

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013463
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	:	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Capillary Tube	Reported On	: 13-Feb-2024 05:28 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

HAEMATOLOGY

ANTE NATEL PROFILE-ELISA

Test Name	Results	Units	Ref. Range	Method
Bleeding Time & Clotting Time				
Bleeding Time (BT)	03:30	Minutes	2 - 5	Capillary Method
Clotting Time (CT)	05:40	Minutes	3 - 7	Capillary Method



Swannabala - M
DR.SWARNA BALA
MD PATHOLOGY

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013461
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:05 PM
Sample Tested In	: Whole Blood EDTA	Reported On	: 13-Feb-2024 04:39 PM
Client Address	: Kimtee colony ,Gokul Nagar, Tarnaka	Report Status	: Final Report

HAEMATOLOGY

ANTE NATEL PROFILE-ELISA

Test Name	Results	Units	Ref. Range	Method
Blood Grouping (A B O)	O			Tube Agglutination
Rh Typing	Positive			Tube Agglutination
Complete Blood Count (CBC)				
Haemoglobin (Hb)	12.3	g/dL	12-15	Cynmeth Method
RBC Count	4.62	10 ¹² /L	4.5-5.5	Cell Impedence
Total WBC Count	5.3	10 ⁹ /L	4.0-10.0	Impedance
Platelet Count (PLT)	248	10 ⁹ /L	150-410	Cell Impedance
Haematocrit (HCT)	38.8	%	40-50	Calculated
MCV	84	fl	81-101	Calculated
MCH	26.6	pg	27-32	Calculated
MCHC	31.6	g/dL	32.5-34.5	Calculated
RDW-CV	15.2	%	11.6-14.0	Calculated
Differential Count by Flowcytometry /Microscopy				
Neutrophils	68	%	40-70	Cell Impedence
Lymphocytes	26	%	20-40	Cell Impedence
Monocytes	03	%	2-10	Microscopy
Eosinophils	03	%	1-6	Microscopy
Basophils	0	%	1-2	Microscopy
Smear				
WBC	Within normal limits.			
RBC	Normocytic normochromic blood picture			
Platelets	Adequate			Microscopy



Swannabala - M
DR. SWARNA BALA
MD PATHOLOGY

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013460
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Serum	Reported On	: 13-Feb-2024 06:37 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

CLINICAL BIOCHEMISTRY

Test Name	Results	Units	Ref. Range	Method
TSH -Thyroid Stimulating Hormone	2.13	µIU/mL	0.35-5.5	CLIA

Pregnancy & Cord Blood

TSH (Thyroid Stimulating Hormone (µIU/mL))	
First Trimester	: 0.24-2.99
Second Trimester	: 0.46-2.95
Third Trimester	: 0.43-2.78
Cord Blood	: 2.3-13.2

- TSH is synthesized and secreted by the anterior pituitary in response to a negative feedback mechanism involving concentrations of FT3 (free T3) and FT4 (free T4). Additionally, the hypothalamic tripeptide, thyrotropin-releasing hormone (TRH), directly stimulates TSH production.
- TSH interacts with specific cell receptors on the thyroid cell surface and exerts two main actions. The first action is to stimulate cell reproduction and hypertrophy. Secondly, TSH stimulates the thyroid gland to synthesize and secrete T3 and T4
- The ability to quantitate circulating levels of TSH is important in evaluating thyroid function. It is especially useful in the differential diagnosis of primary (thyroid) from secondary (pituitary) and tertiary (hypothalamus) hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low
- TRH stimulation differentiates secondary and tertiary hypothyroidism by observing the change in patient TSH levels. Typically, the TSH response to TRH stimulation is absent in cases of secondary hypothyroidism, and normal to exaggerated in tertiary hypothyroidism
- Historically, TRH stimulation has been used to confirm primary hyperthyroidism, indicated by elevated T3 and T4 levels and low or undetectable TSH levels. TSH assays with increased sensitivity and specificity provide a primary diagnostic tool to differentiate hyperthyroid from euthyroid patients.



Dr. Vaishnavi
DR. VAISHNAVI
MD BIOCHEMISTRY

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013460
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Serum	Reported On	: 13-Feb-2024 07:37 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

IMMUNOLOGY & SEROLOGY

ANTE NATEL PROFILE-ELISA

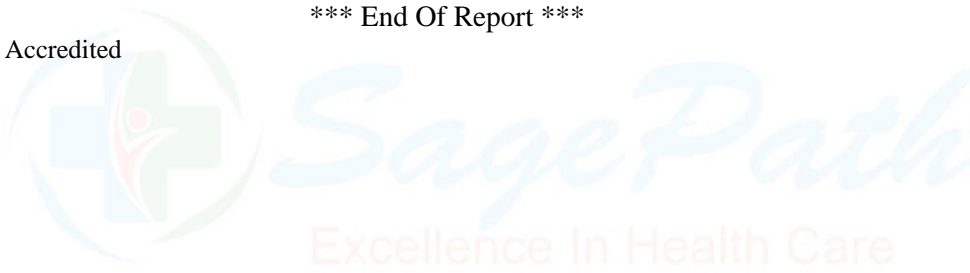
Test Name	Results	Units	Ref. Range	Method
-----------	---------	-------	------------	--------

VDRL- Syphilis Antibodies	Non Reactive		Non Reactive	Slide Flocculation
----------------------------------	--------------	--	--------------	--------------------

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).

*** End Of Report ***

Laboratory is NABL Accredited




DR. RUTURAJ MANIKLAL KOLHAPURE
MD, MICROBIOLOGIST

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013460
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Serum	Reported On	: 13-Feb-2024 07:12 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

IMMUNOLOGY & SEROLOGY

ANTE NATEL PROFILE-ELISA

Test Name	Results	Units	Ref. Range	Method
-----------	---------	-------	------------	--------

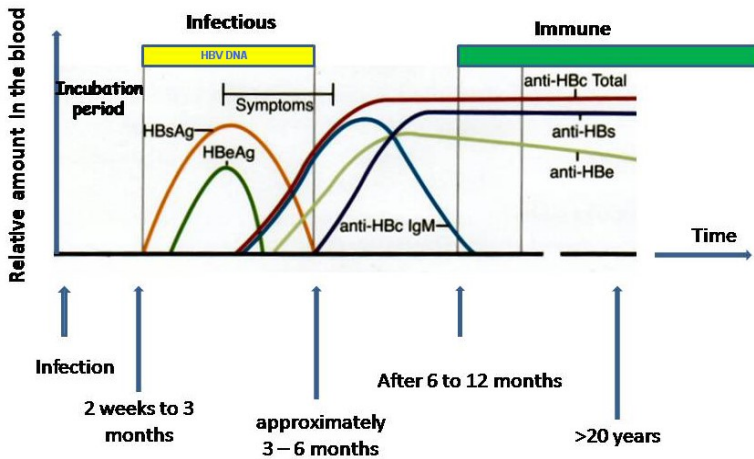
Hepatitis B Surface Antigen (HBsAg)	0.33	S/Co	<1.00 :Negative >1.00 :Positive	ELISA
--	------	------	------------------------------------	-------

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 ***

Laboratory is NABL Accredited



DR. RUTURAJ MANIKLAL KOLHAPURE
MD, MICROBIOLOGIST

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013460
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Serum	Reported On	: 13-Feb-2024 07:37 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

IMMUNOLOGY & SEROLOGY

ANTE NATEL PROFILE-ELISA

Test Name	Results	Units	Ref. Range	Method
-----------	---------	-------	------------	--------

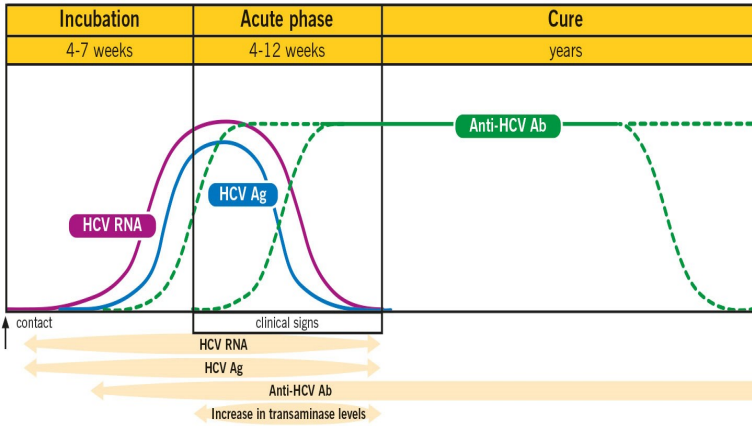
Hepatitis C Virus Antibody	0.25	S/Co	< 1.00 : Negative > 1.00 : Positive	ELISA
-----------------------------------	------	------	--	-------

Interpretation:

- 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.
- 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:

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

*** End Of Report ***

Laboratory is NABL Accredited



DR. RUTURAJ MANIKLAL KOLHAPURE
MD, MICROBIOLOGIST

REPORT

Name	: Mrs. RAJITHA	Sample ID	: A0013460
Age/Gender	: 27 Years/Female	Reg. No	: 0312402130028
Referred by	: Dr. Nivedita Ashrit MD (Obs/Gyn)	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 13-Feb-2024 01:44 PM
Primary Sample	: Whole Blood	Received On	: 13-Feb-2024 04:08 PM
Sample Tested In	: Serum	Reported On	: 13-Feb-2024 07:31 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

IMMUNOLOGY & SEROLOGY

ANTE NATEL PROFILE-ELISA

Test Name	Results	Units	Ref. Range	Method
HIV (1& 2) Antibody	0.21	S/Co	< 1.00 : Negative > 1.00 : Positive	ELISA

Correlate Clinically.

Laboratory is NABL Accredited

*** End Of Report ***



DR. RUTURAJ MANIKLAL KOLHAPURE
MD, MICROBIOLOGIST