

**Specimen Collected: 29-Oct-25 13:34****Chimerism, Post-Transplant, Granulocytes** | **Received: 29-Oct-25 13:34** **Report/Verified: 10-Nov-25 15:00**

Procedure	Result	Units	Reference Interval
Chimerism Post-Granulocytes, Specimen	Whole Blood		
Chimerism Post-Gran, InformativeLoc	9		
Chimerism Post-Gran, Percent Recip	53	%	
Chimerism Post-Gran, Percent Donor	47	%	
Chimerism Post-Gran, Margin Error 3			
Chimerism Post-Gran, Interpretation	Mixed <sup>f1 i1</sup>		

**Result Footnote**

f1: Chimerism Post-Gran, Interpretation

The granulocyte cell content of the isolated fraction typically ranges from 97.2 - 98.8%.

This result has been reviewed and approved by [REDACTED]

**Test Information**

i1: Chimerism Post-Gran, Interpretation

BACKGROUND INFORMATION: Chimerism, Posttransplant, Sorted Cells (Granulocytes)

INDICATION: Monitoring for bone marrow transplant patients; correlation with clinical status and consideration of the interval between bone marrow transplantation and testing is necessary for proper interpretation of results.

METHODOLOGY: PCR followed by capillary electrophoresis. Specimens are analyzed using 15 autosomal markers (D8S1179, D21S11, D7S820, CSF1PO, D3S1358, TH01, D13S317, D16S539, D2S1338, D19S433, vWa, TPOX, D18S51, D5S818, and FGA) and one gender marker (amelogenin).

LIMIT OF DETECTION: 2 percent of minor cell population.

LIMITATIONS: 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 U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

\*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

**Unless otherwise indicated, testing performed at:****ARUP Laboratories**

500 Chipeta Way, Salt Lake City, UT 84108

Laboratory Director: Jonathan R. Genzen, MD, PhD

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