AMPATH
Central Reference Laboratory,
Door No. 1-100/1/CCH Nallagandla
Serilingampally
Hyderabad

AMPATH
ACCURATE DIAGNOSTICS, AMERICAN STANDARDS.

LABORATORY REPORT

NAME	**: MR.SI0817**	REFERRED BY	: SELF	VISIT NO	: VAMP26147885
AGE	: 40Y 0M 0D	ZERO TARIFF CLIENT CODE		COLLECTED ON	: 21-04-2026 10:00
GENDER	: Male	**LAB MR#**	**: AAMP01479164**	RECEIVED ON	: 21-04-2026 18:08
OP / IP / DG #	:			APPROVED ON	: 23-04-2026 16:36
				REPORT STATUS	: Final Report

Test Name	Result	Biological Ref. Interval	Unit
SEROLOGY AND IMMUNOLOGY			
ANA Profile 25 Antigens *(Serum)*			
Nucleosome	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
ds DNA	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Anti Histones	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Sm Ab	1.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
RNP 68kD/A/C	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Sm/RNP	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
SS-A/Ro 60kD	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
SS-A/Ro 52kD	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
SS B	1.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
SCL 70	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
ANTI KU	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
PM-SCL(PM)	1.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Mi - 2	2.00	0 - 6 Negative 6 - 12 Intermediate	AU/mL

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JO 1	3.00	>12 Positive 0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
PL7	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
PL-12	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
SRP-54	1.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Ribosomal-P Protein(PO)	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
CENP-A/B	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
P CNA	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
Sp100	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
gp210	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
M2 Recombinant	1.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
M2 native	2.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL
F-actin	3.00	0 - 6 Negative 6 - 12 Intermediate >12 Positive	AU/mL

Interpretation:

ANA25 Screen IgG is an Immunodot kit intended for the detection, in human sera only, of IgG autoantibodies against the following antigens: Nucleosomes, dsDNA, Histones, Sm, RNP 68kD/A/C, Sm/RNP, SSA/Ro 60kD, SSA/Ro 52kD, SSB, Scl-70, Ku, PM-Scl 100, Mi-2, Jo-1, PL-7, PL-12, SRP-54, Ribosome P0, CENP-A/B, PCNA,

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sp100, gp210, M2 recombinant, M2/nPDC and f-Actin.

Antigens	Disease & Prevalence of Autoantibodies
Nucleosome	SLE 56-90 %
dsDNA	SLE 40-90 %
Histones	Drug Induced LE 56-90 %, SLE 50-80%
Sm	SLE 5-40%
RNP 68kD/A/C	MCTD (95-100%), SLE (30-70%), RA(3%)
Sm/RNP	Same as Sm or RNP
SS-A/Ro 60kD	Sjogren's Syndrome (40-95%), SLE (20-60%), Neonatal LE (95-100%)
SS-A/Ro 52kD	Sjogren's Syndrome (40-95%), SLE (10-20%)
SS-B	Sjogren's Syndrome (40-95%), SLE (10-20%)
Scl-70	Scleroderma(20-59%-all, 70%-diffuse), CREST(13%)
Ku	PM/DM & Progressive Systemic Sclerosis (30-55%), SLE (10%)
PM-Scl 100	PM/DM/Overlap Syndromes(50-70%) & Progressive Systemic Sclerosis (5-10%)
Mi-2	Specific for Dermatomyositis
Jo-1	Polymyositis (23-36%)
PL-7	Idiopathic Myositis (2-3%)
PL-12	Idiopathic Myositis (2-3%)
SRP-54	Polymyositis (5%)
Ribosomes P0	SLE (10-20%)
CENP-A/B	Progressive Systemic Sclerosis (80-95%), CREST(57-82%)
PCNA	SLE (3%)
sp100	Primary Biliary Cholangitis
gp210	Primary Biliary Cholangitis
M2 Recombinant	Primary Biliary Cholangitis
M2 native	Primary Biliary Cholangitis
F-actin	Autoimmune Hepatitis (AIH Type I)

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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 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.

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