

Laboratory Investigation Report

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

Serology Special

Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
Anti Cardiolipin Ab,IgG,Serum			
Anti Cardiolipin IgG FEIA	1.0	GPL-U/mL	< 10.0

Ref. Range

Negative < 10
Weak Positive 10 - 40
Positive > 40

Comment :

Cardiolipin antibodies is detected in autoimmune disorders particularly systemic lupus erythematosus (SLE), vascular thrombosis, thrombocytopenia etc. Elevations of cardiolipin antibody is associated with increased risk in idiopathic thrombocytopenia purpura, rheumatoid, psoriatic, arthritis primary sjogrem's syndrome.

Interpretation :

Cardiolipin IgG is intended for the in vitro quantitative measurement of IgG antibodies directed to cardiolipin in serum and plasma to aid in the diagnosis of antiphospholipid syndrome (APS) and to evaluate the thrombotic risk in patients with systemic lupus erythematosus (SLE). A definitive clinical diagnosis should not be based on the results of a single diagnostic method, but should only be made after all clinical and laboratory findings have been evaluated.

Test Performed at :910 - Max Hospital - Saket M S S H, Press Enclave Road, Mandir Marg, Saket, New Delhi, Delhi 110017

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Booking Centre :1103 - Max Hospital Saket(East Block), 1, 2, Press Enclave Marg, Saket Institutional Area, Saket, New Delhi, 7982100200

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Max Super Speciality Hospital, Saket (West Block), 1, Press Enclave Road, Saket, New Delhi - 110 017, Phone: +91-11-6611 5050

(CIN No.: U85100DL2021PLC381826)

Helpline No. 7982 100 200 www.maxlab.co.in feedback@maxlab.co.in

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MC-2714

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Serology Special

Max-Thrombophilia Profile (Extended)



Test Name	Result	Unit	Bio Ref Interval
Anti Cardiolipin Ab,IgM,Serum			
Anti Cardiolipin IgM	61	MPL-U/mL	< 10.0
Rechecked			

Ref. Range

Negative < 10
 Equivocal 10 - 40
 Positive >40

Comment :

Cardiolipin antibodies is detected in autoimmune disorders particularly systemic lupus erythematosus (SLE), vascular thrombosis, thrombocytopenia etc. Elevations of cardiolipin antibody is associated with increased risk in idiopathic thrombocytopenia purpura, rheumatoid, psoriatic, arthritis primary sjogren's syndrome.

Interpretation :

Cardiolipin IgM is intended for the in vitro quantitative measurement of IgM antibodies directed to cardiolipin in serum and plasma to aid in the diagnosis of antiphospholipid syndrome (APS) and to evaluate the thrombotic risk in patients with systemic lupus erythematosus (SLE). A definitive clinical diagnosis should not be based on the results of a single diagnostic method, but should only be made after all clinical and laboratory findings have been evaluated. Rheumatoid factor (RF) can interfere with the determination of IgM anti-cardiolipin antibodies.

Kindly correlate with clinical findings

*** End Of Report ***



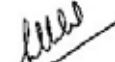
Dr. Poonam.S. Das, M.D.
 Principal Director-
 Max Lab & Blood Bank Services



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 Associate Director
 Microbiology & Molecular Diagnostics



Dr. Sonu Kumari Agrawal, MD
 Associate Consultant
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Dr Nidhi Malik, MD
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Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
Anti Thrombin - III - Functional, Citrate Plasma			
Antithrombin III Functional Chromogenic assay	116	%	10-150


Interpretation Syn – Antithrombin III

Antithrombin is a small protein molecule that inactivates several enzymes of the coagulation system. Low levels of AT are found in 4-5% patients with unexplained VTE.

Reduced levels are seen in Hereditary deficiency, chronic liver diseases, heparin therapy, pregnancy (3rd trimester), acute leukemia, burns and renal diseases.

Kindly correlate with clinical findings


*** End Of Report ***



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Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
Beta-2 Glycoprotein 1, IgG, Serum FEIA			
Beta-2 Glycoprotein 1, IgG	0.7	U/mL	

Ref Range :-

Negative < 7.0
 Equivocal 7 - 10
 Positive > 10

Interpretation :

Detection of Beta-2 Glycoprotein antibodies are indicative of risk for thrombosis in autoimmune diseases.

Beta-2 Glycoprotein I IgG is intended for the in vitro quantitative measurement of IgG antibodies directed to β 2-Glycoprotein I in human serum and plasma to aid in the diagnosis of antiphospholipid syndrome (APS) and to evaluate the thrombotic risk in patients with systemic lupus erythematosus (SLE). A definitive clinical diagnosis should not be based on the results of a single diagnostic method, but should only be made after all clinical and laboratory findings have been evaluated.

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Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
Beta-2 Glycoprotein 1, IgM, Serum FEIA			
Beta-2 Glycoprotein 1, IgM	0.9	U/mL	

Ref Range :-

Negative < 7.0
 Equivocal 7 - 10
 Positive > 10

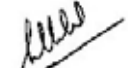
Interpretation :

Detection of Beta-2 Glycoprotein antibodies are indicative of risk for thrombosis in autoimmune diseases.

Beta-2 Glycoprotein I IgM is intended for the in vitro quantitative measurement of IgM antibodies directed to β 2-Glycoprotein I in human serum and plasma to aid in the diagnosis of antiphospholipid syndrome (APS) and to evaluate the thrombotic risk in patients with systemic lupus erythematosus (SLE). A definitive clinical diagnosis should not be based on the results of a single diagnostic method, but should only be made after all clinical and laboratory findings have been evaluated. Rheumatoid factor (RF) can interfere with the determination of IgM anti- β 2-Glycoprotein I antibodies.

Kindly correlate with clinical findings

*** End Of Report ***

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Hematology Special



SIN No:DF1069528

Max-Thrombophilia Profile (Extended)

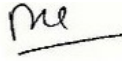
Test Name	Result	Unit	Bio Ref Interval
DRVVT-Lupus Anticoagulant , Plasma Citrate			
Electromechanical Clot Detection			
dRVVT Screen	40.70	Sec	29.9 - 47.1
dRVVT Screen ratio	1.06		
dRVVT Confirm	34.80	Sec	24.8 - 34.1
dRVVT Confirm ratio	1.18		
dRVVT Screen: Confirm ratio	0.90		0.0 - 1.20
Interpretation	No Lupus Like Anticoagulant Present		

Kindly correlate with clinical findings

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Hematology



Max-Thrombophilia Profile (Extended)

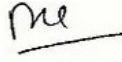
Test Name	Result	Unit	Bio Ref Interval
Factor VIII Studies,Citrate Plasma			
Photo-Optical-Clot Detection			
Factor VIII Assay Based on APTT Assay	142.5	%	50-150

Kindly correlate with clinical findings

*** End Of Report ***



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Max-Thrombophilia Profile (Extended)



Test Name	Result	Unit	Bio Ref Interval
Free Protein S,Citrate Plasma			
Protein S, Free Latex Ligand Immunoassay	83.5	%	74.1-146.1

Interpretation Protein S is a vitamin K-dependent plasma glycoprotein synthesized in the liver. It functions as a cofactor to Protein C in the inactivation of Factors Va and VIIIa and plays a role in anticoagulation pathway.

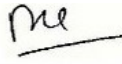
Reduced levels predispose to VTE. It can be seen in hereditary deficiency, pregnancy, Oral anticoagulant e.g. Warfarin, nephritic syndrome and liver diseases.

Kindly correlate with clinical findings

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Clinical Biochemistry
Max-Thrombophilia Profile (Extended)




Homocysteine, Quantitative, Serum

Date	08/Aug/2023 04:46AM	Unit	Bio Ref Interval
Homocysteine, Quantitative Enzymatic Kinetic	23.5	µmol/L	3 - 12


Interpretation Measurement of Homocysteine is considered important to diagnose homocystinuria, to identify individuals with or at a risk of developing cobalamin or folate deficiency, and to assess Homocysteine as a risk factor for cardiovascular disease (CVD) and other disorders.

Kindly correlate with clinical findings

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SIN No:DF1069528

Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
Protein C, Functional, Sodium Citrate Automated Chromagenic Assay			
Protein C, Functional	66	%	70 - 140

Interpretation

Protein C is a zymogen, the activated form of which plays an important role in regulating anticoagulation, inflammation, cell death, and maintaining the permeability of blood vessel walls in humans and other animals.

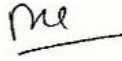
Reduced levels predispose to VTE. It can be seen in hereditary deficiency, pregnancy, Oral anticoagulant e.g. Warfarin, malignancy and liver diseases.

Kindly correlate with clinical findings

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Molecular Diagnostics



Max-Thrombophilia Profile (Extended)

Test Name	Result	Unit	Bio Ref Interval
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Factor V Leiden Mutational Analysis

Real Time PCR

Factor V Leiden Mutation Real Time PCR	Not Detected		
Prothrombin Gene Mutation	Not Detected		
MTHFR Gene Mutation (C677T)	Not Detected		
MTHFR Gene Mutation (A1298C)	Heterozygous Mutation Detected		

Interpretation

Result	Comments
Homozygous Mutation Detected	Both alleles carry mutation
Heterozygous Mutation Detected	Single allele carries mutation
Not Detected	Both alleles do not carry mutation

Note

- This is an in-house developed qualitative assay.
- All results should be interpreted in context of clinical findings.
- This assay detects the following mutations:
 - Factor V Leiden (R506Q)
 - Factor II Prothrombin Gene mutation (G20210A)
 - MTHFR Gene (C677T; A1298C)
- Test conducted on Whole blood.
- Presence of PCR inhibitors if any, might lead to amplification failure.

Comments

The most common identifiable genetic defects in Venous thromboembolism is the factor V (R506Q) Leiden mutation which causes resistance to activated protein C (APC). APC resistance results in thrombotic predisposition via the destruction of the activated protein C cleavage & inactivation site in the factor V procoagulant protein. The factor V Leiden mutation is extremely common; heterozygotes represent 3% to 7% of the general population, approximately 20% of all patients with any venous thrombosis, and approximately 50% of patients with recurrent venous thrombosis.

The second most common thrombophilic genetic defect is the prothrombin G20210A mutation, imparting a 2- to 5-fold increased risk for venous thromboembolism (in heterozygotes) and being present in heterozygous form in 15% to 20% of patients with thrombophilia.

Genetic polymorphism associated with severe MTHFR deficiency is defined by a C to T substitution at position 677 (C677T) and/or A to C substitution at position 1298 (A1298C) of the *MTHFR* gene. These mutations lead to the incorporation of amino acid alanine (A) instead of valine (V) at position 222 and glutamate to alanine substitution at codon 429 respectively of the MTHFR protein. These mutations may lead to hyperhomocysteinemia

Uses

Venous thrombosis is a multifactorial disease frequently related to the interaction of genetic and environmental risk factors. Testing for specific mutations in these patients helps to determine the decision on the duration of anticoagulant therapy and risk stratification for primary or secondary prophylaxis. This test is used as a

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Molecular Diagnostics



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Test Name	Result	Unit	Bio Ref Interval
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thrombosis risk factor in patients prior to major surgery, pregnancy, postpartum, oral contraceptive use, estrogen replacement therapy, transient ischemic attacks, premature stroke, peripheral vascular disease, pulmonary embolism & family history of thrombosis or known Factor V mutations in the family.

Kindly correlate with clinical findings

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