

Client: Example Client ABC123
123 Test Drive
Salt Lake City, UT 84108
UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB: Unknown
Gender: Female
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
Collection Date: 00/00/0000 00:00

Platelet Antigen Genotyping Panel

ARUP test code 3000193

Platelet Antigen Geno Specimen whole blood

Platelet Antigen 1 Genotyping a/b

Platelet Antigen 2 Genotyping a/a

Platelet Antigen 3 Genotyping b/b

Platelet Antigen 4 Genotyping a/a

Platelet Antigen 5 Genotyping a/a

Platelet Antigen 6 Genotyping a/a

Platelet Antigen 15 Genotyping a/b

Platelet Antigen Geno Interpretation See Note

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

Unless otherwise indicated, testing performed at:

Indication for testing: Parental or neonatal genotyping to assess risk for alloimmune thrombocytopenia.

HPA-1a/b Heterozygous: One copy of the common human platelet antigen (HPA)-1 "a" allele and one copy of the less-common HPA-1 "b" allele were identified.

HPA-2a/a Homozygous: Two copies of the common human platelet antigen (HPA)-2 "a" allele were identified.

HPA-3b/b Homozygous: Two copies of the less-common human platelet antigen (HPA)-3 "b" allele were identified.

HPA-4a/a Homozygous: Two copies of the common human platelet antigen (HPA)-4 "a" allele were identified.

HPA-5a/a Homozygous: Two copies of the common human platelet antigen (HPA)-5 "a" allele were identified.

HPA-6a/a Homozygous: Two copies of the common human platelet antigen (HPA)-6 "a" allele were identified.

HPA-15a/b Heterozygous: One copy of the human platelet antigen (HPA)-15 "a" allele and one copy of the HPA-15 "b" allele were identified.

This result has been reviewed and approved by [REDACTED]

BACKGROUND INFORMATION: Platelet Antigen Genotyping Panel

Characteristics: Spontaneous fetal intracranial bleeding may occur in 20 percent of pregnancies affected with severe perinatal alloimmune thrombocytopenia (PAT); there is a risk of fetal death. Post-transfusion purpura may occur in transfusion recipients with antibodies to a specific platelet antigen. Incidence: PAT occurs in 1 in 5000 births.

Inheritance: For women homozygous for the less common "b" HPA allele with antibodies to the common "a" allele, there is a 50 percent risk a pregnancy will be at risk if her partner is heterozygous for the "a" allele and 100 percent risk if her partner is homozygous for the "a" allele.

Cause: Maternal-fetal HPA incompatibility.
Polymorphisms Tested: HPA-1 (ITGB3, GPIIIa) c.176T>C, p.L59P; HPA-2 (GP1BA, GPIba) c.482C>T, p.T161M; HPA-3 (ITGA2B, GPIIb) c.2621T>G, p.I874S; HPA-4 (ITGB3, GPIIIa) c.506G>A, p.R169Q; HPA-5 (ITGA2, GPIa) c.1600G>A, p.E534K; HPA-6 (ITGB3, GPIIIa) c.1544G>A, p.R515Q; HPA-15 (CD109, CD109) c.2108C>A, p.S703Y
Clinical Sensitivity: Variable; dependent on ethnicity.
Methodology: Polymerase Chain Reaction (PCR)/Fluorescence Monitoring.

Analytic Sensitivity and Specificity: 99 percent.
Limitations: Diagnostic errors can occur due to rare sequence variations. Interpretation of this test result may be impacted if this patient has had an allogeneic stem cell transplantation.

This test was developed and its performance characteristics 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.

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

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
Platelet Antigen Geno Specimen	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 1 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 2 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 3 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 4 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 5 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 6 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 15 Genotyping	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen Geno Interpretation	24-248-101776	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

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

Unless otherwise indicated, testing performed at:

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

Patient: Patient, Example
ARUP Accession: 24-248-101776
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
Page 3 of 3 | Printed: 9/4/2024 11:17:50 AM
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