

Patient Name:	Mrs. Dummy	Registered on:	11/08/2022 02:39 PM
Age / Sex:	34 YRS / F	Collected on:	11/08/2022 02:57 PM
Referred By:	Dr. Self	Received on:	11/08/2022 02:57 PM
Reg. no. / UHID:	1015 /	Reported on:	11/08/2022 06:39 PM
Investigations:	Pregnancy Profile - 3		



HAEMATOLOGY
COMPLETE BLOOD COUNT (CBC)

TEST	VALUE	UNIT	REFERENCE
Hemoglobin	L 10.0	g/dl	12 - 15
Total Leukocyte Count	H 12,000	cumm	4,000 - 11,000
Differential Leucocyte Count			
Neutrophils	55	%	40 - 80
Lymphocyte	40	%	20 - 40
Eosinophils	02	%	1 - 6
Monocytes	03	%	2 - 10
Basophils		%	< 2
Platelet Count	3.1	lakhs/cumm	1.5 - 4.5
Total RBC Count	4.1	million/cumm	3.9 - 4.8
Hematocrit Value, Hct	39	%	36 - 46
Mean Corpuscular Volume, MCV	95.1	fL	83 - 101
Mean Cell Haemoglobin, MCH	L 24.4	Pg	27 - 32
Mean Cell Haemoglobin CON, MCHC	L 25.6	%	31.5 - 34.5
Mean Platelet Volume, MPV	8	fL	6.5 - 12
R.D.W. - SD	41	fL	39 - 46
R.D.W. - CV	12	%	11.6 - 14
P-LCR	22.3	%	19.7 - 42.4
P.D.W.	H 18.5	fL	9.6 - 15.2

TEST	VALUE	UNIT	REFERENCE
Blood Group & Rh.			
ABO	B		
Rh (ANTI -D)	POSITIVE		


Sachin
SAMPLE

Mr. Sachin Sharma
DMLT, Lab Incharge



Ak Asthana
SAMPLE

Dr. A. K. Asthana
MBBS, MD Pathologist

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BIOCHEMISTRY

TEST	VALUE	UNIT	REFERENCE
Serum Creatinine	0.7	mg/dl	0.5 - 0.8
Random Blood Sugar	99	mg/dl	60 - 160

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SEROLOGY & IMMUNOLOGY

TEST	VALUE	UNIT	REFERENCE
HIV (Card Test)			
HIV - 1	NEGATIVE		
HIV - 2	NEGATIVE		

1. A negative result implies that no Anti HIV – 1 & HIV - 2 antibodies have been detected in the sample by this method. This means that either the patient has not been exposed to HIV-1 or HIV-2 infection or the sample has been tested during the “WINDOW PHASE” (before the development of detectable levels of antibodies).

2. A positive result suggests the possibilities of HIV-I and / or HIV-II infection. However these results must be verified by a confirmatory test (IFA / WESTERN BLOT I-II) before pronouncing the patient positive for HIV-1 and / or HIV-2 infection.

Hepatitis C Virus, HCV	NEGATIVE
VDRL	NON-REACTIVE

LIMITATION:

- The test described here is primarily valid as a screening procedure. It advisable to confirm positive sample with other methods employing specific treponemal antigen e.g tpha, ftaabs etc.
- This method may give false positive result in presence of disease such as leprosy, malaria, toxoplasma, infectious mononucleosis or lupus erythematosus.



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CLINICAL PATHOLOGY

TEST	VALUE	UNIT	REFERENCE
Urine Routine Examination			
Physical Examination			
Quantity	18	ml	
Colour	Pale Yellow		Pale Yellow
Transparency	Clear		Clear
Specific Gravity	1.020		1.005 - 1.03
pH	6		5 - 7
Leukocytes	Present (+)		Absent
Blood	Present (++)		Absent
Chemical Examination			
Protein / Albumin	Present (+++)		Absent
Sugar / Glucose	Present (++++)		Absent
Ketone Bodies	Absent		Absent
Bilirubin	Absent		Absent
Nitrite	Present (+)		Absent
Microscopic Examination			
R.B.C.	Absent	/HPF	Absent
Pus Cells	Absent	/HPF	Absent
Epithelial Cells	Absent	/HPF	Absent
Casts	Absent		Absent
Crystals	Absent		
Bacteria	Absent		Absent
Others	Absent		

~~~ End of report ~~~

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