

TEST CHANGE

Maternal T Cell Engraftment in SCID, Maternal Specimen 2014704 SCID-MAT

2014704, SCID-MAT	
Specimen Requirements:	
Patient Preparation:	
Collect:	Lavender (EDTA), pPink (K2EDTA), or yYellow (ACD sSolution A). New York State Clients:?-Lavender (EDTA)
Specimen Preparation:	Transport 2 mL whole blood. (Min: 1 mL) New York State Clients: Transport 58 mL whole blood. (Min: 34 mL).
Transport Temperature:	Refrigerated. Also acceptable: Ambient.
Unacceptable Conditions:	
Remarks:	
Stability:	Room Temperature: 1 week; Refrigerated: 1 month; Frozen: unacceptable New York State Clients:?-Room Temperature: 7 days; Refrigerated: 14 days; Frozen: Unacceptable
Methodology:	Polymerase Chain Reaction (PCR) / Hragment Analysis
Performed:	Sun-Sat
Reported:	5-9 days
Note:	To complete Maternal T Cell Engraftment in SCID testing, samples should be collected to perform the following three tests: (1) A buccal brush collected from the patient for Maternal T Cell Engraftment in SCID, Pre-Engraftment Specimen (ARUP test code 2014694), used as a genetic baseline for the patient. (2) A peripheral blood sample from the biological mother for Maternal T Cell Engraftment in SCID, Maternal Specimen (ARUP test code 2014704), used as a genetic baseline for the mother. (3) A peripheral blood sample collected from the patient for Maternal T Cell Engraftment in SCID (ARUP test code 2014699). T cells isolated from the blood sample will be genotyped for comparison to the patient and biological mother baseline genotypes. If T-cell sorting is not completed on the blood sample before submission of Maternal T Cell Engraftment in SCID (ARUP test code 2014699), BMT Cell Isolation (ARUP test code 2005498) will be added to each order of Maternal T Cell Engraftment in SCID (ARUP test code 2014699). Additional charges apply for cell isolation.
CPT Codes:	See CPT code for Maternal T Cell Engraftment in SCID, Pre-

Effective Date: October 6, 2025



Engraftment Specimen (ARUP test code 2014694)

New York DOH Approval Status: Specimens from New York clients will be sent out to a New York DOH approved laboratory, if possible.

Interpretive Data:
Refer to report.

Reference Interval:

Effective Date: October 6, 2025