAJ MANIKLAL KOLHAPURE
MD, MICROBIOLOGIST

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IMMUNOLOGY & SEROLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
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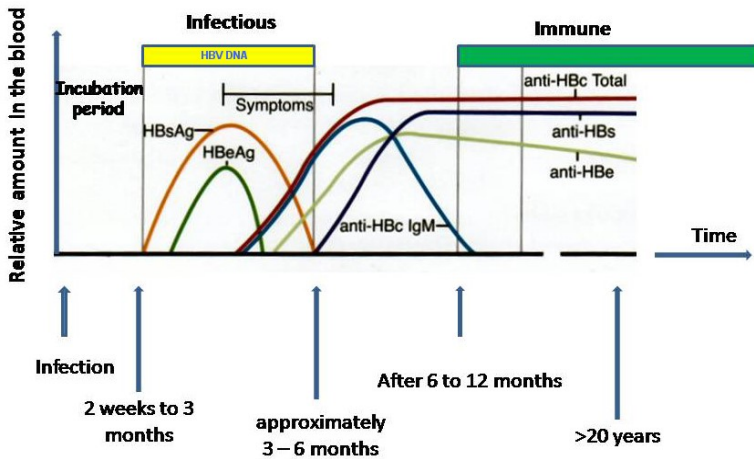
Hepatitis B Surface Antigen (HBsAg)	0.30	S/Co	<1.00 :Negative >1.00 :Positive	ELISA
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Interpretation:

- Negative result implies that antibodies to HBsAg have not been detected in the sample. This means the patient has either not been exposed to HBsAg infection or the sample has been tested during the "window phase" i.e. before the development of detectable levels of antibodies. Hence a Non-Reactive result does not exclude the possibility of exposure or infection with HBsAg.
- Positive result implies that antibodies to HBsAg have been detected in the sample.

Hepatitis B Virus (HBV) is a member of the Hepadna virus family causing infections of the liver with extremely variable clinical features. Hepatitis B is transmitted primarily by body fluids especially serum and also spread effectively sexually and from mother to baby. In most individuals HBV hepatitis is self limiting, but 1-2% normal adolescents and adults develop Chronic Hepatitis. Frequency of chronic HBV infection is 5-10% in immunocompromised patients and 80% in neonates. The initial serological marker of acute infection is HBsAg which typically appears 2-3 months after infection and disappears 12-20 weeks after onset of symptoms. Persistence of HBsAg for more than six months indicates development of carrier state or Chronic liver disease.

HBV antigens and antibodies in the blood



Note:

1. All Reactive results are tested additionally by Specific antibody Neutralization assay . For further confirmation Molecular assays are recommended For diagnostic purposes, results should be used in conjunction with clinical history and other hepatitis markers for Acute or Chronic infection

*** End Of Report ***

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IMMUNOLOGY & SEROLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
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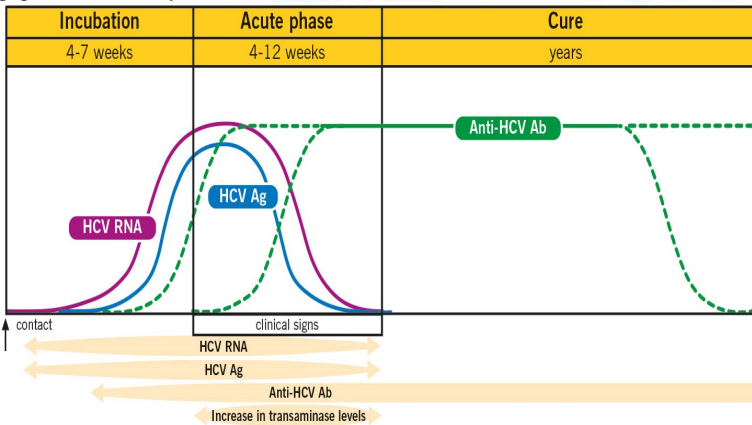
Hepatitis C Virus Antibody	0.21	S/Co	< 1.00 : Negative > 1.00 : Positive	ELISA
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Interpretation:

1. Negative result implies that antibodies to HCV have not been detected in the sample. This means the patient has either not been exposed to HCV infection or the sample has been tested during the "window phase" i.e. before the development of detectable levels of antibodies. Hence a Non-Reactive result does not exclude the possibility of exposure or infection with HCV.
2. Positive result implies that antibodies to HCV have been detected in the sample.

Comments :-

Hepatitis C (HCV) is an RNA virus of Flavivirus group transmitted via blood transfusions, transplantation, injection drug users, accidental needle punctures in healthcare workers, dialysis patients and rarely from mother to infant. 10% of new cases show sexual transmission. As compared to HAV & HBV, chronic infection with HCV occurs in 85% of infected individuals. In high risk populations, the predictive value of Anti HCV for HCV infection is > 99% whereas in low risk populations it is only 25%.



Note:

1. False positive results are seen in Autoimmune diseases, Rheumatoid factor, Hypergammaglobulinemia, Paraproteinemia, passive antibody transfer, Anti- idiotypes & Anti superoxide dismutase
2. False negative results are seen in early Acute infection, Immunosuppression & Immuno-incompetence
3. HCV RNA PCR recommended in all Reactive results to differentiate between past and present infection

*** End Of Report ***

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IMMUNOLOGY & SEROLOGY

SURGICAL PROFILE-II

Test Name	Results	Units	Ref. Range	Method
HIV (1& 2) Antibody	0.43	S/Co	< 1.00 : Negative > 1.00 : Positive	ELISA

Correlate Clinically.

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*** End Of Report ***



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