

## Gurugram

Pathkind Diagnostics Pvt. Ltd. Plot No. 55-56, Udhyog Vihar Ph-IV, Gurugram - 122015

# Processed By Pathkind Diagnostics Pvt. Ltd.

Plot No. 55-56, Udhyog Vihar Ph-IV, Gurugram - 122015

Name : Mrs. PL160		Billing Date :	07/07/202312:25:42
Age : 35 Yrs		Sample Collected on :	10/07/2023 10:01:31
Sex : Female		Sample Received on :	10/07/2023 11:02:13
P. ID No. : P1000100012819		Report Released on :	15/07/2023 17:42:38
Accession No : 10002304875		Barcode No. :	10002304875-02,
Referring Doctor : Self			10002304875-01
Referred By :		Ref no. :	
F	Report Status - Final		
Test Name	Result	Biological Ref. Interval	Unit
	<u>Haematology</u>		
APLA Profile			
PTT & Mixing Studies Plasma			
PTT (Test) Sample: Citrate Plasma	26.40	24.80 - 34.40	Sec
PTT Control (Normal Pooled Plasma) Sample: Citrate Plasma	28.4	24.8 - 34.4	Sec
Lupus Anticoagulant Screen Time			
DRVV Screen Test Sample: Citrate Plasma	34.60	33.80 - 45.80	Sec
DRVV Screen Control Sample: Citrate Plasma	39.0	33.8 - 45.8	Sec
DRVV Screen Ratio Sample: Citrate Plasma	0.92	<1.20	Ratio
Lupus Anticoagulant Sample: Citrate Plasma	Absent		Sec
	<b>SEROLOGY</b>		
Cardiolipin IgG Antibodies	7.20	Negative : < 10	GPL-U/ml
Sample: Serum Method: ELISA	7.20	Positive: >=10	
Cardiolipin IgM Antibodies Sample: Serum Method: ELISA	0.02	Negative : <7 Positive: >/= 7	MPLU/mL
Anti Phospholipid IgG Antibodies Sample: Serum Method: ELISA	3.40	Negative < 10 U/mL Positive : > =10 U/mL	U/mL
Anti Phospholipid IgM Antibodies Sample: Serum Method: ELISA	1.20	Negative : < 10 U/mL	U/mL

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 ☆ care@pathkindlabs.com | ⊕ www.pathkindlabs.com
 ⓒ Customer Care: 75000-75111



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Positive : > = 10 U/mL

## **Lupus Anticoagulant**

Medical Remarks: See Remark - 5. Correlate Clinically.

**Test Description:** Screening of Lupus Anticoagulant Confirmation is done by two different APTT reagents namely Lupus sensitive (LS) APTT Automate and RVVT. Lupus anticoagulant is an antiphospholipid antibody directed against negatively charged phospholipids that is identified functionally by prolongation of in vitro phospholipids dependent coagulation test. Lupus anticoagulant is often associated with a thrombotic tendency.

#### Interpretation :

1. Following is the final Interpretative Table using the findings of both the above mentioned tests:

APTT (LS)	APTT Mixing (Using	DRVVT (Screen)	DRVVT	DRVV screen to	Interpretation
	Rosner Index)		(Confirmation)	confirmation ratio	
Normal	-	Normal	-	-	LAC Absent
Abnormal	Corrected	Normal	-	-	Suspect Factor Deficiency
Abnormal	Not Corrected	Abnormal	Normal	> 1.2	LAC Present
Normal	-	Abnormal	Normal	> 1.2	LAC Present
Abnormal	Corrected/Partially Corrected	Abnormal	Normal	> 1.2	Factor Deficiency + LAC Present
Abnormal	Not Corrected	Abnormal	Abnormal	< 1.2	Either Inhibitor or LAC
Abnormal	Not Corrected	Normal	-	-	Suspect Inhibitor/heparin

2. Interpretation of APTT mixing based on Rosner Index (RI) Rosner Index = Clotting time of mixture - Clotting time of normal (PPP) X 100/Clotting time of patient plasma \* If Rosner Index < 12 then APTT mixing is considered as corrected and Suspects the factor deficiency.

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\* If Rosner Index < 12 then APTT mixing is considered as corrected and Suspects the factor deficiency.

\* If Ronser Index > 12 then APTT mixing is considered as not corrected and suspects the presence of inhibitor or Lupus.

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	-		
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3. All abnormal results of DRVV screening tests are confirmed by neutralization assay using Phospholipids. A normalized ratio > or = Biological Reference Interval confirms the presence of lupus anticoagulant (LAC).

4. Persistent Positive LAC results on two different occasions & 12 weeks apart are essential to suspect & diagnose Lupus anticoagulant.

5. In view of abnormal DRVV Screen, abnormal DRVV confirm and normalised ration <1.2, possibility of Lupus and/or Inhibitor needs exclusion. Advice repeat testing after 12 weeks.

Limitations :

1. False Positive: Patients on heparin or heparin substitute; Coagulation factor VIII inhibitors.

2. False Negative: Elevated factor VIII levels, as may be seen in an acute infection or with replacement therapy when someone has Hemophillia A, may shorten the aPTT time, leading to a temporary false negative test for lupus anticoagulant.

Reference : Update of the guidelines for lupus anticoagulant detection. Pengo V, Tripodi A, et al. J Thromb Haemost 2009;7:1737-40.

## **Cardiolipin IgG Antibodies**

#### **Clinical Significance :**

ANTI-CARDIOLIPIN ANTIBODIES (ACA) are antibodies ofter directed against cardiolipin and found in several diseases

1. The presence of anti cardiolipin antibodies in Systemic Lupus Erythematosus (SLE) can be related to the development of thrombosis and thrombocytopenia.

2. In Gynecology practice they are associated with Intrauterine Death or recurrent abortions and unexplained infertility.

3. They are also found in some non thrombotic neurological disorders e.g. cerebrovacsular insufficiency, cerebral ischemia or chorea. Transient elevation can be seen inother autoimmune & intercurrent diseases. Persistent positive test with titre more than 40 GPL/ml and spaced atleast 12 weeks apart is significant in antiphopholipid antibody syndrome.

## **Cardiolipin IgM Antibodies**

Clinical Significance : ANTI-CARDIOLIPIN ANTIBODIES (ACA) are antibodies ofter directed against cardiolipin and found in several diseases

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0002304875 Mrs. PL160

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## Anti Phospholipid IgG Antibodies

Interpretation:

- Elevated levels of anti phospholipid antibodies(APLA) are seen in cases of Systemic Lupus Erythematosis(SLE) and other Systemic autoimmune disorders like Rheumatoid arthritis, Scleroderma, Sjogrens's syndrome etc.
- The occurence of APLA in patients with SLE and related diseases indicates secondary anti phospholipid syndrome(APS).
- Presence of APS with no other autoimmune disease indicates primary APS.
- Persisting high APLA titres are considered as a risk factor for thrombovascular complications.

## Anti Phospholipid IgM Antibodies

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- Elevated levels of anti phospholipid antibodies(APLA) are seen in cases of Systemic Lupus Erythematosis(SLE) and other Systemic autoimmune disorders like Rheumatoid arthritis, Scleroderma, Sjogrens's syndrome etc.
- The occurence of APLA in patients with SLE and related diseases indicates secondary anti phospholipid syndrome(APS).
- Presence of APS with no other autoimmune disease indicates primary APS.
- Persisting high APLA titres are considered as a risk factor for thrombovascular complications.

\*\* End of Report\*\*

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