



LABORATORY REPORT

NAME	: MR.SI0080	REFERRED BY	: SELF	VISIT NO	: VAMP26147924
AGE	: 40Y 0M 0D	ZERO TARIFF CLIENT CODE		COLLECTED ON	: 21-04-2026 10:00
GENDER	: Male	LAB MR#	: AAMP01479203	RECEIVED ON	: 21-04-2026 18:08
OP / IP / DG #	:			APPROVED ON	: 22-04-2026 20:14
				REPORT STATUS	: Final Report



Test Name	Result	Biological Ref. Interval	Unit
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SEROLOGY AND IMMUNOLOGY

Anti Saccharomyces cerevisae (ASCA) IgA Antibody (Serum)

ASCA-IGA ENZYM IMMUNO ASSAY	5.23	Negative: 0-20 Equivocal: 20.1-24.9 Positive: >=25	RU/mL
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Interpretation:

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1. A positive result indicates the presence of ASCA IgA antibodies and suggests the possibility Crohn's disease.
2. A specimen with equivocal levels of ASCA IgA cannot be assessed for antibody status. If results remain equivocal after repeat testing, the result should be reported as equivocal and/or an additional sample should be taken.
3. A negative results indicates no ASCA IgA antibody or levels below the negative cut-off of the assay.

Limitations:

1. The presence of immune complexes or other immunoglobulin aggregates in the patient sample may cause an increased level of non-specific binding and produce false positives in this assay
2. A negative ASCA IgA result does not rule out the presence of Crohn's disease.
3. A negative ASCA IgA antibody does not rule out the presence of ASCA antibodies, Because the concentration of antibody may be below the detection limit of the assay.
4. A positive test result only indicates the presence of antibody to S.cerevisiae and does not necessarily indicate the presence of Crohn's disease.
5. This test may be used to complement, But not to substitute for ASCA IgG antibody screening. ASCA IgA results on specimens should not be repeated without the corresponding ASCA IgG results.
6. Results of this assay should be used in conjunction with clinical findings and other serological tests.

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





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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 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.

