

PR.

292/05, Tulsidas Marg, Basement Chowk, Lucknow-226 003

: CHA250040331

(BCG)

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CMO Reg. No. RMEE 2445133 NABLReg. No.MC-2491 Certificate No. MIS-2023-0218

Patient Name : B/O.POONAM YADAV Visit No

Registration ON Age/Gender : 6 Y/M 06/Mar/2025 10:52AM Lab No : 10137626 Sample Collected ON 06/Mar/2025 10:54AM Referred By : Dr.KGMU Sample Received ON : 06/Mar/2025 11:09AM

Refer Lab/Hosp : CHARAK NA Report Generated ON 06/Mar/2025 01:21PM

. BLOOD GROUP, CALCIUM, Albumin, PROTEIN, HCV, HBSAg, HIV, RANDOM, PT/PC/INR, NA+K+, CREATININE, UREA, LFT, CBC (WHOLE BLOOD) Doctor Advice

| Test Name | Result | Unit | Bio. Ref. Range | Method |
|---------------|----------|-------|-----------------|----------------------|
| BLOOD GROUP | | | | |
| Blood Group | "B" | | | _ |
| Rh (Anti -D) | POSITIVE | | | |
| SERUM CALCIUM | | | | |
| CALCIUM | 10 | mg/dl | 8.8 - 10.8 | dapta / arsenazo III |
| PROTEIN | | | | |
| PROTEIN Serum | 7.10 | mg/dl | 6.8 - 8.5 | |
| | A second | 10 | | |
| SERUM ALBUMIN | | | | |
| ALBUMIN | 4.7 | gm/dl | 3.20 - 5.50 | Bromcresol Green |

PT/PC/INR PROTHROMBIN TIME 13 Second 13 Second Clotting Assay 100 % 100 % Protrhromin concentration INR (International Normalized Ratio) 1.00 1.0

CHARAK







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| Test Name | Result | Unit | Bio. Ref. Range | Method |
|-------------------------------------|--------|------|-----------------|--------|
| HEPATITIS B SURFACE ANTIGEN (HBsAg) | | | | |
| Sample Type : SERUM | | | | |

HEPATITIS B SURFACE ANTIGEN NON REACTIVE <1 - Non Reactive **CMIA**

>1 - Reactive

Note: This is only a Screening test. Confirmation of the result (Non Reactive/Reactive) should be done by performing a PCR based test.

COMMENTS:

-HBsAg is the first serological marker after infection with Hepatitis B Virus appearing one to ten weeks after exposure and two to eight weeks before the onset of clinical symptoms. HBsAg persists during the acute phase and clears late in the convalescence phase. Failure to clear HBsAg within six months indicates a chronic HBsAg carrier state. HBsAg assays are used to identify the persons infected with HBV and to prevent transmission of the virus by blood and blood products as well as to monitor the status of infected individuals in combination with other hepatitis B serological markers

-Borderline cases must be confirmed with confirmatory neutralizing assay

LIMITATIONS:

- -Results should be used in conjunction with patient history and other hepatitis markers for diagnosis of acute and chronic infections
- -Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may contain human anti-mouse antibodies (HAMA) which may produce anomalous values when tested with assay kits that employs mouse monoclonal
- -Heterophilic antibodies in human serum can react with reagent immunoglobulins, interfering with in vitro immunoassays. Patients routinely exposed to animals or animal serum products can be prone to this interference and anomalous results may be observed.

 -Cross reactivity for specimens from individual with medical conditions (Pregnancy, HIV etc) has been observed.

 -HBsAg mutations may result in a false negative result in some HBsAg assays.

- -If HBsAg results are inconsistent with clinical evidence, additional testing is suggested to confirm the result.



DR. ADITI D AGARWAL



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Patient Name : B/O.POONAM YADAV

Age/Gender : 6 Y/M **Lab No** : 10137626

Referred By : Dr.KGMU
Refer Lab/Hosp : CHARAK NA

: Dr.KGMU : CHARAK NA Sample Received ON : 06/Mar/2025 11:09AM

Visit No

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| Test Name | Result | Unit | Bio. Ref. Range | Method |
|-----------|--------|------|-----------------|--------|
| HIV | | | | |

HIV-SEROLOGY

NON REACTIVE

<1.0 : NON REACTIVE >1.0 : REACTIVE

Done by: Vitros ECI (Sandwich Assay)

Note:-Elisa test is a screening method for HIV.It is known to give false Positive & Negative result.

Hence confirmation: "Western Blot" method is advised.

HEPATITIS C VIRUS (HCV) ANTIBODIES

HEPATITIS C VIRUS (HCV) ANTIBODIES NON REACTIVE

Non Reactive

(TRIO DOT ASSAY)

Note: This is only a Screening test. Confirmation of the result (Non Reactive/Reactive) should be done by performing a PCR based test.

CHARAK



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P.R.

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Age/Gender : 6 Y/M : 06/Mar/2025 10:52AM Registration ON Lab No : 10137626 Sample Collected ON 06/Mar/2025 10:54AM Referred By : Dr.KGMU Sample Received ON : 06/Mar/2025 11:08AM Refer Lab/Hosp : CHARAK NA Report Generated ON 06/Mar/2025 12:23PM

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| Test Name | Result | Unit | Bio. Ref. Range | Method |
|------------------------------|---------|---------|-----------------|----------------|
| CBC (COMPLETE BLOOD COUNT) | | | | |
| Hb | 10.0 | g/dl | 11 - 15 | Non Cyanide |
| R.B.C. COUNT | 6.10 | mil/cmm | 3.8 - 5.2 | Electrical |
| | | | | Impedence |
| PCV | 34.2 | % | 31 - 43 | Pulse hieght |
| | | | | detection |
| MCV | 55.9 | fL | 78 - 81 | calculated |
| MCH | 16.3 | pg | 26 - 28 | Calculated |
| MCHC | 29.2 | g/dL | 33 - 35 | Calculated |
| RDW | 22.8 | % | 11 - 15 | RBC histogram |
| | | | | derivation |
| RETIC | 2.0 % | % | 0.3 - 1 | Microscopy |
| TOTAL LEUCOCYTES COUNT | 12930 | /cmm | 5000 - 15000 | Flocytrometry |
| DIFFERENTIAL LEUCOCYTE COUNT | | | | |
| NEUTROPHIL | 63 | % | 40 - 70 | Flowcytrometry |
| LYMPHOCYTES | 29 | % | 25 - 55 | Flowcytrometry |
| EOSINOPHIL | 4 | % | 1 - 6 | Flowcytrometry |
| MONOCYTE | 4 | % | 0 - 8 | Flowcytrometry |
| BASOPHIL | 0 | % | 00 - 01 | Flowcytrometry |
| PLATELET COUNT | 272,000 | /cmm | 150000 - 500000 | Elect Imped |
| PLATELET COUNT (MANUAL) | 272000 | /cmm | 150000 - 500000 | Microscopy. |
| Absolute Neutrophils Count | 8,146 | /cmm | 2000 - 7000 | Calculated |
| Absolute Lymphocytes Count | 3,750 | /cmm | 1000-3000 | Calculated |
| Absolute Eosinophils Count | 517 | /cmm | 20-500 | Calculated |
| Absolute Monocytes Count | 517 | /cmm | 200-1000 | Calculated |
| Mentzer Index | 9 | | | |
| Peripheral Blood Picture | : | | | |

Red blood cells are microcytic hypochromic with anisocytosis+. Platelets are adequate. No immature cells or parasite seen.









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| Test Name | Result | Unit | Bio. Ref. Range | Method |
|--------------------------------|--------|-------|-----------------|--|
| BLOOD SUGAR RANDOM | , | 1 | | <u>, </u> |
| BLOOD SUGAR RANDOM | 98 | mg/dl | 70 - 170 | Hexokinase |
| NA+K+ | | | | |
| SODIUM Serum | 136.0 | MEq/L | 135 - 155 | ISE Direct |
| POTASSIUM Serum | 4.1 | MEq/L | 3.5 - 5.5 | ISE Direct |
| BLOOD UREA | | | | |
| BLOOD UREA | 35.00 | mg/dl | 15 - 45 | Urease, UV, Serum |
| SERUM CREATININE | | | | |
| CREATININE | 0.60 | mg/dl | 0.50 - 1.40 | Alkaline picrate- kinetic |
| LIVER FUNCTION TEST | | | | |
| TOTAL BILIRUBIN | 0.85 | mg/dl | 0.4 - 1.1 | Diazonium Ion |
| CONJUGATED (D. Bilirubin) | 0.22 | mg/dL | 0.00-0.30 | Diazotization |
| UNCONJUGATED (I.D. Bilirubin) | 0.63 | mg/dL | 0.1 - 1.0 | Calculated |
| ALK PHOS | 378.30 | U/L | 93 - 309 | PNPP, AMP Buffer |
| SGPT | 12.0 | U/L | 5 - 40 | UV without P5P |
| SGOT | 27.0 | U/L | 5 - 40 | UV without P5P |

*** End Of Report ***





