



LABORATORY REPORT

NAME : MR.SI0432 REFERRED BY : SELF VISIT NO : VAMP26148262
AGE : 40Y 0M 0D ZERO TARIFF CLIENT CODE COLLECTED ON : 21-04-2026 10:00
GENDER : Male LAB MR# : AAMP01479541 RECEIVED ON : 21-04-2026 18:06
OP / IP / DG # : APPROVED ON : 21-04-2026 21:16
REPORT STATUS : Final Report



| Test Name | Result | Biological Ref. Interval | Unit |
|-----------|--------|--------------------------|------|
|-----------|--------|--------------------------|------|

SEROLOGY AND IMMUNOLOGY

SS-A (Ro) IgG Antibodies (Serum)

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|-----------------------------------|------|--|-------|
| SS-A (Ro) IgG Antibodies ELISA | 6.30 | Negative: < 20 Weak Positive: 20-39 Moderate Positive: 40-80 Strong Positive: >80 | RU/mL |
|-----------------------------------|------|--|-------|

Interpretation:

Patients with SLE may have antibodies to SSA/Ro alone or may have both SSA/Ro & SSB/La antibodies. Presence of SSA/Ro antibody alone is commonly seen in association with HLA DR2 in patients less than 22 years of age at onset. Presence of both SSA/Ro & SSB/La in SLE is associated with HLA DR3 and is seen in older patients more than 50 years of age at onset. SLE patients with SSA/Ro antibodies develop a much more serious renal disease and have a higher incidence of concomitant anti DNA antibodies.

Increased levels

- Subacute cutaneous Lupus erthematosus
- Neonatal Lupus erthematosus syndrome with congenial heart block and cutaneous lesions
- Homozygous C2 & C4 deficiency with SLE like disease
- Primary Sjorgen's syndrome vasculitis, Rheumatoid factor positivity & severe systemic symptoms
- ANA negative SLE patients
- SLE with Interstitial patients

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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.





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| <p>5. Test results may show inter-laboratory as well as intra-laboratory variations as per the acceptable norms.</p> <p>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.</p> <p>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.</p> <p>8. If accidental damage, loss, or destruction of the specimen is not attributable to any direct or negligent act or omission on 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.</p> | | | |

