



Medical Coverage Policy

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Laboratory Testing for Transplantation Rejection

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Related Coverage Resources

- [Heart, Lung, and Heart-Lung Transplantation](#)
- [Kidney Transplantation, Pancreas-Kidney Transplantation, and Pancreas Transplantation Alone](#)
- [Liver and Liver-Kidney Transplantation](#)
- [Magnetic Resonance Imaging \(MRI\), Cardiac Laboratory Management Clinical Guidelines](#)

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covered under this Coverage Policy (see "Coding Information" below). When billing, providers must use the most appropriate codes as of the effective date of the submission. Claims submitted for services that are not accompanied by covered code(s) under the applicable Coverage Policy will be denied as not covered. Coverage Policies relate exclusively to the administration of health benefit plans. Coverage Policies are not recommendations for treatment and should never be used as treatment guidelines. In certain markets, delegated vendor guidelines may be used to support medical necessity and other coverage determinations.

Overview

This coverage policy addresses the clinical indications for laboratory testing for transplantation rejection.

Coverage Policy

Gene expression profile (i.e., AlloMap®) is considered medically necessary when ALL of the following criteria are met:

- age 15 years or older
- two months to five years post-heart transplantation
- performed in lieu of routinely scheduled endomyocardial biopsies
- result will be used to determine the need for subsequent endomyocardial biopsy to clarify rejection status
- heart allograft function is stable as demonstrated by ALL of the following:
 - absence of signs or symptoms of congestive heart failure
 - current echocardiogram with left ventricular ejection fraction (LVEF) \geq 45%
 - absence of severe cardiac allograft vasculopathy (CAV)
- low probability of moderate or severe rejection as demonstrated by BOTH of the following:
 - no history or evidence of acute cellular rejection that required treatment
 - no history or evidence of antibody mediated rejection
- no history of gene expression profile (i.e., AlloMap) that did not correlate with endomyocardial biopsy

Gene expression profile (i.e., AlloMap®) for any other indication is considered not medically necessary.

All of the following are considered considered experimental, investigational or unproven for any transplantation indication:

- other gene expression profiling tests (e.g. HepatoTrack™, TruGraf, Molecular Microscope [MMDx])
- donor-derived cell-free DNA testing (e.g., AlloSure®, OmniGraf® Liver, Prospera™, VitaGraft™ Kidney 2.0)
- combined gene expression profiling and donor-derived cell-free DNA testing (i.e. HeartCare®)

Coding Information

Notes:

1. This list of codes may not be all-inclusive since the American Medical Association (AMA) and Centers for Medicare & Medicaid Services (CMS) code updates may occur more frequently than policy updates.

2. Deleted codes and codes which are not effective at the time the service is rendered may not be eligible for reimbursement.

Considered Medically Necessary when criteria in the applicable policy statements listed above are met:

CPT®* Codes	Description
81595	Cardiology (heart transplant), mRNA, gene expression profiling by real-time quantitative PCR of 20 genes (11 content and 9 housekeeping), utilizing subfraction of peripheral blood, algorithm reported as a rejection risk score.

Considered Experimental/Investigational/Unproven:

CPT®* Codes	Description
81558	Transplantation medicine (allograft rejection, kidney), mRNA, gene expression profiling by quantitative polymerase chain reaction (qPCR) of 139 genes, utilizing whole blood, algorithm reported as a binary categorization as transplant excellence, which indicates immune quiescence, or not transplant excellence, indicating subclinical rejection
0087U	Cardiology (heart transplant), mRNA gene expression profiling by microarray of 1283 genes, transplant biopsy tissue, allograft rejection and injury algorithm reported as a probability score
0088U	Transplantation medicine (kidney allograft rejection), microarray gene expression profiling of 1494 genes, utilizing transplant biopsy tissue, algorithm reported as a probability score for rejection
0493U	Transplantation medicine, quantification of donor-derived cell-free DNA (cfDNA) using next-generation sequencing, plasma, reported as percentage of donor-derived cell-free DNA
0540U	Transplantation medicine, quantification of donor- derived cell-free DNA using next-generation sequencing analysis of plasma, reported as percentage of donor-derived cell-free DNA to determine probability of rejection
0544U	Nephrology (transplant monitoring), 48 variants by digital PCR, using cell-free DNA from plasma, donor-derived cell-free DNA, percentage reported as risk for rejection
0575U	Transplantation medicine (liver allograft rejection), miRNA gene expression profiling by RT-PCR of 4 genes (miR-122, miR-885, miR-23a housekeeping, spike-in control), serum, algorithm reported as risk of liver allograft rejection
0576U	Transplantation medicine (liver allograft rejection), quantitative donor-derived cell-free DNA (cfDNA) by whole genome next- generation sequencing, plasma and mRNA gene expression profiling by multiplex real-time PCR of 56 genes, whole blood, combined algorithm reported as a rejection risk score

***Current Procedural Terminology (CPT®) ©2024 American Medical Association: Chicago, IL.**

General Background

Organ transplant recipients are at risk for allograft rejection, even with modern immunosuppressive therapies. Traditionally, diagnosis of allograft rejection has relied on nonspecific biochemical markers and histologic examination of the grafted tissue. Since this

requires an invasive tissue biopsy, there is great interest among those in the field of transplantation medicine to develop a noninvasive method of detecting organ transplant rejection (Verhoeven, et al., 2018). Gene testing is being investigated for the detection of post transplantation rejections following various types of transplants including heart, kidney, lung, and liver.

Based on U.S. Organ Procurement and Transplantation Network data (2025):

- In 2025 (as of Oct 26, 2025), 20,695 kidney transplants, 3,461 heart transplants, 9,227 liver transplants, and 64 intestine transplants were completed in the United States.
- 93,410 people were on the waiting list for a kidney transplant, 3,885 for a heart, 9,094 patients for a liver and 165 patients for an intestinal transplant.

Gene Expression Profile (GEP) Tests

AlloMap®: AlloMap® (CareDx®, South San Francisco, CA) has evolved into an established alternative to endomyocardial biopsy (EMB) in a defined subgroup of heart transplant recipients. The blood test measures gene expression by quantifying the gene-specific messenger RNA (mRNA) that is present in the sample. The expression level of 11 informative genes and nine normalization genes is measured using quantitative real-time polymerase chain reaction (qRT-PCR). The results are reported as an integer ranging from 0–40, and the lower the score the less the likelihood that the patient will experience rejection (i.e., AlloMap detects a low risk of rejection). It is proposed that circulating peripheral blood mononuclear cells may be indicative of rejection earlier than changes seen at local sites.

AlloMap may be used to help identify patients, age 15 years or older, who are between two months and five years post-heart transplantation, have a stable heart allograft function, and are at low risk of moderate or severe rejection, have no history or evidence of acute cellular or antibody mediated rejection and therefore, may not need to undergo endomyocardial biopsies. The test is recommended for use in conjunction with standard clinical evaluation and assessment (e.g., history and physical, echocardiography, endomyocardial biopsy) of graft function.

Stable heart allograft function is determined by the absence of congestive heart failure, left ventricular ejection fraction (LVEF) $\geq 45\%$ and absence of severe cardiac allograft vasculopathy (CAV). Symptoms of congestive heart failure include exertional dyspnea, orthopnea, paroxysmal nocturnal dyspnea, edema, syncope, palpitation and/or arrhythmias. LVEF measures the amount of blood being pumped out of the left ventricle of the heart. A LVEF below 45 may be evidence of CHF or cardiomyopathy. Rejection can be associated with hemodynamic compromise as indicated by a decrease in the LVEF. LVEF can be assessed by echocardiography, cardiac catheterization, MRI, multiple gated acquisition (MUGA) scan, or ventriculography (Pham, et al., 2010; Yamani and Taylor, 2010).

Cardiac allograft vasculopathy (CAV) is a disease that affects the entire network of coronary arteries in a transplanted heart. It is characterized by diffuse, concentric narrowing caused by thickening of the inner layer (intimal hyperplasia) in the larger epicardial arteries, as well as disease in the middle layer (medial) of the smaller microvascular vessels. This results in progressive luminal stenosis throughout the graft. Cardiac allograft vasculopathy (CAV) is also called transplant coronary artery disease or cardiac transplant vasculopathy and is one of the top three causes of death in heart transplant patients after the first year (Eisen, 2025). Invasive coronary angiography or intravascular ultrasound (IVUS) is used to assess CAV. CAV may be diagnosed when there is stenosis of 50% within any major epicardial coronary vessel or branches on angiography, severe diffuse or distal vessel tapering on angiography, maximal intimal thickness ≥ 0.5 mm in any major epicardial coronary vessel at the time of intravascular ultrasound, evidence of recent ischemic injury on biopsy, graft dysfunction and/or an epicardial

stenosis of < 50% that does not respond to anti-rejection therapy. Clinical manifestation of CAV may be silent, or occur as acute myocardial infarction, congestive heart failure, arrhythmias, and/or wall motion abnormalities (Eisen, 2025; Pham, et al., 2010; Yamani and Taylor, 2010; Schmauss and Weis, 2008).

The 2010 International Society for Heart and Lung Transplantation standardized nomenclature for cardiac allograft vasculopathy (CAV) (Mehra, et al., 2010) is as follows:

- ISHLT CAV₀ (Not significant): No detectable angiographic lesion
- ISHLT CAV₁ (Mild): Angiographic left main (LM) <50%, or primary vessel with maximum lesion of <70%, or any branch stenosis <70% (including diffuse narrowing) without allograft dysfunction
- ISHLT CAV₂ (Moderate): Angiographic LM <50%; a single primary vessel ≥70%, or isolated branch stenosis ≥70% in branches of two systems, without allograft dysfunction
- ISHLT CAV₃ (Severe): Angiographic LM ≥50%, or two or more primary vessels ≥70% stenosis, or isolated branch stenosis ≥70% in all three systems; or ISHLT CAV₁ or CAV₂ with allograft dysfunction (defined as LVEF ≤45% usually in the presence of regional wall motion abnormalities) or evidence of significant restrictive physiology (which is common but not specific; see text for definitions)

Definitions:

- Primary Vessel denotes the proximal and middle 33% of the left anterior descending artery, the left circumflex, the ramus and the dominant or co-dominant right coronary artery with the posterior descending and posterolateral branches
- Secondary Branch Vessel includes the distal 33% of the primary vessels or any segment within a large septal perforator, diagonals and obtuse marginal branches or any portion of a non-dominant right coronary artery
- Restrictive cardiac allograft physiology is defined as symptomatic heart failure with echocardiographic E to A velocity ratio >2 (>1.5 in children), shortened isovolumetric relaxation time (<60 msec), shortened deceleration time (<150 msec), or restrictive hemodynamic values (Right Atrial Pressure >12mmHg, Pulmonary Capillary Wedge Pressure >25 mmHg, Cardiac Index <2 l/min/m²)”

Pham, et al. (2010) identified severe CAV as “> 50% left main stenosis; ≥ 50% stenosis in ≥ 2 primary vessels (proximal 1/3 or middle 1/3 of the left anterior descending or left circumflex, right coronary artery to takeoff of posterior descending artery in right-dominant coronary circulations) or isolated branch stenosis of > 50% in all three systems (diagonal branches, obtuse marginal branches, distal 1/3 of left anterior descending or left circumflex, posterior descending artery, posterior lateral branch, and right coronary artery to takeoff of posterior descending artery in non-dominant systems)”.

Acute cellular rejection (ACR), or cell-mediated rejection, is the clinical syndrome that occurs as the result of an alloimmune response against a transplanted organ and can be caused by either a cellular or humoral response. ACR is the most common form of rejection and occurs at least once in approximately 50% of heart transplant recipients during the first year. Patients with a low probability of moderate or severe acute cellular rejection have a histological International Society of Heart and Lung Transplantation (ISHLT) grade 0 or grade 1. Typically, these grades are not treated for rejection. Grades 2R (moderate) and 3R (severe) are indicative of rejection and treated per institutional protocol. The two key elements of acute cellular rejection are the presence of lymphocytes and myocyte injury. Low probability of moderate or severe acute cellular rejection has been defined as patients who do not have treatable rejection on two consecutive biopsies over a period of 3–9 months. Treated rejection includes the administration of anti-rejection therapy (e.g., steroids, antibody therapy) (Eisen and Khush, 2024; Velleca, 2023; Acker and Jessup, 2011; Pham, et al., 2010; Stewart, et al., 2005).

Antibody mediated rejection (AMR), or humoral rejection, is initiated by antibodies rather than by T cells. AMR is manifested as graft dysfunction or hemodynamic compromise (e.g., shock, hypotension, decreased cardiac output, and/or a rise in pulmonary capillary wedge pressure) in the absence of cellular rejection on biopsy. The diagnosis is based on histologic findings indicative of acute myocardial capillary injury. Per Pham et al. (2010), AMR may be associated with hemodynamic compromise including: "LVEF \leq 30% or at least 25% below baseline, cardiac index < 2 L/min/m² or administration of inotropic agents to support circulation" (Berry, et al., 2013; Acker and Jessup, 2011; Pham, et al., 2010).

AlloMap has been proposed as an alternative to EMB in patients for whom biopsy is contraindicated or cannot be performed. One complication of biopsy is tricuspid regurgitation caused by repeated passing of the biopptome across the tricuspid valve into the right ventricular to obtain tissue specimens. Repeated biopsies may further damage the valve and increase regurgitation (Strecker, et al., 2013; Badiwala and Rao, 2007). Other contraindications for EMB include profound hemodynamic compromise, coagulopathy, and mechanical tricuspid prosthesis (Bennett and Tang, 2013). Lack of adequate vascular access, malignant arrhythmic events (e.g., unstable ventricular arrhythmias) or intracavitary mass or thrombus may also be contraindications to EMB. Clinical trials investigating AlloMap in this subset of patients are lacking.

AlloMap has not been validated for use in patients who demonstrate antibody-mediated rejection or noncellular rejection accompanied by hemodynamic compromise, pregnant women, patients who have recently (i.e., less than 30 days) received a blood transfusion, patients recently (i.e., less than 20 days) treated with high-dose steroids, patients recently treated for rejection, or patients who are being treated with ≥ 20 milligrams per day of prednisone or equivalent (Caideiras, et al., 2007; Mehra and Uber, 2007).

Moayed et al. (2019) reported on gene expression profiling (GEP) and racial disparities in outcomes after heart transplantation from the Outcomes AlloMap Registry. The data revealed that African Americans have lower survival rates after heart transplantation than Caucasians. There were 933 eligible recipients with 737 (79%) Caucasian and 196 (21%) African Americans. Compared to Caucasian recipients, African Americans were younger (age 55 vs 59 years, $p < 0.001$), had higher rates of non-ischemic cardiomyopathy (68% vs 50%, $p < 0.001$), lower rates of ischemic cardiomyopathy (27.0% vs 39.5%, $p < 0.001$), more likely sensitized with + 10% panel reactive antibody (PRA) (15.8% vs 9.1%, $p = 0.009$), and less likely to have primary cytomegalovirus serology (CMV) mismatch (14.3% vs 27.3%, $p < 0.001$). It was noted that African Americans were more likely than Caucasians to have eight or more human leukocyte antigen (HLA) mismatches (46.9% vs. 35.7%, $p = 0.013$). This is significant because there is increasing evidence that recipients with a higher number of mismatches have worse outcomes. The three-year survival rate was significantly lower in African Americans at 83.5% vs Caucasians at 97.5% ($p = 0.006$). There were no significant differences in the primary composite endpoint of left ventricular dysfunction, rejection with hemodynamic compromise, retransplantation, and mortality between races ($p = 0.36$). Higher tacrolimus levels were associated with decreased mortality in African American recipients ($p = 0.009$). On an individual gene level, it was found that MARCH8 gene expression was significantly correlated with mortality in Caucasians and elevated levels of FLT3 gene expression were associated with an increased risk in African Americans. GEP scores were similar for both racial groups until 24 months with higher scores in African American recipients after that. The disparity in survival based on race is confirmed by this registry data, but the reasons remain unclear.

U. S. Food and Drug Administration (FDA): In 2008, XDx Laboratories received 510(k) Class II approval for AlloMap Molecular Expression Testing "to aid in the identification of heart transplant recipients with stable allograft function who have a low probability of moderate/severe acute cellular rejection (ACR) at the time of testing in conjunction with standard clinical assessment".

AlloMap is indicated for use in heart transplant recipients who are 15 years of age or older and at least 2 months (≥ 55 days) post-transplantation (FDA, Nov 2008).

Device or Product	Identifier	Manufacturer	Decision Date
AlloMap Heart Molecular Expression Testing	K221640	CareDx, Inc.	9/13/2023
AlloMap [®] Molecular Expression Testing	K073482	CareDx, Inc.	8/8/2008

*FDA product codes: OJQ

Note: Coverage decisions are not based solely on FDA approval. Device or product names are provided for example purposes only. Their inclusion does not indicate endorsement or preference for any specific brand or model. This list is not intended to reflect all available products or technologies.

Literature Review: Clinical trials have reported that AlloMap was effective in identifying those patients with a low probability of rejection following heart transplantation at greater than six months following transplantation. Crespo-Leiro et al. (2016) conducted the Cardiac Allograft Rejection Gene Expression Observational II Study (CARGO II) to further clinically validate the GEP test performance using an independent study population. AlloMap testing was performed during post-transplant surveillance. Using a cutoff GEP score of 34 at six months post-transplantation, 95.5% (381/399) of GEP tests were true negatives, 4.5% (18/399) were false negatives, 10.2% (6/59) were true positives, and 89.8% (53/59) were false positives. Based on 938 paired biopsies, the GEP test score receiver operating characteristic curve (AUC-ROC) for distinguishing $\geq 3A$ rejection was 0.69 for 6 months post-transplantation. Depending on the chosen threshold score, the NPV and PPV ranged from 98.1%–100% and 2.0%–4.7%, respectively. In the CARGO (Cardiac Allograft Rejection Gene Expression Observational) study, Deng, et al. (2006) reported that the gene test significantly distinguished the absence of moderate/severe acute rejection from quiescence ($p=0.0018$). At a threshold of 30, the test predicted that patients with low molecular scores at or after one year following transplant were at low risk of current moderate/severe rejection (negative predictive value $>99\%$). Using CARGO samples ($n=127$), Mehra et al. (2008) reported that the use of AlloMap allowed for the identification and separation of patients into low-, intermediate-, and high-risk groups. Baseline scores were significantly higher for those who went on to reject, remained high during an episode of rejection, and dropped post-treatment for rejection ($p<0.01$). The Pham et al. randomized controlled trial (2010a; 2010b) (Invasive Monitoring Attenuation through Gene Expression [IMAGE]) compared outcomes of monitoring for rejection following heart transplantation using AlloMap gene-testing ($n=297$) compared to routine endomyocardial biopsy ($n=305$). Outcomes reported that this selected subpopulation of patients more than six months post transplantation were not at an increased risk of serious adverse outcomes and significantly fewer biopsies ($p=0.001$) were performed.

AlloMap Testing at Two to Six Months Following Transplantation: A prospective, observational, multi-center study, the Cardiac Allograft Rejection Gene Expression Observational II Study (CARGO II) (Crespo-Leiro, et al., 2016), was conducted to validate the clinical performance of AlloMap in an independent population. The performance of AlloMap was evaluated in two post-transplantation windows, ≥ 2 –6 months and > 6 months. Included samples were obtained at least 55 days post-transplantation, 30 days after transfusion of blood products, 21 days after administration of ≥ 20 mg/day of prednisone; and 60 days after treating a prior rejection. The main metric for validating the GEP test performance was the receiver operating characteristic curve (AUC-ROC). Values for the AUC-ROC range from 0.5 (uninformative) to 1.0 (perfect discrimination). All samples that had local biopsy grades of 1B, 2, and $\geq 3A$ were sent to an independent central pathology for grading. If more than one grade 0 or grade 1A local sample was obtained on a subject, one random sample was sent to central pathology. The ≥ 2 –6 months

post-transplantation specimens with histopathology slides selected for independent central panel pathology rejection grading included 328 patients with 480 GEP test scores. The mean receiver AUC-ROC for the GEP test scores in the ≥ 2 –6 months post-transplantation period was 0.70 with a 95% confidence interval from 0.67 to 0.73. The mean AUC-ROC for the GEP test scores for the > 6 months post-transplantation period was 0.69 with 95% confidence interval from 0.66 to 0.72. A GEP test score of ≥ 34 corresponded to histology-based grade $\geq 3A$ (2R) rejection with a PPV of 4.0% at months ≥ 2 –6 post-transplantation vs. 4.3% at > 6 months post-transplantation. The negative predictive values (NPVs) were 98.4% at months ≥ 2 –6 post-transplantation and 98.3% at > 6 months post-transplantation. The rates of rejection were similar in the early post-transplantation period (3.2% for months ≥ 2 to 6) compared with the > 6 post-transplantation population (3.2%). However, it was noted by the authors that the biopsies performed after month six post-transplantation may have been performed more often due to clinical suspicion of rejection, whereas biopsies performed prior to six months post-transplantation were probably conducted for surveillance of asymptomatic patients. There were several limitations of the study discussed by the authors including: 1) the study excluded GEP data and biopsies from visits earlier than 55 days post-transplantation; therefore, GEP performance is not known in this population; 2) patients treated for rejection within the prior 30 days were excluded which may have led to an underestimation of the incidence of rejection; 3) not all GEP tests had paired biopsy results; 4) because of the very small number and proportion of clinically suspected, biopsy confirmed, rejections of grade $\geq 3A$ (2R), rejections discovered during routine surveillance and rejections confirmed with biopsies performed for clinical suspicion were pooled together in the analyses; and 5) the central pathology feature of this study did not reflect practical clinical care.

Kobashigawa et al. (2015) conducted a single center randomized controlled trial (n=60), the Early Invasive Monitoring Attenuation through Gene Expression (EIMAGE) study, to evaluate the safety and efficacy of gene expression profiling (GEP; AlloMap) (n=30), compared to endomyocardial biopsy (n=30), in monitoring heart rejection from day 55 and onward post-transplantation. Patients were ≥ 18 years of age, had undergone cardiac transplantation within the past 2–6 months (55–185 days), and in a clinically stable condition with an echocardiographic LVEF of $\geq 50\%$. Exclusion criteria included any clinical signs of declining graft function defined by symptoms of congestive heart failure at the first study surveillance visit, rejection therapy for biopsy-proven ISHLT Grade 2R (3A) or higher during the preceding two months, previous or current evidence of antibody-mediated rejection (AMR) defined according to the ISHLT 2004 guidelines, presence of donor-specific antibodies (DSAs), high corticosteroid doses of > 20 mg at the time of the first study visit, receiving hematopoietic growth factors or blood transfusion during the previous 30 days, and pregnancy. A positive GEP was considered ≥ 30 at 2–6 months and ≥ 34 after six months and prompted a biopsy. The primary outcome was a composite of first occurrence of death/retransplant, rejection with hemodynamic compromise, or allograft dysfunction because of other causes ≤ 18 months post-transplant. There was no statistically significant difference in the primary outcome between the groups at 18 months follow-up. Intravascular ultrasound (IVUS) results showed no significant differences between the groups in the average change in maximal intimal thickness (p=0.944) or plaque burden in the first year. LVEF results at 18 months were not significantly different (p=0.522), nor were the cardiac index score (p=0.413) or other cardiac hemodynamics. The number of biopsy-proven ISHLT $\geq 2R$ (3A) rejection episodes within the first 18 months post-transplant was not significantly different between the groups (p=0.31). At 18 months, 42 biopsies were performed in the GEP group, compared with 253 in the EMB group. There was no significant difference in prednisone weaning or average prednisone dosage between the two groups. Satisfaction of method rejection surveillance was statistically significant in the GEP group (p=0.003) but there were no significant differences in the SF-12 mental health and physical-health summary scores. The results of this pilot study should be viewed with caution due to the low risk, small patient population included in the trial. The authors noted that firm conclusions could not be made about noninferiority of GEP versus EMB, further large-scale clinical trials to confirm these findings are required; and the study

was limited by the few primary end points experienced as only eight patients reached the composite primary end point.

Other AlloMap indications: Studies have also been conducted to determine if AlloMap could identify different forms of rejection, identify patients who may develop Grade \geq 3A rejection, what impact coronary allograft vasculopathy (CAV) and blood transfusion would have on AlloMap scoring. AlloMap has also been analyzed for the ability to predict clinical events \geq 315 days following transplantation and as a predictor for long-term survival after heart transplant.

In a retrospective, single center study, Fujita et al. (2017) investigated AlloMap as a predictor of long-term survival following heart transplantation. Data on 46 patients from the CARGO II trial were analyzed. Patients who had survived for at least one year and in whom AlloMap scores were available at six, nine, 12 and 18 months after transplant were included. The primary outcome measure was long term all-cause mortality. Survival data were gathered by review of medical records and contact with the patient/family. Six to nine months following transplantation 23 patients showed a decreased AlloMap score and 23 showed an increased score. The scores at individual time points (6, 9, 12, 18 months) did not correlate with long-term survival. Patients with an increased score from six to nine months after transplant compared to a decreased score showed a significant elevation in all-cause mortality ($p=0.005$). It was also noted that long-term mortality may have been associated with side effects or inadequate immunosuppression. Limitations of the study are the retrospective design, small patient populations and heterogeneity in baseline characteristics. Prospective studies with larger patient populations are required to validate the predictive value of AlloMap in identifying patients at high risk for death during long-term follow-up.

Crespo-Leiro et al. (2015) conducted a retrospective analysis ($n=91$) of an independent patient population from the Cardiac Allograft Rejection Gene Expression Observational (CARGO) II study to determine the prognostic utility of within patient variability of GEP scores in predicting future, significant clinical events. A second objective was to determine the negative predictive value (NPV) and the positive predictive value (PPV) of GEP score variability in predicting future significant clinical events. The GEP score variability was defined as the standard deviation of four GEP scores collected \geq 315 days post-transplant. Analyzed clinical outcome data included CARGO II patients who had four AlloMap scores preceding a first clinical event (event group) or had four sequential GEP scores without a subsequent event (control group). Out of the 737 patients in the CARGO II study, 55 patients who did not have events and had at least a three year follow-up were chosen as controls and 36 patients who experienced at least one predefined cardiac event were included in the event group. There was no statistically significant difference between the groups in number of days between the first and the last AlloMap test. The estimated prevalence of events was 17%. Events occurred at a median of 391 days after the final GEP Test. The NPV increased from 87.4% at a score variability of 1.0 to 97% at a score variability of 0.6. The PPVs for the same score decreased from 26% at a score variability of 1.0 to 23.3% for a GEP score variability of 0.6. For a GEP score variability cutoff of 1.5, the estimated PPV was 35.4%. In the event group 58% of the patients died, 31% experienced graft failure and 11% underwent cardiac re-transplantation. The results of the study proposed that the AlloMap score variability may be useful in estimating the probability of future events of death, re-transplantation or graft failure in heart transplant recipients tested with AlloMap \geq 315 days post-transplantation. The NPV \geq 97% indicated that clinical utility of AlloMap score variability may help identify patients at a low risk for future clinical events. Given, the limitations of the study, the results should be viewed with caution. Limitations of the study include the retrospective study design, small patient population, potential selection bias of patients, imbalance of number of patients per group and heterogeneity of clinical conditions.

Bernstein et al. (2007) conducted a subanalysis of the CARGO study to determine if gene expression (GE) (i.e., AlloMap genetic testing) could distinguish different forms of mild heart transplant rejection. Inclusion criteria were met by 265 of the 737 adult and pediatric CARGO patients. Reinterpretation of the tissue identified: 176 grade 0 biopsies, 17 grade 1As, 12 grade 1Bs, 21 grade 2, and 24 grade 3As. The mean GE scores differentiated moderate-to-severe rejection (grades $\geq 3A$) (32 ± 0.9) from grades 0 (25.3 ± 0.5), 1A (23.8 ± 2.1) and 2 (26.9 ± 1.5) ($p < 0.00001$, $p < 0.001$ and $p < 0.01$, respectively). The mean GE score for grade 1B was indistinguishable from that for grades $\geq 3A$, (29.8 ± 2.0 vs. 32.0 ± 0.9) ($p = 0.25$). Based on a calculation of the fold-difference of each gene, grade 1B was identified as a subgroup of rejection with a peripheral gene expression profile that more closely resembled moderate-to-severe rejection. The study also analyzed whether or not the time from transplantation influenced the GE scores compared to the grades. For the two- to six-month period following transplantation, the mean GE score for grade $\geq 3A$ (30.8 ± 1.4) was not significantly different from that for grade 1B (28.5 ± 3.9) ($p = 0.49$). The mean GE scores differentiated grades 0, 1A, and 2 from grades $\geq 3A$. EMBs obtained more than six months following transplantation indicated grades $\geq 3A$ demonstrated mean GE scores similar to grade 1B scores ($p = 0.19$). Mean GE scores for grades 0, 1A, and 2 were significantly lower than for grades $\geq 3A$ scores.

Mehra et al. (2007) also conducted a subanalysis of cardiac allograft recipients ($n = 104$) from the CARGO study to determine if the AlloMap test could distinguish between rejection-free stable patients and patients who develop Grade $\geq 3A$ rejection within 12 weeks following transplantation. In addition, the study characterized the associations with rejection within 180 days of transplantation, identified individual classifier genes associated with the risk of future rejection and explored the pathways and functions of the genes. Patients with grades 0 or 1A at baseline and free of \geq grade 2 rejection for at least the first 12 weeks post-transplantation were designated as the matched control group ($n = 65$). The rejection group included 39 patients, clinically stable at baseline, who experienced an episode of grade $\geq 3A$ within 12 weeks following sample collection. Data for the study was analyzed from blood samples and EMB obtained during the same visit. Analysis of the data demonstrated a significant difference in the mean GE score of 27.4 ± 6.3 for the study group and 23.9 ± 7.1 for the control group ($p = 0.01$). The study also analyzed a subgroup of these patients who were ≤ 180 days post-transplant and reported a significant difference in the mean GE score of 28.4 ± 4.9 for the study group ($n = 28$) and 22.4 ± 7.5 for the control group ($n = 46$) ($p < 0.001$). To explore the molecular pathways associated with steroid sensitivity and T-cell activation, the expression levels of 33 additional genes were measured, and the data demonstrated that "transcriptional signals of genes regulated by corticosteroids or involved in T-cell activation in peripheral blood of heart transplant recipients are associated with the presence or absence of future clinically relevant rejection." The authors stated that the data from this study "must be interpreted with care and in the context of the case-control study in which they were derived." They further explained that case-control studies include "inherent spectrum bias, preventing generalization," and noted that milder rejection grades (i.e., 1B and 2) were not addressed.

In 2007, Yamani et al. conducted two retrospective reviews. The first study (Apr 2007a) included 69 patients and evaluated the impact of transplant coronary allograft vasculopathy (CAV) on AlloMap gene expression analysis. Evidence of CAV within 4.3 ± 3 months of AlloMap testing was demonstrated in 20 patients by coronary angiography. The control group had a mean AlloMap score of 26.1 ± 6.5 compared to $> 32.2 \pm 3.9$ in the CAV group ($p < 0.001$). Fifteen control patients and 14 CAV patients had an AlloMap score of greater than 30 ($p = 0.0026$). CAV was associated with a significantly increased AlloMap score in the absence of significant rejection ($p = 0.0002$). The second review (2007b) investigated the impact of early post-transfusion ischemic injury on subsequent AlloMap testing from data retrieved from a transplant database ($n = 67$). The subjects were evaluated at a mean 34 ± 20 months following heart transplantation. Compared to the control group, the injury group demonstrated worse five-year freedom from vasculopathy,

lower left ventricular ejection fraction (LVEF), and higher percentage of AlloMap scores. The presence of ischemic injury was associated with a significant increase in AlloMap scores ($p < 0.0001$).

HepatoTrack™: The HepatoTrack (LuminoDx Inc., San Diego, CA) assay analyzes cell-free microRNAs (miRNAs) in blood samples and has been proposed to predict acute cellular rejection (ACR) in liver transplant patients.

Literature Review: There is insufficient evidence to support the accuracy and clinical utility of HepatoTrack.

TruGraf®: The TruGraf® assay (Eurofins Transplant Genomics, Framingham, MA) is a DNA microarray-based gene expression blood test proposed for use in renal transplant recipients. It is proposed as an alternative to surveillance biopsies to rule out subclinical rejection in patients with stable graft function (Anglicheau, et al., 2024; Marsh, et al., 2019). To date, the literature reports that TruGraf has only been tested on patients whose transplant graft was known to be doing well. The test has not been tested on patients whose graft was starting to show signs of rejection. How the test performs in this subpopulation needs to be studied before TruGraf can be recommended for use.

Literature Review: In a multicenter retrospective study, Marsh et al. (2019) conducted simultaneous blood tests and clinical assessments in 192 patients from seven transplant centers to assess the clinical utility of the TruGraf DNA microarray-based gene expression blood test in the serial assessment of kidney transplant recipients with stable renal function. The accuracy or concordance between TruGraf result and clinical and/or histologic assessment was 74% (142/192). The test was accurate in 93% (116/125) of patients identified as Transplant eXcellence (TX: stable serum creatinine, normal biopsy results, indicative of immune quiescence). The negative predictive value for TruGraf was 90%, with a sensitivity of 74% and specificity of 73%. Results did not differ significantly in patients with a biopsy-confirmed diagnosis vs those without a biopsy.

Molecular Microscope (MMDx) Kidney and Heart: The molecular microscope diagnostic system (MMDx) (One Lambda, Inc [Thermo Fisher Scientific, Inc], West Hills, CA) is a microarray-based system. It utilizes transplant biopsy tissue and analyzes it for messenger RNA (mRNA) expression patterns to predict the diagnosis of acute T cell-mediated rejection (TCMR) or antibody-mediated rejection (ABMR) (Anglicheau, et al., 2024). According to the manufacturer's website, the laboratory developed tests are used for clinical purposes by the CLIA-certified laboratory performing the test. These tests are not cleared or approved for clinical use by the FDA as an in vitro diagnostic test (Thermo Fisher Scientific, Inc, 2025).

Literature Review: There is insufficient evidence to support the accuracy and clinical utility of the molecular microscope diagnostic system. The literature is primarily in the form of an editorial, reviews and test accuracy analysis.

Donor-Derived Cell-Free DNA (dd-cfDNA) Tests: It is hypothesized that transplant patients experiencing organ injury associated with acute rejection will have higher levels of donor-derived cell free DNA (dd-cfDNA) which is thought to be due to cell-free DNA being an indicator of dying cells. In recent years, dd-cfDNA tests have become clinically available to quantify the amount of dd-cfDNA in kidney and heart transplant recipients.

There are multiple techniques for measuring dd-cfDNA with each test measuring a different number of single-nucleotide polymorphisms (SNPs). These tests are intended to assess the probability of allograft rejection in a particular patient. The technology does not rely on previous

genotyping of either the patient or the donor, which is a benefit over previous methods that have been used to measure dd-cfDNA. Additionally, these tests do not require invasive tissue biopsy, which is necessary for the standard methods of histopathological interpretation that are used to diagnose allograft rejection. However, biopsy is still necessary to confirm and establish the type of active rejection in affected patients (Jordan et al., 2018). It has been proposed that these tests be used for serial monitoring in order to detect new onset injury or rejection prior to clinical symptoms, however the optimal time interval has yet to be established (Bloom et al., 2017). The use of dd-cfDNA to evaluate transplant rejection is not a new development in the field of transplant medicine, however the clinical utility of this technology has yet to be established.

AlloSure®: Allosure® (CareDx®, South San Francisco, CA) is a targeted next-generation sequencing test that evaluates single nucleotide polymorphisms (SNP) in cell-free DNA samples. AlloSure Kidney is proposed for use in renal transplant patients who are 18 years or older and a minimum of 14 days post-transplant. AlloSure Heart is proposed for use in conjunction with AlloMap in heart transplant patients who are 15 years or older and at least 55 days post-transplant (CareDx, 2020).

Literature Review-Heart: There is insufficient evidence to support the accuracy and clinical utility of donor-derived cell free DNA for assessing and monitoring the probability of allograft rejection in heart transplant patients. Studies are primarily in the form of an author manuscript, a conference abstract, a proof-of-concept study, retrospective reviews, and registry data (Henricksen et al., 2023; Kamath, et al., 2022; Khush, et al., 2019; Crespo-Leiro et al., 2015; De Vlaminck, et al., 2014; Hidestrand, et al., 2014).

Literature Review-Kidney: Aubert et al. (2024) conducted a multinational, observational cohort study to evaluate donor-derived cell-free DNA (dd-cfDNA) as a biomarker for kidney allograft rejection. A total of 2,882 transplant recipients were enrolled across 14 centers; the derivation cohort included 1,134 adults (1,415 biopsies), and the external validation cohort comprised 1,748 patients. The intervention was dd-cfDNA measurement (AlloSure/AlloSeq cfDNA assays) added to standard monitoring (eGFR, proteinuria, anti-HLA DSA), with biopsy-proven rejection as the comparator. Inclusion criteria were kidney transplant recipients ≥ 18 years; exclusions included combined organ transplants, pregnancy, monozygotic twin donors, and prior bone marrow transplant. Outcomes were biopsy-confirmed antibody-mediated rejection (AMR), T cell-mediated rejection (TCMR), or mixed rejection, assessed at protocol (three months, one year) and for-cause biopsies. Median follow-up was approximately one year. Donor-derived cell-free DNA levels strongly correlated with allograft rejection, including antibody-mediated rejection ($p < 0.0001$), T cell-mediated rejection ($p < 0.0001$) and mixed rejection ($p < 0.0001$). Limitations include observational design, lack of randomization, incomplete assessment of longitudinal dd-cfDNA trends, and absence of adherence data

Bu et al. (2022) conducted a multicenter, observational study (the ADMIRAL study: Assessing AlloSure Dd-cfDNA, Monitoring Insights of Renal Allografts with Longitudinal Surveillance) of 1,092 adult kidney transplant recipients who were monitored with donor-derived cell-free DNA (dd-cfDNA) testing over the course of three years. The objective of the study was to see how effective dd-cfDNA was in identifying allograft rejection, subclinical changes and to evaluate the relationship between dd-cfDNA measurements and nonimmune allograft injury. The study also sought to evaluate the relationship between elevated dd-cfDNA and predictors of long-term graft survival, including estimated glomerular filtration rate (eGFR) and formation of de novo donor-specific antibodies (dnDSAs). The plasma cfDNA fraction was measured using a targeted, single nucleotide polymorphism (SNP)-based assay. Pathological diagnosis was made according to the 2019 Banff Kidney Rejection Classification. To calculate the performance characteristics of the assay (sensitivity, specificity, negative predictive value, positive predictive value), the discriminatory power was considered at previously published thresholds of 0.5% and 1%. Patients were

categorized as high dd-cfDNA ($\geq 0.5\%$) versus low dd-cfDNA ($\leq 0.5\%$). Elevation of dd-cfDNA (0.5% or more) was significantly correlated with clinical and subclinical allograft rejection. Patients with antibody mediated rejection (ABMR) had higher levels of dd-cfDNA with clinical rejection (2.2% vs. 0.34%) and subclinical rejection (0.91% vs. 0.23%). Patients with T cell mediated rejection (TCMR) also had higher levels of dd-cfDNA with clinical rejection (1.30% vs. 0.34%) and subclinical rejection (0.52% vs. 0.23%). dd-cfDNA values of 0.5% or more were associated with a nearly three-fold increase in risk development of de novo donor-specific antibodies (hazard ratio 2.71) and were determined to be elevated a median of 91 days (interquartile range of 30-125 days) ahead of donor specific antibody identification. Persistently elevated dd-cfDNA (more than one result above the 0.5% threshold) predicted over a 25% decline in the estimated glomerular filtration rate over three years (hazard ratio 1.97). Author noted limitations of the study include the observational real-world design. Additional studies are needed to define how the information provided by dd-cfDNA can be used to guide clinical practice.

Zhang et al. (2020) conducted a prospective single center observational study to evaluate the diagnostic performance of donor-derived plasma cell-free DNA (cfDNA) in discriminating antibody-mediated rejection (ABMR) or de novo donor-specific antibodies (DSA) without histological lesions in kidney allograft recipients. The plasma cfDNA fraction was measured using a targeted, single nucleotide polymorphism (SNP)-based assay. Pathological diagnosis was made according to the 2015 Banff Kidney Rejection Classification. The area under the ROC curve (AUC-ROC) was determined using the bootstrapping method to estimate median and 95% confidence interval (95% CI). The sensitivity, specificity and Youden index, positive predictive value (PPV), and negative predictive value (NPV) were calculated for specific cfDNA fractions. The study included 37 consecutive patients who received kidney allografts, including 18 recipients in the ABMR group and 19 recipients in the stable allograft group (seven DSA-positive and 12 DSA-negative). All patients in the ABMR group were DSA positive and seven patients in the stable group were DSA positive but had no pathologically proven ABMR. The median donor-derived plasma cfDNA fraction was 2.4% (Q1 1.52% -Q3 3.70%) in the ABMR group, and was significantly higher than that of the stable group (0.65%, Q1 0.57% -Q3 0.97%; $P < 0.001$), but comparable with that of the DSA-positive patients in the stable allograft group ($P = 0.074$). The AUC-ROC of cfDNA was 0.90 (95% CI, 0.79-0.98). When a cfDNA threshold of 1% was chosen, it had a sensitivity of 88.9% and a specificity of 73.7%. The PPV was 76.2% and the NPV was 87.5%. The authors concluded that donor-derived plasma cfDNA fraction increased in kidney allograft recipients with ABMR. The study is limited by the small number of participants.

Bloom et al. (2017) conducted a prospective observational study: Circulating Donor-Derived Cell-Free DNA in Blood for Diagnosing Acute Rejection in Kidney Transplant Recipients (DART) study. Blood specimens were collected from 102 patients at scheduled intervals and at the time of clinically indicated biopsies. Plasma levels of donor-derived cell-free DNA (dd-cfDNA) were measured and correlated with allograft rejection status ascertained by histology in 107 biopsy specimens. The dd-cfDNA level discriminated between biopsy specimens showing any rejection (T cell-mediated rejection or antibody-mediated rejection [ABMR]) and controls (no rejection histologically), $P < 0.001$ (receiver operating characteristic area under the curve [AUC], 0.74; 95% confidence interval [95% CI], 0.61 to 0.86). Positive and negative predictive values for active rejection at a cutoff of 1.0% dd-cfDNA were 61% and 84%, respectively. The AUC for discriminating ABMR from samples without ABMR was 0.87 (95% CI, 0.75 to 0.97). Positive and negative predictive values for ABMR at a cutoff of 1.0% dd-cfDNA were 44% and 96%, respectively. Median dd-cfDNA was 2.9% (ABMR), 1.2% (T cell-mediated types \geq IB), 0.2% (T cell-mediated type IA), and 0.3% in controls ($P = 0.05$ for T cell-mediated rejection types \geq IB versus controls).

Jordan et al. (2018) conducted a study with a cohort from above DART study to assess the combined use of donor-derived cell-free DNA (dd-cfDNA) and Donor-specific antibodies (DSA)

testing to diagnose active antibody-mediated rejection (ABMR). Donor-derived cell-free DNA was assayed in 90 blood samples with paired DSA and clinically indicated biopsies from 87 kidney transplant patients. Sixteen cases met criteria for active ABMR. Performance characteristics of dd-cfDNA for diagnosis of active ABMR were determined for samples with prior or current positive DSA (DSA+, n = 33). The median level of dd-cfDNA (2.9%) in DSA+ patients with active ABMR was significantly higher than the median level (0.34%) in DSA+ patients without ABMR (P<0.001). The median level of dd-cfDNA in DSA- patients was 0.29%. The positive predictive value of dd-cfDNA (at 1%) to detect active ABMR in DSA+ patients was 81%, whereas the negative predictive value was 83%. The positive predictive value for DSA+ alone was 48%.

Bromberg et al. (2017) conducted an observational study of a cohort of the above DART study to establish biological variation and clinical reference intervals of dd-cfDNA in renal transplant recipients by using an analytically validated assay that has a CV of 6.8%. Venous blood was sampled at patient surveillance visits (typically at posttransplant months 1–4, 6, 9, and 12). Patients with stable renal allograft function spanning ≥ 3 serial visits were selected. AlloSure was used to measure dd-cfDNA in the plasma and computed the intraindividual CV (CVI) and interindividual CV (CVG), the index of individuality (II), and reference change value (RCV). The study included 93 patients with 61% men, 56% Caucasian, mean ages 49 years, and 63% were deceased donor kidney recipients. Of the 380 blood samples, the dd-cfDNA median value was 0.21% (interquartile range 0.12%–0.39%) and the 97.5th percentile was 1.20%. In 18 patients with an average of 4.1 tests, the CVI was 21%, CVG was 37%, II was 0.57, and RCV was 61%. The authors concluded that in a renal transplant recipient, a dd-cfDNA level above 1.2% is out of range and potentially abnormal. A serial increase of up to 61% in level of dd-cfDNA in a patient may be attributable to biological variation.

OmniGraf® Liver: OmniGraf Liver (Eurofins Transplant Genomics, LLC, Lenexa, KS) is a donor derived cell free DNA (dd-cfDNA) blood test proposed to detect allograft rejection in liver transplant recipients.

Literature Review: There is insufficient evidence to support the accuracy and clinical utility of OmniGraf Liver.

Prospera™: Prospera (Natera™, Austin, TX) is a suite of donor derived cell free DNA (dd-cfDNA) blood tests that are proposed to detect antibody-mediated rejection (AMR) and acute cellular rejection (ACR) rejection in kidney, heart or lung post-transplant recipients. According to the manufacturer's website, Prospera has not been cleared or approved by the FDA (Natera, 2025). The platform is run in a CLIA-certified, College of American Pathologists-accredited laboratory.

Literature Review-Heart: There is insufficient evidence to support the accuracy and clinical utility of donor-derived cell free DNA for assessing and monitoring the probability of allograft rejection in heart transplant patients. Studies are primarily in the form of an observational study (Deshpande et al., 2024).

Literature Review-Kidney: There is insufficient evidence to support the accuracy and clinical utility of donor-derived cell free DNA for assessing and monitoring the probability of allograft rejection in kidney transplant patients. Studies are primarily in the form of retrospective reviews (Bromberg et al., 2024; Kim et al., 2022).

Literature Review-Lung: There is insufficient evidence to support the accuracy and clinical utility of donor-derived cell free DNA for assessing and monitoring the probability of allograft rejection in lung transplant patients.

VitaGraft™ Kidney: VitaGraft™ Kidney (Oncocyte Corporation, Irvine, CA) is a blood-based test proposed to monitor transplant rejection that quantifies the concentration of donor-derived cell-free DNA following kidney transplantation. Beginning testing with VitaGraft Kidney requires a urine sample for the initial test to set the assay for the patient, all subsequent tests only require a blood sample. VitaGraft™ Kidney 2.0 evaluates eight additional single-nucleotide polymorphisms (SNPs) than the first version of VitaGraft Kidney. This test represents digital PCR, using cell-free DNA from plasma, donor-derived cell-free DNA, and gives a percentage reported as risk for rejection. VitaGraft is not indicated in patients who are: pregnant, less than two weeks post-transplant for subsequent testing, recipients of an allograft from an identical twin, bone marrow transplant recipients, recipients of an allogeneic stem cell transplant, recipients of a non-liver or non-kidney organ transplant, or recipients of multiple organ transplants. According to the manufacturer's website, "The VitaGraft Kidney Test has not been cleared or approved by the US Food and Drug Administration (FDA). Oncocyte's laboratory offering the VitaGraft Kidney Test is CAP-accredited and CLIA-certified" (Oncocyte Corporation, 2024).

Literature Review: There is insufficient evidence to support the accuracy and clinical utility of VitaGraft Kidney for the identification of post-transplant subclinical acute or acute rejection of a kidney.

Combined Gene Expression Profile and Donor-Derived Cell-Free DNA (dd-cfDNA) Tests

HeartCare®: HeartCare (CareDx®, South San Francisco, CA) combines AlloMap Heart gene expression profiling with AlloSure Heart dd-cfDNA. As noted above, the clinical utility of AlloSure Heart has yet to be established.

Khush et al. (2024) reported on a prospective, multicenter observational study (SHORE registry; NCT03695601) which evaluated 2,077 heart transplant recipients across 67 U.S. centers between 2018–2021 to assess dual noninvasive surveillance for acute cellular rejection (ACR) using gene-expression profiling (GEP; AlloMap) and donor-derived cell-free DNA (dd-cfDNA; AlloSure). Eligible patients were ≥15 years old, ≥55 days post-transplant, and had at least one paired GEP/dd-cfDNA test; pregnant women and multiorgan transplant recipients were excluded. Participants were categorized into four groups: GEP–/dd-cfDNA–, GEP+/dd-cfDNA–, GEP–/dd-cfDNA+, and GEP+/dd-cfDNA+. Primary outcomes included biopsy-proven ACR incidence, test performance (sensitivity, specificity, positive likelihood ratios [LR+]), and follow-up biopsy rates; secondary outcomes were graft function and survival. Median follow-up was 40 months. ACR incidence ranged from 1.5% (dual negative), 1.9% (GEP+/dd-cfDNA–), 4.3% (GEP–/dd-cfDNA+,), and 9.2% (dual positive). The sensitivity of dual positive testing with GEP/dd-cfDNA for ACR was 32.3% compared to 59.0% for GEP alone and 44.7% for dd-cfDNA alone. The specificity was 91.7% for dual-positive test results, 56.8% for GEP alone, and 84.6% for dd-cfDNA alone. Two-year survival was 94.9%, with graft dysfunction in 2.7%. Limitations include exclusion of antibody-mediated rejection from performance analyses, lack of centralized biopsy review, potential bias from variable timing of biopsies, and incomplete site verification early in the study.

Technology Assessments

BlueCross BlueShield Association (BCBSA) Technology Evaluation Center (TEC): BCBSA (2011) conducted a systematic review of the literature to determine if AlloMap testing improved health outcomes compared to other methods used for monitoring rejection following heart transplantation. Validation studies that only included patients with no rejection or class 3A rejection reported a sensitivity of 76–84% and specificity of 38–41% at a cutoff score of 20. Post hoc analyses of subgroups (n < 30 patients) who were > 6 months or > 12 months post-transplant with higher cutoff scores reported sensitivities of 71.4–80% and specificities of 77.8%–78.7%. Depending on the cutoff score used to denote a positive test, other studies reported a positive predictive value (PPV) typically < 7% and a negative predictive value (NPV) > 98%. According to

BCBSA, the data used for these values was not available and the results were not consistent with results from actual patient samples. One study reported a 7.8% PPV and a 100% NPV using a cutoff score of 34 but out of the total 243 samples, only five were rejection samples. The authors reported that higher AlloMap scores were associated with a greater likelihood of rejection in class 3A or higher patients but the “diagnostic characteristics of AlloMap testing were uncertain”, study methods were unclear, study samples were not completely described, number of cases of rejection were small and cutoff scores appeared “to be determined post hoc.” According to BCBSA, the “sensitivity of the test for detecting rejection is uncertain.” The available evidence was insufficient to permit conclusions regarding the effects of AlloMap on net health outcomes, or if the test is as beneficial as any established alternatives for monitoring heart transplant patients.

Professional Societies/Organizations

In their 2010 guidelines on the care of heart transplant (HT) recipients, the International Society for Heart and Lung Transplantation (ISHLT) stated, “gene expression profiling (AlloMap) can be used to rule out the presence of ACR [acute cardiac rejection] of grade 2R [i.e., an infiltrate plus the presence of multifocal myocyte damage] or greater in appropriate low-risk patients, between 6 months and 5 years after HT”. This recommendation is based on data from the CARGO and IMAGE clinical trials. The ISHLT guidelines were updated in 2023 to state “Gene Expression Profiling (GEP) (i.e., Allomap) of peripheral blood can be used in low-risk patients between 2 months and 5 years after HT to identify adult recipients who have low risk of current ACR to reduce the frequency of endomyocardial biopsy (EMB). Data in children does not allow a general recommendation of GEP as a routine tool at present.” The 2023 ISHLT guidelines for monitoring of rejection may include in addition to surveillance EMB several noninvasive rejection monitoring tests including Gene Expression Profiling (Allomap), donor specific antibody (DSA), brain natriuretic peptide (BNP), high sensitivity troponins, and donor-derived cell-free DNA (dd-cfDNA). The guidelines are limited in that dd-cfDNA is neither specifically recommended nor discouraged; the evaluation of the recommendation and evidence includes all seven actions that may be part of follow-up visits to monitor for rejection and screen for adverse events.

The evaluation of donor-derived cell free DNA has not yet been addressed by the Kidney Disease: Improving Global Outcomes (KDIGO) clinical practice guideline for care of kidney transplant recipients.

In 2023 (updated Oct 2024), the American Society of Transplant Surgeons (ASTS) released a statement on donor-derived cell-free DNA (dd-cf-DNA) which included the following:

- Suggested that clinicians consider measuring serial dd-cfDNA levels in kidney transplant recipients with stable renal allograft function to exclude the presence of subclinical antibody mediated rejection.
- Recommend that clinicians measure dd-cfDNA levels in kidney transplant recipients with acute allograft dysfunction to exclude the presence of rejection, particularly antibody-mediated rejection (ABMR).
- Recommended that dd-cfDNA may be utilized to rule out subclinical rejection in heart transplant recipients.
- Recommended that clinicians who choose to utilize dd-cfDNA in heart transplant recipients also utilize peripheral blood GEP as a non-invasive diagnostic tool to rule out acute cellular rejection in stable, low-risk, adult heart transplant recipients who are over 55 days status post heart transplantation.

Overall, there is no grade assigned for the level of evidentiary rigor or the strength of recommendations provided, and there is a lack of background, rationale, or literature cited for the recommended use in heart transplant recipients. The specific limitations identified in the ASTS statement include: the need for further work to enhance the detection of T-cell-mediated rejection (TCMR); the necessity for additional investigation to establish the optimal monitoring interval; and the requirement for further studies to clarify how the information provided by dd-cfDNA can

inform clinical practice and decisions regarding immunomodulation, infection management, treatment of all types of rejection, and the control or prevention of dnDSA formation.

Health Equity Considerations

Health equity is the highest level of health for all people; health inequity is the avoidable difference in health status or distribution of health resources due to the social conditions in which people are born, grow, live, work, and age.

Social determinants of health are the conditions in the environment that affect a wide range of health, functioning, and quality of life outcomes and risks. Examples include safe housing, transportation, and neighborhoods; racism, discrimination and violence; education, job opportunities and income; access to nutritious foods and physical activity opportunities; access to clean air and water; and language and literacy skills.

Medicare Coverage Determinations

	Contractor	Determination Name/Number	Revision Effective Date
NCD	National	No National Determination found	
LCD	CGS Administrators, LLC	MoIDX: Molecular Testing for Solid Organ Allograft Rejection (L38582)	7/10/2025
LCD	Noridian	MoIDX: Molecular Testing for Solid Organ Allograft Rejection (L38629)	7/10/2025
LCD	Noridian	MoIDX: Molecular Testing for Solid Organ Allograft Rejection (L38671)	7/10/2025
LCD	Wisconsin Physicians Service Insurance Corporation	MoIDX: Molecular Testing for Solid Organ Allograft Rejection (L38680)	7/10/2025
LCD	Palmetto GBA	MoIDX: Molecular Testing for Solid Organ Allograft Rejection (L38568)	7/10/2025

Note: Please review the current Medicare Policy for the most up-to-date information. (NCD = National Coverage Determination; LCD = Local Coverage Determination)

References

1. Acker MA, Jessup M. Ch 31 Surgical management of heart failure. In: Bonow: Braunwald's Heart Disease - A Textbook of Cardiovascular Medicine, 9th ed. Saunders, Philadelphia, 2011.
2. American Society of Transplant Surgeons (ASTS) Statement on donor-derived cell-free DNA (dd-cf-DNA). March 6, 2023, updated October 2024. Accessed on Nov 3, 2025. Available at URL address: <https://asts.org/advocacy/position-statements>
3. Anglicheau D, Malone A, Chon WJ. Kidney transplantation in adults: Investigational methods in the diagnosis of acute renal allograft rejection. In: UpToDate, Lam AQ, ed. Dec 16, 2024. UpToDate, Waltham, MA. Accessed on Oct 29, 2025.

4. Aubert O, Ursule-Dufait C, Brousse R, Gueguen J, Racapé M, Raynaud M, Van Loon E, Pagliuzzi A, Huang E, Jordan SC, Chavin KD, Gupta G, Kumar D, Alhamad T, Anand S, Sanchez-Garcia J, Abdalla BA, Hogan J, Garro R, Dadhania DM, Jain P, Mandelbrot DA, Naesens M, Dandamudi R, Dharnidharka VR, Anglicheau D, Lefaucheur C, Loupy A. Cell-free DNA for the detection of kidney allograft rejection. *Nat Med*. 2024 Aug;30(8):2320-2327. doi: 10.1038/s41591-024-03087-3. Epub 2024 Jun 2. PMID: 38824959; PMCID: PMC11333280.
5. Badiwala MV, Rao V. Tricuspid valve replacement after cardiac transplantation. *Curr Opin Cardiol*. 2007 Mar;22(2):123-7.
6. Bennett MK, Tang, WHW, Ch 61 endomyocardial biopsy. In: *Manual of Cardiovascular Medicine*. 4th ed. Lippincott Williams & Wilkins, Philadelphia; 2013.
7. Berger Y, Har Zahav Y, Kassif Y, Kogan A, Kuperstein R, Freimark D, Lavee J. Tricuspid Valve Regurgitation after Orthotopic Heart Transplantation: Prevalence and Etiology. *J Transplant*. 2012; 2012: 120702. Published online 2012 October 14.
8. Bernstein D. Section 7 – cardiac therapeutics. Chapter 434 – heart failure. Heart transplantation. In Behrman: *Nelson Textbook of Pediatrics*, 17th ed., St. Louis. W.B. Saunders; 2004.
9. Bernstein D, Williams GE, Eisen H, Mital S, Wohlgemuth JG, Klingler TM, Fang KC, Deng MC, Kobashigawa J. Gene expression profiling distinguishes a molecular signature for grade 1B mild acute cellular rejection in cardiac allograft recipients. *J Heart Lung Transplant*. 2007 Dec;26(12):1270-80.
10. Berry GJ, Burke MM, Andersen C, Bruneval P, Fedrigo M, Fishbein MC, Goddard M, Hammond EH, Leone O, Marboe C, Miller D, Neil D, Rassl D, Revelo MP, Rice A, Rene Rodriguez E, Stewart S, Tan CD, Winters GL, West L, Mehra MR, Angelini A. The 2013 International Society for Heart and Lung Transplantation Working Formulation for the standardization of nomenclature in the pathologic diagnosis of antibody-mediated rejection in heart transplantation. *J Heart Lung Transplant*. 2013 Dec;32(12):1147-62.
11. Bloom RD, Bromberg JS, Poggio ED, Bunnapradist S, Langone AJ, Sood P, et al.; Circulating Donor-Derived Cell-Free DNA in Blood for Diagnosing Active Rejection in Kidney Transplant Recipients (DART) Study Investigators. Cell-Free DNA and Active Rejection in Kidney Allografts. *J Am Soc Nephrol*. 2017 Jul;28(7):2221-2232.
12. BlueCross BlueShield Association (BCBSA). Technology Evaluation Center (TEC). Gene expression profiling as a noninvasive method to monitor for cardiac allograft rejection. TEC Assessment Program. Vol 26. No. 8. Chicago,IL. BCBSA. 2011 Nov.
13. Brennan DC, Alhamad T, Malone A. Kidney transplantation in adults: Clinical features and diagnosis of acute kidney allograft rejection. In: *UpToDate*, Legendre A and Vella J, eds. Sep 30, 2025. UpToDate, Waltham, MA. Accessed Oct 29, 2025.
14. Bromberg JS, Brennan DC, Poggio E, Bunnapradist S, Langone A, Sood P, Matas AJ, Mannon RB, Mehta S, Sharfuddin A, Fischbach B, Narayanan M, Jordan SC, Cohen DJ, Zaky ZS, Hiller D, Woodward RN, Grskovic M, Sninsky JJ, Yee JP, Bloom RD. Biological Variation of Donor-Derived Cell-Free DNA in Renal Transplant Recipients: Clinical Implications. *J Appl Lab Med*. 2017 Nov 1;2(3):309-321. doi: 10.1373/jalm.2016.022731. PMID: 33636851.

15. Bromberg JS, Bunnapradist S, Samaniego-Picota M, Anand S, Stites E, Gauthier P, Demko Z, Prewett A, Armer-Cabral M, Marshall K, Kaur N, Bloom MS, Tabriziani H, Bhorade S, Cooper M; ProActive Investigators. Elevation of Donor-derived Cell-free DNA Before Biopsy-proven Rejection in Kidney Transplant. *Transplantation*. 2024 Sep 1;108(9):1994-2004. doi: 10.1097/TP.0000000000005007. Epub 2024 Aug 20. PMID: 38595232; PMCID: PMC11335081.
16. Bu L, Gupta G, Pai A, Anand S, Stites E, Moinuddin I, Bowers V, Jain P, Axelrod DA, Weir MR, Wolf-Doty TK, Zeng J, Tian W, Qu K, Woodward R, Dholakia S, De Golovine A, Bromberg JS, Murad H, Alhamad T. Clinical outcomes from the Assessing Donor-derived cell-free DNA Monitoring Insights of kidney Allografts with Longitudinal surveillance (ADMIRAL) study. *Kidney Int*. 2022 Apr;101(4):793-803. doi: 10.1016/j.kint.2021.11.034. Epub 2021 Dec 22. PMID: 34953773.
17. Cadeiras M, Shahzad K, John MM, Gruber D, Bayern M, Auerbach S, Sinha A, Latif F, Unniachan S, Memon S, Mital S, Restaino S, Marboe CC, Addonizio LJ, Deng MC. Relationship between a validated molecular cardiac transplant rejection classifier and routine organ function parameters. *Clin Transplant*. 2010 May-Jun;24(3):321-7.
18. Cadeiras, M, von Bayern M, Sinha A, John M, Baron H; Restaino S, Deng, MC. Noninvasive diagnosis of acute cardiac allograft rejection. *Current Opinion in Organ Transplantation*. 12(5):543-550, October 2007.
19. CareDx. AlloMap® Heart. Nov 2024. Accessed Nov 5, 2025. Available at URL address: <https://caredx.com/products-and-services/transplant-services/heart/allomap/>
20. CareDx. HeartCare. 2024. Accessed Nov 5, 2025. Available at URL address: <https://caredx.com/products-and-services/transplant-services/heart/heartcare/>
21. Carey SA, Tecson KM, Jamil AK, Felius J, Wolf-Doty TK, Hall SA. Gene expression profiling scores in dual organ transplant patients are similar to those in heart-only recipients. *Transpl Immunol*. 2018 Aug;49:28-32. doi: 10.1016/j.trim.2018.03.003. Epub 2018 Mar 26.
22. Centers for Medicare and Medicaid Services (CMS). Medicare Coverage Database. Accessed Nov 5, 2025. Available at URL address: <https://www.cms.gov/medicare-coverage-database/search.aspx>
23. Costanzo MR, Dipchand A, Starling R, Anderson A, Chan M, Desai S, Fedson S, Fisher P, Gonzales-Stawinski G, Martinelli L, McGiffin D, Smith J, Taylor D, Meiser B, Webber S, Baran D, Carboni M, Dengler T, Feldman D, Frigerio M, Kfoury A, Kim D, Kobashigawa J, Shullo M, Stehlik J, Teuteberg J, Uber P, Zuckermann A, Hunt S, Burch M, Bhat G, Canter C, Chinnock R, Crespo-Leiro M, Delgado R, Dobbels F, Grady K, Kao W, Lamour J, Parry G, Patel J, Pini D, Towbin J, Wolfel G, Delgado D, Eisen H, Goldberg L, Hosenpud J, Johnson M, Keogh A, Lewis C, O'Connell J, Rogers J, Ross H, Russell S, Vanhaecke J; International Society of Heart and Lung Transplantation Guidelines. The International Society of Heart and Lung Transplantation Guidelines for the care of heart transplant recipients. *J Heart Lung Transplant*. 2010 Aug;29(8):914-56.
24. Crespo-Leiro MG, Stypmann J, Schulz U, Zuckermann A, Mohacsi P, Bara C, Ross H, Parameshwar J, Zakliczyński M, Fiocchi R, Hofer D, Colvin M, Deng MC, Leprince P, Elashoff B, Yee JP, Vanhaecke J. Clinical usefulness of gene-expression profile to rule out

acute rejection after heart transplantation: CARGO II. *Eur Heart J*. 2016 Sep 1;37(33):2591-601.

25. Crespo-Leiro MG, Stypmann J, Schulz U, Zuckermann A, Mohacsi P, Bara C, Ross H, Parameshwar J, Zakliczyński M, Fiocchi R, Hoefer D, Deng M, Leprince P, Hiller D, Eubank L, Deljkich E, Yee JP, Vanhaecke J. Performance of gene-expression profiling test score variability to predict future clinical events in heart transplant recipients. *BMC Cardiovasc Disord*. 2015 Oct 9;15:120.
26. Crespo-Leiro M, Zuckermann A, Stypmann J, et al. Increased plasma levels of donor-derived cell-free DNA correlate with rejection in heart transplant recipients: The CARGO II multicenter trial. *The Journal of Heart and Lung Transplantation*. 2015;34(4S):S31.
27. Dedrick, RL. Understanding gene expression patterns in immune-mediated disorders. *J Immunol*. 2007;4:3: 201-207.
28. Deng MC, Eisen HJ, Mehra MR, Billingham M, Marboe CC, Berry G, Kobashigawa J, Johnson FL, Starling RC, Murali S, Pauly DF, Baron H, Wohlgemuth JG, Woodward RN, Klingler TM, Walther D, Lal PG, Rosenberg S, Hunt S; CARGO Investigators. Noninvasive discrimination of rejection in cardiac allograft recipients using gene expression profiling. *Am J Transplant*. 2006 Jan;6(1):150-60.
29. Deng MC, Elashoff B, Pham MX, Teuteberg JJ, Kfoury AG, Starling RC, Cappola TP, Kao A, Anderson AS, Cotts WG, Ewald GA, Baran DA, Bogaev RC, Shahzad K, Hiller D, Yee J, Valentine HA; IMAGE Study Group. Utility of gene expression profiling score variability to predict clinical events in heart transplant recipients. *Transplantation*. 2014 Mar 27;97(6):708-14.
30. Deshpande SR, Zangwill SD, Richmond ME, Kindel SJ, Schroder JN, Gaglianella N, Bichell DP, Wigger MA, Knecht KR, Thrush PT, Mahle WT, North PE, Simpson PM, Zhang L, Dasgupta M, Tomita-Mitchell A, Mitchell ME. Evaluating threshold for donor fraction cell-free DNA using clinically available assay for rejection in pediatric and adult heart transplantation. *Pediatr Transplant*. 2024 May;28(3):e14708. doi: 10.1111/petr.14708. PMID: 38553812; PMCID: PMC10987078.
31. De Vlaminck I, Valentine HA, Snyder TM, Strehl C, Cohen G, Luikart H, Neff NF, Okamoto J, Bernstein D, Weisshaar D, Quake SR, Khush KK. Circulating cell-free DNA enables noninvasive diagnosis of heart transplant rejection. *Sci Transl Med*. 2014 Jun 18;6(241):241ra77. doi: 10.1126/scitranslmed.3007803. PMID: 24944192; PMCID: PMC4326260.
32. Eisen HJ. Heart transplantation in adults: Cardiac allograft vasculopathy risk factors, surveillance, and diagnosis. In: *UpToDate*, Borlaug BA, ed. May 7, 2025. UpToDate, Waltham, MA. Accessed Oct 24, 2025.
33. Eisen, HJ. Heart transplantation in adults: Diagnosis of acute allograft rejection. In: *UpToDate*, Borlaug BA, ed. Jun 23, 2025. UpToDate, Waltham, MA. Accessed Oct 28, 2025.
34. Eisen HJ and Khush KK. Heart transplantation in adults: Treatment of rejection. In: *UpToDate*, Borlaug BA, ed. Feb 2, 2024. UpToDate, Waltham, MA. Accessed Oct 28, 2025.
35. Fang KC. Clinical utilities of peripheral blood gene expression profiling in the management of cardiac transplant patients. *J Immunol* 2007;4:3:209-217.

36. Fujita B, Prashovikj E, Schulz U, Börgermann J, Sunavsky J, Fuchs U, Gummert J, Ensminger S. Predictive value of gene expression profiling for long-term survival after heart transplantation. *Transpl Immunol*. 2017 Mar;41:27-31.
37. Ginsburg GS, Haga SB. Translating genomic biomarkers into clinically useful diagnostics. *Expert Rev Mol Diagn*. 2006 Mar;6(2):179-91.
38. Grskovic M, Hiller DJ, Eubank LA, Sninsky JJ, Christopherson C, Collins JP, et al. Validation of a Clinical-Grade Assay to Measure Donor-Derived Cell-Free DNA in Solid Organ Transplant Recipients. *J Mol Diagn*. 2016 Nov;18(6):890-902.
39. Henricksen EJ, Moayedi Y, Purewal S, Twiggs JV, Waddell K, Luikart H, Han J, Feng K, Wayda B, Lee R, Shudo Y, Jimenez S, Khush KK, Teuteberg JJ. Combining donor derived cell free DNA and gene expression profiling for non-invasive surveillance after heart transplantation. *Clin Transplant*. 2023 Mar;37(3):e14699. doi: 10.1111/ctr.14699. Epub 2022 May 23. PMID: 35559582.
40. Hidestrand M, Tomita-Mitchell A, Hidestrand PM, Oliphant A, Goetsch M, Stamm K, Liang HL, Castleberry C, Benson DW, Stendahl G, Simpson PM, Berger S, Tweddell JS, Zangwill S, Mitchell ME. Highly sensitive noninvasive cardiac transplant rejection monitoring using targeted quantification of donor-specific cell-free deoxyribonucleic acid. *J Am Coll Cardiol*. 2014 Apr 1;63(12):1224-1226. doi: 10.1016/j.jacc.2013.09.029. Epub 2013 Oct 16. PMID: 24140666; PMCID: PMC4988656.
41. Jordan SC, Bunnapradist S, Bromberg JS, Langone AJ, Hiller D, Yee JP, et al. Donor-derived Cell-free DNA Identifies Antibody-mediated Rejection in Donor Specific Antibody Positive Kidney Transplant Recipients. *Transplant Direct*. 2018 Aug 20;4(9):e379.
42. Kamath M, Shekhtman G, Grogan T, Hickey MJ, Silacheva I, Shah KS, Shah KS, Hairapetian A, Gonzalez D, Godoy G, Reed EF, Elashoff D, Bondar G, Deng MC. Variability in Donor-Derived Cell-Free DNA Scores to Predict Mortality in Heart Transplant Recipients - A Proof-of-Concept Study. *Front Immunol*. 2022 Feb 18;13:825108. doi: 10.3389/fimmu.2022.825108. PMID: 35251005; PMCID: PMC8895247.
43. Keslar K, Rodriguez ER, Tan CD, Starling RC, Heeger PS. Complement gene expression in human cardiac allograft biopsies as a correlate of histologic grade of injury. *Transplantation*. 2008 Nov 15;86(9):1319-21.
44. Khush K, Hall S, Kao A, Raval N, Dhingra R, Shah P, Bellumkonda L, Ravichandran A, Van Bakel A, Uriel N, Patel S, Pinney S, DePasquale E, Baran DA, Pinney K, Oreschak K, Kobulnik J, Shen L, Teuteberg J. Surveillance with dual noninvasive testing for acute cellular rejection after heart transplantation: Outcomes from the Surveillance HeartCare Outcomes Registry. *J Heart Lung Transplant*. 2024 Sep;43(9):1409-1421. doi: 10.1016/j.healun.2024.05.003. Epub 2024 May 15. PMID: 38759766.
45. Khush KK, Patel J, Pinney S, Kao A, Alharethi R, DePasquale E, Ewald G, Berman P, Kanwar M, Hiller D, Yee JP, Woodward RN, Hall S, Kobashigawa J. Noninvasive detection of graft injury after heart transplant using donor-derived cell-free DNA: A prospective multicenter study. *Am J Transplant*. 2019 Oct;19(10):2889-2899. doi: 10.1111/ajt.15339. Epub 2019 Apr 8. PMID: 30835940; PMCID: PMC6790566.

46. Kidney Disease: Improving Global Outcomes (KDIGO). Clinical Practice Guideline for Care of Kidney Transplant Recipients. 2009. Accessed Nov 5, 2025. Available at URL address: <https://kdigo.org/guidelines/transplant-recipient/>
47. Kim PJ, Olymbios M, Siu A, Wever Pinzon O, Adler E, Liang N, Swenerton R, Sternberg J, Kaur N, Ahmed E, Chen YA, Fehringer G, Demko ZP, Billings PR, Stehlik J. A novel donor-derived cell-free DNA assay for the detection of acute rejection in heart transplantation. *J Heart Lung Transplant*. 2022 Jul;41(7):919-927. doi: 10.1016/j.healun.2022.04.002. Epub 2022 Apr 10. PMID: 35577713; PMCID: PMC9670834.
48. Kobashigawa J, Patel J, Azarbal B, Kittleson M, Chang D, Czer L, Daun T, Luu M, Trento A, Cheng R, Esmailian F. Randomized pilot trial of gene expression profiling versus heart biopsy in the first year after heart transplant: early invasive monitoring attenuation through gene expression trial. *Circ Heart Fail*. 2015 May;8(3):557-64.
49. LuminoDx Inc. HepatoTrack. 2025. Accessed Nov 5, 2025. Available at URL address: <https://luminodx.com/acute-allograft-rejection/>
50. Marsh CL, Kurian SM, Rice JC, Whisenant TC, David J, Rose S, Schieve C, Lee D, Case J, Barrick B, Peddi VR, Mannon RB, Knight R, Maluf D, Mandelbrot D, Patel A, Friedewald JJ, Abecassis MM, First MR. Application of TruGraf v1: A Novel Molecular Biomarker for Managing Kidney Transplant Recipients With Stable Renal Function. *Transplant Proc*. 2019 Apr;51(3):722-728. doi: 10.1016/j.transproceed.2019.01.054. Epub 2019 Jan 26. PMID: 30979456.
51. Mehra MR. The emergence of genomic and proteomic biomarkers in heart transplantation. *J Heart Lung Transplant*. 2005 Jul;24(7 Suppl):S213-8.
52. Mehra MR, Crespo-Leiro MG, Dipchand A, Ensminger SM, Hiemann NE, Kobashigawa JA, Madsen J, Parameshwar J, Starling RC, Uber PA. International Society for Heart and Lung Transplantation working formulation of a standardized nomenclature for cardiac allograft vasculopathy-2010. *J Heart Lung Transplant*. 2010 Jul;29(7):717-27. doi: 10.1016/j.healun.2010.05.017. Erratum in: *J Heart Lung Transplant*. 2011 Mar;30(3):360. PMID: 20620917.
53. Mehra MR, Kobashigawa JA, Deng MC, Fang KC, Klingler TM, Lal PG, Rosenberg S, Uber PA, Starling RC, Murali S, Pauly DF, Dedrick R, Walker MG, Zeevi A, Eisen HJ; CARGO Investigators. Clinical implications and longitudinal alteration of peripheral blood transcriptional signals indicative of future cardiac allograft rejection. *J Heart Lung Transplant*. 2008 Mar;27(3):297-301.
54. Mehra MR, Kobashigawa JA, Deng MC, Fang KC, Klingler TM, Lal PG, Rosenberg S, Uber PA, Starling RC, Murali S, Pauly DF, Dedrick R, Walker MG, Zeevi A, Eisen HJ; CARGO Investigators. Transcriptional signals of T-cell and corticosteroid-sensitive genes are associated with future acute cellular rejection in cardiac allografts. *J Heart Lung Transplant*. 2007 Dec;26(12):1255-63.
55. Mehra MR, Parameshwar J. Gene expression profiling and cardiac allograft rejection monitoring: is IMAGE just a mirage? *J Heart Lung Transplant*. 2010 Jun;29(6):599-602.
56. Mehra MR, Uber PA. Genomic biomarkers and heart transplantation. *Heart Fail Clin*. 2007 Jan;3(1):83-6.

57. Mehra MR, Uber PA, Walther D, Vesely M, Wohlgemuth JG, Prentice J, Tayama D, Billingham M. Gene expression profiles and B-type natriuretic peptide elevation in heart transplantation: more than a hemodynamic marker. *Circulation*. 2006 Jul 4;114(1 Suppl):I21-6.
58. Menon MC, Murphy B, Heeger PS. Moving Biomarkers toward Clinical Implementation in Kidney Transplantation. *J Am Soc Nephrol*. 2017 Mar;28(3):735-747.
59. Moayed Y, Foroutan F, Miller RJH, Fan CS, Posada JGD, Alhussein M, Tremblay-Gravel M, Oro G, Luikart HI, Yee J, Shullo MA, Khush KK, Ross HJ, Teuteberg JJ. Risk evaluation using gene expression screening to monitor for acute cellular rejection in heart transplant recipients. *J Heart Lung Transplant*. 2019 Jan;38(1):51-58. doi: 10.1016/j.healun.2018.09.004. Epub 2018 Sep 12.
60. Natera. Prospera™ Heart. 2025. Accessed on Nov 5, 2025. Available at URL address: <https://www.natera.com/organ-health/prospera-heart-transplantation-assessment/>
61. Oellerich M, Schütz E, Kanzow P, Schmitz J, Beck J, Kollmar O, et al. Use of graft-derived cell-free DNA as an organ integrity biomarker to reexamine effective tacrolimus trough concentrations after liver transplantation. *Ther Drug Monit*. 2014 Apr;36(2):136-40.
62. Patel JK, Kobashigawa JA. Should we be doing routine biopsy after heart transplantation in a new era of anti-rejection? *Curr Opin Cardiol*. 2006 Mar;21(2):127-31.
63. Pham MX, Deng MC, Kfoury AG, Teuteberg JJ, Starling RC, Valantine H. Molecular testing for long-term rejection surveillance in heart transplant recipients: design of the Invasive Monitoring Attenuation Through Gene Expression (IMAGE) trial. *J Heart Lung Transplant*. 2007 Aug;26(8):808-14.
64. Pham MX, Teuteberg JJ, Kfoury AG, Starling RC, Deng MC, Cappola TP, Kao A, Anderson AS, Cotts WG, Ewald GA, Baran DA, Bogaev RC, Elashoff B, Baron H, Yee J, Valantine HA; IMAGE Study Group. Gene-expression profiling for rejection surveillance after cardiac transplantation. *N Engl J Med*. 2010a May 20;362(20):1890-900.
65. Pham MX, Teuteberg JJ, Kfoury AG, et al. Gene-expression profiling for rejection surveillance after cardiac transplantation. *N Engl J Med* 2010b;362:1890-900. DOI: 10.1056/NEJMoa0912965; supplement.
66. Renlund DG, Taylor DO, Smedira NG. Chapter 90 cardiac transplantation and mechanical circulatory support. Cardiac allograft rejection. In: Topol EJ, editor. *Textbook of Cardiovascular Medicine*. Philadelphia: Lippincott Williams & Wilkins: 2007. p 1429-36.
67. Schmauss D, Weis M. Cardiac allograft vasculopathy: recent developments. *Circulation*. 2008 Apr 22;117(16):2131-41.
68. Sharon E, Shi H, Kharbanda S, Koh W, Martin LR, Khush KK, et al. Quantification of transplant-derived circulating cell-free DNA in absence of a donor genotype. *PLoS Comput Biol*. 2017 Aug 3;13(8):e1005629.
69. Snyder TM, Khush KK, Valantine HA, Quake SR. Universal noninvasive detection of solid organ transplant rejection. *Proc Natl Acad Sci U S A*. 2011 Apr 12;108(15):6229-34.

70. Society for Cardiovascular Pathology. International Society for Heart and Lung Transplantation (ISHLT) revised grading criteria. 2025. Accessed Oct 28, 2025. Available at URL address: <https://www.scvp.net/acute-cellular-rejection-tutorial/#grading-criteria>
71. Starling RC, Pham M, Valantine H, Miller L, Eisen H, Rodriguez ER, Taylor DO, Yamani MH, Kobashigawa J, McCurry K, Marboe C, Mehra MR, Zuckerman A, Deng MC; Working Group on Molecular Testing in Cardiac Transplantation. Molecular testing in the management of cardiac transplant recipients: initial clinical experience. *J Heart Lung Transplant*. 2006 Dec;25(12):1389-95.
72. Stewart S, Winters GL, Fishbein MC, Tazelaar HD, Kobashigawa J, Abrams J, Andersen CB, Angelini A, Berry GJ, Burke MM, Demetris AJ, Hammond E, Itescu S, Marboe CC, McManus B, Reed EF, Reinsmoen NL, Rodriguez ER, Rose AG, Rose M, Suciu-Focia N, Zeevi A, Billingham ME. Revision of the 1990 working formulation for the standardization of nomenclature in the diagnosis of heart rejection. *J Heart Lung Transplant*. 2005 Nov;24(11):1710-20. doi: 10.1016/j.healun.2005.03.019. Epub 2005 Jun 20. PMID: 16297770.
73. Strecker T, Rösch J, Weyand M, Agaimy A. Endomyocardial biopsy for monitoring heart transplant patients: 11-years-experience at a German heart center. *Int J Clin Exp Pathol*. 2013;6(1):55-65.
74. Thermo Fisher Scientific, Inc. MMDx® for Molecular Biopsy Assessment. 2006-2025. Accessed on Oct 29, 2025. Available at URL address: <https://www.thermofisher.com/onelambda/us/en/post-transplant/molecular-biopsy-assessment.html#:~:text=Molecular%20Microscope%20Diagnostics%20System%20%28MMDx%29%20is%20new%20microarray,assessment%20of%20rejection%20and%20injury%20in%20transplanted%20organs>.
75. U. S. Food and Drug Administration (FDA). 510(k) substantial equivalence determination decision summary. AlloMap® Molecular Expression Testing. K073482. Aug 8, 2008. Accessed Oct 29, 2025. Available at URL address: http://www.accessdata.fda.gov/cdrh_docs/reviews/K073482.pdf
76. U. S. Food and Drug Administration (FDA). Guidance for Industry and FDA Staff - Class II Special Controls Guidance Document: Cardiac Allograft Gene Expression Profiling Test Systems. Oct 21, 2009, current as of Aug 20, 2018. Accessed Oct 29, 2025. Available at URL address: <http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/GuidanceDocuments/ucm187084.htm>
77. U.S. Organ Procurement and Transplantation Network and the Scientific Registry of Transplant Recipients. National data as of Oct 26, 2025. Accessed on Oct 27, 2025. Available at URL address: <https://optn.transplant.hrsa.gov/data/view-data-reports/national-data/>
78. Velleca A, Shullo MA, Dhital K, Azeka E, Colvin M, DePasquale E, Farrero M, García-Guereta L, Jamero G, Khush K, Lavee J, Pouch S, Patel J, Michaud CJ, Shullo MA, Schubert S, Angelini A, Carlos L, Mirabet S, Patel J, Pham M, Urschel S, Kim KH, Miyamoto S, Chih S, Daly K, Grossi P, Jennings DL, Kim IC, Lim HS, Miller T, Potena L, Velleca A, Eisen H, Bellumkonda L, Danziger-Isakov L, Dobbels F, Harkess M, Kim D, Lyster H, Peled Y, Reinhardt Z. The International Society for Heart and Lung Transplantation (ISHLT) guidelines for the care of heart transplant recipients. *J Heart Lung Transplant*. 2023

May;42(5):e1-e141. doi: 10.1016/j.healun.2022.10.015. Epub 2022 Dec 20. PMID: 37080658.

79. Verhoeven JGHP, Boer K, Van Schaik RHN, Manintveld OC, Huibers MMH, Baan CC, et al. Liquid Biopsies to Monitor Solid Organ Transplant Function: A Review of New Biomarkers. *Ther Drug Monit.* 2018 Oct;40(5):515-525.
80. Yamani MH, Taylor DO. Sec 2 Heart Transplantation. In: *Cleveland Clinic: Current Clinical Medicine*, 2nd ed. Saunders, Maryland Heights, MO. 2010 pgs 180-186.
81. Yamani MH, Taylor DO, Rodriguez ER, Cook DJ, Zhou L, Smedira N, Starling RC. Transplant vasculopathy is associated with increased AlloMap gene expression score. *J Heart Lung Transplant.* 2007a Apr;26(4):403-6.
82. Yamani MH, Taylor DO, Haire C, Smedira N, Starling RC. Post-transplant ischemic injury is associated with up-regulated AlloMap gene expression. *Clin Transplant.* 2007b Jul-Aug;21(4):523-5.
83. Zhang H, Zheng C, Li X, et al. Diagnostic Performance of Donor-Derived Plasma Cell-Free DNA Fraction for Antibody-Mediated Rejection in Post Renal Transplant Recipients: A Prospective Observational Study. *Front Immunol.* 2020;11:342. Published 2020 Feb 28.
84. Zhou Y, Yang G, Liu H, Chen Y, Li X, Ge J, et al. A Noninvasive and Donor-independent Method Simultaneously Monitors Rejection and Infection in Patients With Organ Transplant. *Transplant Proc.* 2019 Jul - Aug;51(6):1699-1705.

Revision Details

Type of Revision	Summary of Changes	Date
Annual review	<ul style="list-style-type: none"> Added example of HepatoTrack™ to other gene expression profiling tests Added example of OmniGraf® Liver to donor-derived cell-free DNA testing Removed examples VitaGraft™ Kidney Baseline + 1st Plasma Test, VitaGraft™ Kidney Subsequent from donor-derived cell-free DNA testing 	12/15/2025
Focused review	<ul style="list-style-type: none"> Added example of Prospera™ to donor-derived cell-free DNA testing 	9/15/2025
Focused review	<ul style="list-style-type: none"> Added examples of AlloSure® and VitaGraft™ Kidney 2.0 to donor-derived cell-free DNA testing 	4/15/2025
Focused review	<ul style="list-style-type: none"> Added example of Trugraf back to non-coverage statement for other gene expression profiling tests 	2/15/2025
Annual review	<ul style="list-style-type: none"> Added coverage for Allomap starting at two months post-transplant Removed examples TruGraf, AlloSure®, Prospera™, and the coverage statement for combined gene expression profiling and donor-derived cell-free DNA testing (i.e. HeartCare®, OmniGraf™). These are 	11/1/2024

	<p>addressed in the Cigna/EviCore cobranded guidelines, and therefore removed from this policy.</p> <ul style="list-style-type: none">• Removed the example Viracor TRAC from the coverage statement for donor-derived cell-free DNA testing• Added VitaGraft™ Kidney to examples of donor-derived cell-free DNA testing	
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