

Client

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	: Mr. CO19		Billing Date	:	07/07/202312:21:21	
Age	: 35 Yrs		Sample Collected on	:	10/07/2023 10:01:31	
Sex	: Male		Sample Received on	:	10/07/2023 11:02:13	
P. ID No.	: P1000100012726		Report Released on	:	14/07/2023 15:43:03	
Accession No	: 10002304782		Barcode No.	:	10002304782-01	
Referring Doctor : Self						
Referred By	:		Ref no.	:		
		Report Status - Final				
Test Name		Result	Biological Ref. Interva	al	Unit	
HAEMATOLOGY						

Protein S Antigen- Free	101.0	70.0 - 148.0	%
Sample: Citrate Plasma			

Protein S Antigen- Free

Free protein S measures the antigen and not the activity.

1. Protein S is a vitamin K dependent plasma glycoprotein: 60% bound to C4bBP-b chain, 40% free. Protein S possesses both APCdependent and independent anticoagulant properties and thus is an important guardian in controlling thrombin generation and fibrinolysis.

2. Protein S deficiency may be associated with 3 to 10 fold increased risk of venous thrombosis, recurrent miscarriage, complications of pregnancy (preeclampsia, abruptio, placentae, intrauterine growth restriction, and stilbirth) and possibly arterial thrombosis.

3. Three subtypes of PS deficiency are recognized, types I and III (also known as type IIa) are quantitative defects while type II is very rare and is a quanlitative defect. It can be inherited or acquired.

4. Acquired Protein S deficeincy - vitamin K-antagonis therapy, oral contraceptives, pregnancy and various disorders, such as liver diseases, nephritic syndrome, disseminated intravascular coagluation and chronic infections.

5. Congenital Protein S deficiency - is rare autosomal dominant disorder with 1 to 3% incidence of venous thromboembolism in adults.

6. Using lower cut-off levels of Protein S activity that is 40% which is equal to the highest Protein S activity value found in

heterozygouscarriers with mutations in PROS1, the diagnostic specificity for risk of thromboembolism increases.

7. Associated APC Resistance (Factor V leiden), increased factor VIII levels, warfarin therapy can cause false decrease in protein S activity.

8. Associated heparin therapy, lupus anticoagulant can cause false increase in protein S activity.

9. Repeat testing is recommended to confirm diagnosis after at least 4-6 weeks.

** End of Report**

Dr. Aarti Khanna Nagpal DNB (Pathology) Senior Consultant

Page No: 1 of 1



