



LABORATORY REPORT

NAME : MR.SI0031 REFERRED BY : SELF VISIT NO : VAMP26148094
AGE : 40Y 0M 0D ZERO TARIFF CLIENT CODE COLLECTED ON : 21-04-2026 10:00
GENDER : Male LAB MR# : AAMP01479373 RECEIVED ON : 21-04-2026 18:11
OP / IP / DG # : APPROVED ON : 22-04-2026 16:30
REPORT STATUS : Final Report



Test Name	Result	Biological Ref. Interval	Unit
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SEROLOGY AND IMMUNOLOGY

Hepatitis A virus (HAV) IgM (Serum)

Hepatitis A virus (HAV) IgM ELISA 0.02 Negative : <0.8 Index
Equivocal : 0.8-1.2
Positive : >1.2

Interpretation:

Hepatitis A Virus (HAV) is a RNA virus of Picornavirus family transmitted by fecal- oral route. Infection with HAV is self limiting though 5-10% cases may show a secondary rise in enzymes. Since symptomatic Hepatitis A virus infections are clinically indistinguishable from Hepatitis B or C virus, serological testing is an extremely important tool to achieve proper diagnosis. During the acute phase of HAV infection, IgM appears in patient's serum in nearly all cases at the onset of symptoms, peaks within the first month of illness and persists for 3-6 months. It declines to undetectable levels within 12 months. The most effective diagnostic determination of HAV acute infection is the detection of Anti HAV- IgM. Patients exhibiting Borderline Reactivity should be monitored at weekly intervals. This will distinguish rising Anti HAV- IgM levels associated with Acute Hepatitis A infection from decreasing or unchanging levels associated with recovery. Rheumatoid factor can give rise to false positive results. False negative / positive results are observed in patients receiving mouse monoclonal antibodies for diagnosis or therapy.

Dr. G. Amitha
MBBS, MD (MICROBIOLOGY)
Consultant Microbiologist

Disclaimer:

1. All results released pertain to the specimen as received by the lab for testing and under the assumption that the patient indicated or identified on the bill/test requisition form is the owner of the specimen.
2. Clinical details and consent forms, especially in Genetic testing, histopathology, as well as wherever applicable, are mandatory to be accompanied with the test requisition form. The non-availability of such information may lead to delay in reporting as well as misinterpretation of test results. The lab will not be responsible for any such delays or misinterpretations thereof.
3. Test results are dependent on the quality of the sample received by the lab. In case the samples are preprocessed elsewhere (e.g., paraffin blocks), results may be compromised.
4. Tests are performed as per the schedule given in the test listing and in any unforeseen circumstances, report delivery may be affected.
5. Test results may show inter-laboratory as well as intra-laboratory variations as per the acceptable norms.
6. Genetic reports as well as reports of other tests should be correlated with clinical details and other available test reports by a qualified medical practitioner. Genetic counselling is advised in genetic test reports by a qualified genetic counsellor, medical practitioner or both.
7. Samples will be discarded post processing after a specified period as per the laboratory's retention policy. Kindly get in touch with the lab for more information.
8. If accidental damage, loss, or destruction of the specimen is not attributable to any direct or negligent act or omission on

Generated On 25-Apr-2026 11:28:00

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Sin No: 20385391

AMPATH
Central Reference Labor
Door No. 1-100/1/CCH N
Serilingampally
Hyderabad



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the part of Ampath Labs or its employees, Ampath shall in no event be liable. Ampath lab's liability for a lack of services, or other mistakes and omissions, shall be restricted to the amount of the patient's payment for the pertinent laboratory services.			

