

Client**Pathkind Labs (Hansrajpur Varanasi)**

House No. 113, Tehsil Jakhaniya

Nasirpur, Hansrajpur, Ghazipur, UP

Processed By**Pathkind Diagnostic Pvt. Ltd.**

S.No 40, Badi Bagh Lanka, Ghazipur-233001

Name	: Mrs. Dummy	Billing Date	: 01/01/2020 11:16:22
Age	: 42 Yrs	Sample Collected on	: 01/01/2020 11:27:45
Sex	: Female	Sample Received on	: 01/01/2020 13:15:35
P. ID No.	: P1201123210	Report Released on	: 01/01/2020 13:39:41
Accession No	: 12011910051010	Barcode No.	: 4548545
Referring Doctor	: Dr.Pradeep maurya		
Referred By	:		

Report Status - Final

Test Name	Result	Biological Ref. Interval	Unit
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HAEMATOLOGY**Fever Panel Basic****Complete Blood Count (CBC)****Haemoglobin (Hb)**

Sample: Whole Blood EDTA

Method: Photometric measurement

13.9

12.0 - 15.0

gm/dL

Total WBC Count

Sample: Whole Blood EDTA

Method: Impedance

14.7 H

4.0 - 10.0

thou/ μ L**RBC Count**

Sample: Whole Blood EDTA

Method: Impedance

4.4

3.8 - 4.8

million/ μ L**PCV / Hematocrit**

Sample: Whole Blood EDTA

Method: Impedance

39.6

36.0 - 46.0

%

MCV

Sample: Whole Blood EDTA

Method: Calculated

89.0

83.0 - 101.0

fL

MCH

Sample: Whole Blood EDTA

Method: Calculated

31.3

27.0 - 32.0

pg

MCHC

Sample: Whole Blood EDTA

Method: Calculated

35.0 H

31.5 - 34.5

g/dL

RDW (Red Cell Distribution Width)

Sample: Whole Blood EDTA

Method: Calculated

13.7

11.9 - 15.5

%

DLC (Differential Leucocyte Count)

Method: Flowcytometry/Microscopy

Neutrophils

Sample: Whole Blood EDTA

Method: VCS Technology & Microscopy

82 H

40 - 80

%

12011910051010 Mrs. Dummy



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Lymphocytes <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology & Microscopy</i>	14 L	20 - 40	%
Eosinophils <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology & Microscopy</i>	02	01 - 06	%
Monocytes <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology & Microscopy</i>	02	02 - 10	%
Basophils <i>Sample: Whole Blood EDTA</i> <i>Method: VCS Technology & Microscopy</i>	00	00 - 02	%
Absolute Neutrophil Count <i>Sample: Whole Blood EDTA</i>	12054 H	2000 - 7000	/μL
Absolute Lymphocyte Count <i>Sample: Whole Blood EDTA</i>	2058	1000 - 3000	/μL
Absolute Eosinophil Count <i>Sample: Whole Blood EDTA</i>	294	20 - 500	/μL
Absolute Monocyte Count <i>Sample: Whole Blood EDTA</i>	294	200 - 1000	/μL
Absolute Basophil Count <i>Sample: Whole Blood EDTA</i>	00 L	20 - 100	/μL
DLC Performed By <i>Sample: Whole Blood EDTA</i>	EDTA Smear		
Platelet Count <i>Sample: Whole Blood EDTA</i> <i>Method: Impedance</i>	338	150 - 410	thou/μL
MPV (Mean Platelet Volume) <i>Sample: Whole Blood EDTA</i> <i>Method: Calculated</i>	8.7	6.8 - 10.9	fL

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Malarial Parasite (MP) Smear*Method: Method: Microscopy***Thin Smear***Sample: Whole Blood EDTA*

Not Detected

Not Detected

Thick Smear*Sample: Whole Blood EDTA*

Not Detected

Not Detected

Widal*Sample: Serum*

Salmonella Typhi 'O'	1:320	< 1:80
Salmonella Typhi 'H'	1:160	< 1:80
Salmonella Paratyphi 'AH'	< 1:80	< 1:80
Salmonella Paratyphi 'BH'	< 1:80	< 1:80
Result :	Positive	



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CLINICAL PATHOLOGY**Urine Routine & Microscopic Examination***Method: Reflectance Photometry***Physical Examination****Colour***Sample: Urine**Method: Physical Examination*

Pale Yellow

Pale Yellow

Appearance*Sample: Urine**Method: Physical Examination*

Slightly Hazy

Clear

Specific Gravity*Sample: Urine**Method: pKa change of pretreated polyelectrolytes*

1.010

1.003 - 1.035

pH*Sample: Urine**Method: Double indicator principle*

5.0

4.7 - 7.5

Chemical Examination**Glucose***Sample: Urine**Method: Glucose oxidase/peroxidase*

Greenish Yellow (++)

Not Detected

Protein*Sample: Urine**Method: Protein-error-of-indicators principle*

Not Detected

Not Detected

Ketones*Sample: Urine**Method: Sodium nitroprusside reaction*

Not Detected

Not Detected

Blood*Sample: Urine**Method: Peroxidase*

Not Detected

Not Detected

Bilirubin*Sample: Urine**Method: Diazo reaction*

Not Detected

Not Detected

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Urobilinogen <i>Sample: Urine</i> <i>Method: Ehrlich's reaction</i>	Normal	Normal	
Nitrite <i>Sample: Urine</i> <i>Method: Nitrite Test</i>	Not Detected	Not Detected	
Microscopic Examination <i>Method: Microscopy</i>			
Pus Cells <i>Sample: Urine</i>	1-2	0 - 5	/hpf
RBC <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
Epithelial Cells <i>Sample: Urine</i>	2-3	0 - 5	/hpf
Casts <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
Crystals <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
Bacteria <i>Sample: Urine</i>	Not Detected	Not Detected	/hpf
Remarks <i>Sample: Urine</i>			

Remarks : Microscopic Examination is performed on urine sediment**Complete Blood Count (CBC)**Clinical Significance :

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CBC comprises of estimation of the cellular components of blood including RBCs, WBCs and Platelets. Mean corpuscular volume (MCV) is a measure of the size of the average RBC, MCH is a measure of the hemoglobin content of the average RBC and MCHC is the hemoglobin concentration per RBC. The red cell distribution width (RDW) is a measure of the degree of variation in RBC size (anisocytosis) and is helpful in distinguishing between some anemias. CBC examination is used as a screening tool to confirm a hematologic disorder, to establish or rule out a diagnosis, to detect an unsuspected hematologic disorder, or to monitor effects of radiation or chemotherapy. Abnormal results may be due to a primary disorder of the cell-producing organs or an underlying disease. Results should be interpreted in conjunction with the patient's clinical picture and appropriate additional testing performed.

Widal

While the definitive diagnosis of typhoid fever depends on the isolation of *S typhi* from blood, stools, urine or other body fluids, the role of the Widal test had been to increase the index of suspicion for the presence of typhoid fever by demonstrating a positive agglutination during the acute and convalescent period of infection with evidence of a four-fold rise of antibody titre. In many developing countries, including India, the Widal test appears to be the only laboratory means employed in the diagnosis of typhoid fever among suspected patients. As the test suffers from serious cross-reactivity with other infectious agents, it may produce false-positive results, leading to a misdiagnosis of typhoid fever. The Widal test reaction involves the use of bacterial suspensions of *S typhi* and *S paratyphi* 'A' and 'B', treated to retain only the 'O' and 'H' antigens. These antigens are employed to detect corresponding antibodies in the serum of a patient suspected of having typhoid fever. The IgM somatic O antibody appears first and represents the initial serologic response in acute typhoid fever, while the IgG flagella H antibody usually develops more slowly but persists for longer.

In an individual with no prior exposure to *S typhi* infection (either lack of active infection or absence of passive immunisation), a higher than 1:80 or 1:160 titre on an initial single test, usually correlates fairly well with exposure to typhoid fever. However, even these single high value titres in an endemic area where repeated exposures to *S typhi* may have occurred, do not have any clinical relevance in the absence of a positive isolate of the causative organism.

Researchers from different parts of India have reported that in normally healthy blood donors, the baseline titre for antibodies to "O" and "H" antigens of *Salmonella enterica* serotype typhi was 1:40 and hence, based on the above results, it could be recommended to use a cutoff level of $\geq 1:80$ for a single antibody test titre. Similarly, baseline titre for antibody to H antigen of *Salmonella enterica* serotype paratyphi A and paratyphi B was 1:80 and the cutoff level was $\geq 1:160$ for a single antibody test titre.

Urine Routine & Microscopic ExaminationClinical Significance :

Urine routine examination and microscopy comprises of a set of screening tests that can detect some common diseases like urinary tract infections, kidney disorders, liver problems, diabetes or other metabolic conditions. Physical characteristics (colour and appearance), chemical composition (glucose, protein, ketone, blood, bilirubin and urobilinogen) and microscopic content (pus cells, epithelial cells, RBCs, casts and crystals) are analyzed



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and reported.

** End of Report**

**Dr. Vikas Gupta**MD Pathologist
Lab Head

Consultant

