



Name : **MRS. MOUNIKA**
Age/Gender : **28YEARS/FEMALE**
Sample Type : **SERUM**
Reff By : **DR.SELF**
TypedBy : **Md Masud Ansari**

Bill Number : **M3396**
Bill Date : **15-Jul-2024 10:06 AM**
Sample Collection : **15-Jul-2024 10:11 AM**
Sample Received : **15-Jul-2024 10:16 AM**
Reporting Date : **15-Jul-2024 02:34 PM**

LIVER FUNCTION TEST (LFT)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
TOTAL BILIRUBIN (Method: Jendrassik and Grof)	0.9	mg/dl	0.4 - 1.2
DIRECT BILIRUBIN (Method: Modified Jendrassik)	0.2	mg/dl	Up to 0.25
INDIRECT BILIRUBIN (Method: Calculated)	0.7	mg/dl	up to 1
SGPT(ALT) (Method: Kinetic: IFCC)	19	U/L	Male :Upto 40 Female :Upto 31
SGOT(AST) (Method: Kinetic IFCC)	27	U/L	Male: Upto 37 Female: Upto 31
ALKALINE PHOSPHATASE(ALP) (Method: PNPP AMP Buffer)	88	U/L	Adults : 30-120 Children: 47 - 406
TOTAL PROTEINS (Method: Biuret)	7.8	gm/dl	6.4 - 8.3
ALBUMIN (Method: BCG)	4.2	gm/dl	3.8 - 4.4 gm/dL
GLOBULIN (Method: Calculated)	3.6	gm/dl	2.6 - 3.9
A/G Ratio (Method: Calculated)	1.2		1.2-2.2

Total Bilirubin reference range in case of Premature neonates is :0 - 1day: 1.0 - 8.0, 1 - 2day: 6.0 - 12.0, 3 - 5day: 10.0 - 14.0

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

Authorized Signatory



LAB INCHARGE



Name : **MRS. MOUNIKA**
Age/Gender : **28YEARS/FEMALE**
Sample Type : **BLOOD**
Reff By : **DR.SELF**
TypedBy : Md Masud Ansari

Bill Number : **M3396**
Bill Date : 15-Jul-2024 10:06 AM
Sample Collection : 15-Jul-2024 10:11 AM
Sample Received : 15-Jul-2024 10:16 AM
Reporting Date : 17-Jul-2024 05:45 PM

ANAEROBIC BLOOD CULTURE

INVESTIGATION

SPECIMEN

RESULTS

RESULT

Blood

No bacterial growth seen after 36 to 48 hrs of aerobic incubation at 37 °C

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

Authorized Signatory



LAB INCHARGE



Name : **MRS. MOUNIKA**
Age/Gender : **28YEARS/FEMALE**
Sample Type : **URINE**
Reff By : **DR.SELF**
TypedBy : **Md Masud Ansari**

Bill Number : **M3396**
Bill Date : **15-Jul-2024 10:06 AM**
Sample Collection : **15-Jul-2024 10:11 AM**
Sample Received : **15-Jul-2024 10:16 AM**
Reporting Date : **17-Jul-2024 05:40 PM**

CULTURE & SENSITIVITY OF URINE

INVESTIGATION

SPECIMEN

Results

RESULT

Urine

No bacterial growth seen after 36 to 48 hrs of aerobic incubation at 37 °C

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

Authorized Signatory



LAB INCHARGE



Name : **MRS. MOUNIKA**
Age/Gender : **28YEARS/FEMALE**
Sample Type : **WB EDTA**
Reff By : **DR.SELF**
TypedBy : **Md Masud Ansari**

Bill Number : **M3396**
Bill Date : **15-Jul-2024 10:06 AM**
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Sample Received : **15-Jul-2024 10:16 AM**
Reporting Date : **15-Jul-2024 02:37 PM**

COMPLETE BLOOD PICTURE (CBP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
HAEMOGLOBIN (Method: Cell Counter)	11.2	gm/dL	12.0 - 15.0
RBC Count (Method: Cell Counter)	4.1	Millions/Cumm	3.8 - 4.8
WBC Count (Method: Cell Counter)	19,800	Cells/cumm	4,000 - 11,000
RDW (Method: Cell Counter)	13.9	%	11.0 - 16.0
DIFFERENTIAL COUNT			
NEUTROPHILS (Method: Cell Counter)	78	%	40 - 75
LYMPHOCYTES (Method: Cell Counter)	18	%	20 - 40
EOSINOPHILS (Method: Cell Counter)	01	%	01 - 06
MONOCYTES (Method: Cell Counter)	03	%	02 - 10
BASOPHILS (Method: Cell Counter)	00	%	0 - 0
PCV (Haematocrit) (Method: Cell Counter)	33	%	35 - 45
MCV (Method: Cell Counter)	80	FL	83 - 101
MCH (Method: Cell Counter)	27	pg	27 - 32
MCHC (Method: Cell Counter)	33	%	32 - 35
PLATELET COUNT (Method: Cell Counter)	2.20	Lakhs/Cumm	1.5 - 4.5
PERIPHERAL SMEAR			
RBCs	NORMOCYTIC MILD HYPOCHROMIC		
WBCs	LEUCOCYTOSIS WITH NEUTROPHILIA		
PLATELETS	ADEQUATE		



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Sugessted Clinical Correlation If necesarry Kindly Discuss.

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



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RAPID MALARIA TEST (PV PF)

INVESTIGATION	RESULT	NORMAL RANGE
MALARIAL PARASITE PLASMODIUM VIVAX(P.V) (Method: Immunochromotography)	NEGATIVE	NEGATIVE
MALARIAL PARASITE PLASMODIUM FALCIPARUM(PF) (Method: Immunochromotography)	NEGATIVE	NEGATIVE
Method: Immunochromotography		

Note :

1. This test detects P.falciparum and P.Vivax Malarial antigens targeted by highly specific monoclonal antibodies of pLDH and Aldolase respectively.
2. Sensitivity and Specificity of this test are 98.2% & 99.6% for P.falciparum and 91.8% & 99.6% for P.Vivax detection.
3. 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.

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WIDAL

INVESTIGATION

SALMONELLA TYPHI' O '

SALMONELLA TYPI ' H '

SALMONELLA PARA TYPHI' AH'

SALMONELLA PARA TYPHI' BH'

BIOLOGICAL REFERENCE

RESULT

1 in 160 DILUTION

1 in 80 DILUTION

1 in 20 DILUTION

1 in 20 DILUTION

1:80 and above titers considered as positive

Method: SEMI QUANTITAVE SLIDE AGGLUTINATION

Interpretation and Remarks:

- 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-----

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C - REACTIVE PROTEINS (CRP)

INVESTIGATION	RESULT	UNITS	NORMAL RANGE
C – REACTIVE PROTEINS (Method: Immunoturbidimetry)	24.76	mg/L	0.0 - 6.0
INTERPRETAION	POSITIVE		

Note :

- 1.The CRP test is a sensitive indicator of inflammatory processes.
- 2.The determination of the CRP level can be used in therapy control
- 3.As with all the diagnostic methods, the final diagnosis should not be based on the result of a single test, but on a correlation of test results with other clinical findings.
- 4.The strength of agglutination is not indicative of the CRP concentration in the samples tested.

Please Correlate With Clinical Findings If Necessary Discuss

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

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 TypedBy : Md Masud Ansari

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CUE(COMPLETE URINE EXAMINATION)

INVESTIGATION	RESULT	NORMAL RANGE
PHYSICAL EXAMINATION		
QUANTITY	20 ml	
COLOUR	PALE YELLOW	
APPEARANCE	SLIGHTLY TURBID	
REACTION (PH)	6.0	4.6 - 8.0
SPECIFIC GRAVITY	1.025	1.005 - 1.030
CHEMICAL EXAMINATION		
ALBUMIN	TRACE	NEGATIVE
SUGAR	NIL	NIL
UROBILINOGEN	NEGATIVE	NEGATIVE
BILE SALT	NEGATIVE	NEGATIVE
BILE PIGMENT	NEGATIVE	NEGATIVE
KETONE BODIES	NEGATIVE	NEGATIVE
MICROSCOPIC EXAMINATION		
PUS CELLS	6 - 8 / HPF	0 - 5 / HPF
RBC	NIL	NIL
EPETHILIAL CELLS	2 - 3 / HPF	0 - 5 / HPF
CRYSTALS	NIL	NIL
CASTS	NIL	NIL



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OTHERS

NIL

NIL

Method: Multi Reagent Strip / Chemical / Microscopy

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



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