





Name : MRS. MAHESHWARI Bill Number

Age/Gender : 24YEARS/FEMALE Bill Date : 01-Sep-2024 10:34 AM

Sample Type : WB EDTA Sample Collection : 01-Sep-2024 11:19 AM Reff By : DR.L.RAJ KUMAR Sample Received : 01-Sep-2024 11:20 AM

TypedBy : Tanveer Fathima : 01-Sep-2024 02:33 PM Reporting Date

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	<u>UNITS</u>	NORMAL RANGE
HAEMOGLOBIN (Method: Cell Counter)	11.7	gm/dL	12.0 - 15.0
RBC Count (Method: Cell Counter)	3.9	Millions/Cumm	3.8 - 4.8
WBC Count (Method: Cell Counter)	7,800	Cells/cumm	4,000 - 11,000
RDW - CV (Method: Cell Counter)	12.8	%	11.0 - 16.0
DIFFERENTIAL COUNT			
NEUTROPHILS (Method: Cell Counter)	74	%	40 - 75
LYMPHOCYTES (Method: Cell Counter)	18	%	20 - 40
EOSINOPHILS (Method: Cell Counter)	01	%	01 - 06
MONOCYTES (Method: Cell Counter)	07	%	02 - 10
BASOPHILS (Method: Cell Counter)	00	%	0 - 0
PCV (Haematocrit) (Method: Cell Counter)	35	%	35 - 45
MCV (Method: Cell Counter)	89	FL	83 - 101
MCH (Method: Cell Counter)	30	pg	27 - 32
MCHC (Method: Cell Counter)	34	%	32 - 35
PLATELET COUNT (Method: Cell Counter)	2.11	Lakhs/Cumm	1.5 - 4.5
PERIPHERAL SMEAR			
BBCs	NORMOCYTIC NORMOCHROMIC		

RBCs NORMOCYTIC NORMOCHROMIC

WBCs WITHIN NORMAL LIMITS

PLATELETS ADEQUATE

Sugessted Clinical Correlation If necesarry Kindly Discuss.

-----End of the Report-----







Name : MRS. MAHESHWARI Bill Number : M610

Age/Gender : 24YEARS/FEMALE Bill Date : 01-Sep-2024 10:34 AM

Sample Type : WB EDTA Sample Collection : 01-Sep-2024 11:19 AM

Reff By : **DR.L.RAJ KUMAR** Sample Received : 01-Sep-2024 11:20 AM TypedBy : Tanveer Fathima Reporting Date : 01-Sep-2024 02:33 PM

は対象 の対象を含

LAB INCHARGE

Authorized Signatory







Name : MRS. MAHESHWARI Bill Number : M6162

Age/Gender : 24YEARS/FEMALE Bill Date : 01-Sep-2024 10:34 AM

Sample Type : URINE Sample Collection : 01-Sep-2024 11:19 AM

Reff By : **DR.L.RAJ KUMAR** Sample Received : 01-Sep-2024 11:20 AM

TypedBy : Tanveer Fathima Reporting Date : 01-Sep-2024 02:34 PM

CUE(COMPLETE URINE EXAMINATION)

INVESTIGATION RESULT NORMAL RANGE

PHYSICAL EXAMINATION

QUANTITY 20 ml

COLOUR YELLOW

APPEARANCE SLIGHTLY TURBID

REACTION (PH) 7.0 4.6 - 8.0

SPECIFIC GRAVITY 1.025 1.005 - 1.030

CHEMICAL EXAMINATION

ALBUMIN PRESENT (+) NEGATIVE

SUGAR NIL NIL

UROBILINOGEN NEGATIVE NEGATIVE

BILE SALT NEGATIVE NEGATIVE

BILE PIGMENT NEGATIVE NEGATIVE NEGATIVE

KETONE BODIES NEGATIVE NEGATIVE NEGATIVE

MICROSCOPIC EXAMINATION

PUS CELLS 3 - 4 / HPF 0 - 5 / HPF

RBC NIL NIL

EPETHILIAL CELLS 3 - 4 / HPF 0 - 5 / HPF

CRYSTALS NIL NIL

CASTS NIL NIL

OTHERS NIL NIL

Method: Multi Reagent Strip / Chemical / Microscopy

Sugessted Clinical Correlation If necesarry Kindly Discuss.







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RAPID MALARIA TEST (PV PF)

INVESTIGATION **NORMAL RANGE RESULT**

MALARIAL PARASITE PLASMODIUM **NEGATIVE NEGATIVE**

VIVAX(P.V)

(Method: Immunochromotography)

NEGATIVE NEGATIVE MALARIAL PARASITE PLASMODIUM

FALCIPARUM(PF) (Method: Immunochromotography)

Method: Immunochromotography

Note:

This test detects P.falciparum and P.Vivax Malarial antigens targeted by highly specific monoclonal antibodies of pLDH and Aldolase respectively.

Sensitivity and Specificity of this test are 98.2% &99.6% for P.falciparum and 91.8% &99.6% for P.Vivax detection.

This kit can provide fast and easy way to get a result, but do not completely exclude the possibility of false positive or false negative result caused by various factors.

Sugessted Clinical Correlation If necesarry Kindly Discuss.

-----End of the Report-----

Authorized Signatory



: DR.L.RAJ KUMAR



: 01-Sep-2024 11:20 AM

Name : MRS. MAHESHWARI Bill Number : M6162

Age/Gender : 24YEARS/FEMALE Bill Date : 01-Sep-2024 10:34 AM

Sample Type : **SERUM** Sample Collection : 01-Sep-2024 11:19 AM

TypedBy : Tanveer Fathima Reporting Date : 01-Sep-2024 02:36 PM

WIDAL

Sample Received

INVESTIGATION RESULT

SALMONELLA TYPHI' O ' 1 in 160 DILUTION

SALMONELLA TYPI 'H' 1 in 80 DILUTION

SALMONELLA PARA TYPHI' AH' 1 in 20 DILUTION

SALMONELLA PARA TYPHI' BH' 1 in 20 DILUTION

BIOLOGICAL REFERENCE 1:80 and above titers considered as positive

Method: SEMI QUANTITAVE SLIDE AGGLUTINATION

Interpretation and Remarks:

Reff By

- The Widal test is applied for the diagnosis of enteric fever that includes typhoid and paratyphoid caused by Salmonella? typhi and Salmonella paratyphi respectively.
- For the slide agglutination test, stained Salmonella antigens are used to detect the presence of specific agglutinin in the patient's serum.
- The slide agglutination test is used as a primary screening procedure.
- Widal test measures somatic O and flagellar H antibodies against Typhoid and Paratyphoid bacilli. The Agglutinins appears usually at the end of the first week of infection and increases steadily till third / fourth week after which the decline starts. A positive Widal test may occur because of typhoid vaccination or past typhoid infection and in certain autoimmune diseases. Nonspecific febrile disease may cause titre to increase. The test may be falsely negative in cases of Enteric fever treated with antibiotics in the early stages

Sugessted Clinical Correlation If necesarry Kindly Discuss.

-----End of the Report-----

Authorized Signatory



: DR.L.RAJ KUMAR





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Sample Type : **SERUM** Sample Collection : 01-Sep-2024 11:19 AM

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Dengue NS1 antigen IgG IgM

Sample Received

INVESTIGATION RESULT NORMAL RANGE

DENGUE NS1 NEGATIVE NEGATIVE

(Method: Rapid)

Reff By

DENGUE - IGG NEGATIVE NEGATIVE

(Method: Rapid)

DENGUE - IGM NEGATIVE NEGATIVE

(Method: Rapid)

TECHNOLOGY RAPID VISUAL TEST FOR THE DETECTIONOF DENGUE

Interpretation:

The NS1 antigen is typically detectable within 1 to 2 days following infection and up to 9 days following symptom onset.NS1 antigen may also be detectable during secondary dengue virus infection, but for a shorter duration of time (1-4 days following symptom onset).

False-positive Dengue NS results may occur in individuals with active infection due to other flaviviruses, including West Nile virus and yellow fever virus.

Negative NS1 antigen results may occur if the specimen was collected greater than 7 days following symptom onset

Sugessted Clinical Correlation If necesarry Kindly Discuss.

-----End of the Report-----

Authorized Signatory