

Client: ARUP Example Report Only 500 Chipeta Way Salt Lake City, UT 84108 UNITED STATES

Physician: arup, arup

Patient: Test, PAI-1 GENO 5G

DOB

Sex: Male **Patient Identifiers:** 46500 Visit Number (FIN): 46829

Collection Date: 3/3/2023 10:36

Plasminogen Activator Inhibitor-1, PAI-1 (SERPINE1) Genotyping

ARUP test code 2004980

PAI-1 Specimen Whole Blood

PAI-1 Interpretation

5G/5G

Indication for testing: Assess genetic risk for thrombosis.

Result: PAI-1 Genotype 5G/5G

Interpretation: Two copies of the PAI-1 5G allele (5G/5G) were detected. The 4G allele, which is associated with increased PAI-1 plasma activity levels, was not detected. This result does not exclude other genetic, or non-genetic, causes for thrombosis.

Recommendations: If clinically indicated, testing for other inherited or acquired thrombophilic disorders is recommended. For additional testing information, refer to the Antiphospholipid Syndrome, Hypercoagulable States, Thrombophilia, or Venous Thromboembolism topics at arupconsult.com.

This result has been reviewed and approved by

Background Information: Plasminogen Activator Inhibitor-1, PAI-1 (SERPINE1) Genotyping:

Characteristics: The 4G allele within in the promoter region of the PAI-1 (SERPINE1)gene is associated with higher plasma PAI-1 activity when compared with the 5G allele. Heterozygosity or homozygosity for the 4G allele confers a risk for venous thromboembolism (VTE), especially in individuals with other thrombophilic risk factors, as well as a risk for myocardial

Frequency of the 4G Allele: Caucasian 0.52, Hispanic 0.38,

African-American 0.13-0.28.

Variant Tested: The PAI-1 promoter 4G/5G polymorphism located in the promoter region of the SERPINE1 gene. NM_000602.3(SERPINE1) c.-817dupG (from start of translation).

Inheritance: Autosomal dominant.

Clinical sensitivity: Unknown.
Methodology: Polymerase chain reaction and fluorescence

monitoring.

Analytical Sensitivity and Specificity: 99 percent.

Limitations: Variants in the PAI-1 (SERPINE1) gene, other than the 4G/5G polymorphism, are not evaluated. Diagnostic errors can occur due to rare sequence variations.

This test was developed and its performance characteristics

H=High, L=Low, *=Abnormal, C=Critical



determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Counseling and informed consent are recommended for genetic testing. Consent forms are available online.

VERIFIED/REPORTED DATES				
Procedure	Accession	Collected	Received	Verified/Reported
PAI-1 Specimen	23-062-104017	3/3/2023 10:36:00 AM	3/3/2023 10:36:21 AM	4/6/2023 2:30:00 PM
PAI-1 Interpretation	23-062-104017	3/3/2023 10:36:00 AM	3/3/2023 10:36:21 AM	4/6/2023 2:30:00 PM

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

ARUP LABORATORIES | 800-522-2787 | aruplab.com 500 Chipeta Way, Salt Lake City, UT 84108-1221 Jonathan R. Genzen, MD, PhD, Laboratory Director