

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, Fetal

ARUP test code 3016673

Platelet Antigen 1 Genotyping a/a

Platelet Antigen 2 Genotyping a/a

Platelet Antigen 3 Genotyping a/a

Platelet Antigen 4 Genotyping a/a

Platelet Antigen 5 Genotyping a/a

Platelet Antigen 6 Genotyping a/a

Platelet Antigen 15 Genotyping a/a

Maternal Contamination Study Fetal Spec Fetal Cells

Single fetal genotype present; no maternal cells present. Fetal and maternal samples were tested using STR markers to rule out maternal cell contamination.

Maternal Contam Study, Maternal Spec whole Blood

Platelet Antigen Geno, Fetal Specimen Direct Amnio

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

Unless otherwise indicated, testing performed at:

Platelet Antigen Geno Fetal, Interp

See Note

Indication for testing: Fetal genotyping to assess risk for fetal alloimmune thrombocytopenia.

Interpretation: Fetal human platelet antigen (HPA) genotypes should be correlated with parental HPA genotypes for interpretation. A genetic discrepancy between maternal and fetal HPA is necessary, but not sufficient, to diagnose fetal alloimmune thrombocytopenia. Maternal antibody testing for genetically discrepant antigen(s) is recommended.

HPA-1a/a Homozygous: Two copies of the common human platelet antigen (HPA)-1 "a" allele were identified.

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

HPA-3a/a Homozygous: Two copies of the common human platelet antigen (HPA)-3 "a" 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/a Homozygous: Two copies of the human platelet antigen (HPA)-15 "a" allele were identified.

This result has been reviewed and approved by [REDACTED]

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

BACKGROUND INFORMATION: Platelet Antigen Genotyping Panel, Fetal

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, GPIIb) 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/Fragment Analysis.

Analytic Sensitivity and Specificity: 99 percent.

Limitations: Bloody amniotic fluid specimens may give false-negative results because of maternal cell contamination. Diagnostic errors can occur due to rare sequence variations.

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 1 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 2 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 3 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 4 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 5 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 6 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen 15 Genotyping	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Maternal Contamination Study Fetal Spec	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Maternal Contam Study, Maternal Spec	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen Geno, Fetal Specimen	23-235-100853	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Platelet Antigen Geno Fetal, Interp	23-235-100853	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: 23-235-100853
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
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