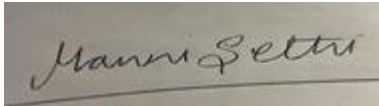


**Prior Authorization Review Panel
MCO Policy Submission**

A separate copy of this form must accompany each policy submitted for review.
Policies submitted without this form will not be considered for review.

Plan: AmeriHealth Caritas Pennsylvania Community HealthChoices & Keystone First Community HealthChoices	Submission Date: 1/2/2025
Policy Number: ccp.1202	Effective Date: 1/2016 Revision Date: November 1, 2024
Policy Name: Lung transplants	
Type of Submission – Check all that apply: <div style="margin-left: 20px;"><input type="checkbox"/> New Policy <input checked="" type="checkbox"/> Revised Policy* <input type="checkbox"/> Annual Review – No Revisions <input type="checkbox"/> Statewide PDL</div>	
<p>*All revisions to the policy <u>must</u> be highlighted using track changes throughout the document.</p> <p>Please provide any clarifying information for the policy below:</p> <p style="color: red;">See tracked changes below.</p>	
Name of Authorized Individual (Please type or print): Manni Sethi, MD, MBA, CHCQM	Signature of Authorized Individual: 

Lung transplants

Clinical Policy ID: CCP.1202

Recent review date: 11/2024

Next review date: 3/2026

Policy contains: Heart-lung transplant; lung transplant, pulmonary transplant.

AmeriHealth Caritas Pennsylvania Community HealthChoices has developed clinical policies to assist with making coverage determinations. AmeriHealth Caritas Pennsylvania Community HealthChoices' clinical policies are based on guidelines from established industry sources, such as the Centers for Medicare & Medicaid Services (CMS), state regulatory agencies, the American Medical Association (AMA), medical specialty professional societies, and peer-reviewed professional literature. These clinical policies along with other sources, such as plan benefits and state and federal laws and regulatory requirements, including any state- or plan-specific definition of "medically necessary," and the specific facts of the particular situation are considered by AmeriHealth Caritas Pennsylvania Community HealthChoices on a case by case basis, when making coverage determinations. In the event of conflict between this clinical policy and plan benefits and/or state or federal laws and/or regulatory requirements, the plan benefits and/or state and federal laws and/or regulatory requirements shall control. AmeriHealth Caritas Pennsylvania Community HealthChoices' clinical policies are for informational purposes only and not intended as medical advice or to direct treatment. Physicians and other health care providers are solely responsible for the treatment decisions for their patients. AmeriHealth Caritas Pennsylvania Community HealthChoices' clinical policies are reflective of evidence-based medicine at the time of review. As medical science evolves, AmeriHealth Caritas Pennsylvania Community HealthChoices will update its clinical policies as necessary. AmeriHealth Caritas Pennsylvania Community HealthChoices' clinical policies are not guarantees of payment.

Coverage policy

Lung transplantation is clinically proven and, therefore, may be medically necessary in cases of end-stage lung disease, when all of the following general selection criteria are met:

- A > 50% risk of death due to lung disease within two years if lung transplantation is not performed.
- A > 80% likelihood of five-year post-transplant survival from a general medical perspective provided there is adequate graft function.
- No absolute contraindications (See limitations) (Leard, 2021).

Heart-lung transplantation is clinically proven and, therefore, may be medically necessary for members with intrinsic lung disease or severe pulmonary artery hypertension and severe structural or cardiac dysfunction unlikely to improve with normalization of pulmonary pressures (Bermudez, 2021; Leard, 2021).

Limitations

The transplanting institution may require additional general selection criteria or disease-specific criteria for medical necessity determination.

Absolute contradictions to lung transplantation include (Leard, 2021):

- Lack of patient willingness or acceptance of transplant.
- Malignancy with high risk of recurrence or death related to cancer.

- Glomerular filtration rate < 40 mL/min/1.73m² unless being considered for multi-organ transplant.
- Acute coronary syndrome or myocardial infarction within 30 days (excluding demand ischemia).
- Stroke within 30 days.
- Liver cirrhosis with portal hypertension or synthetic dysfunction unless being considered for multi-organ transplant.
- Acute liver failure.
- Acute renal failure with rising creatinine or on dialysis and low likelihood of recovery.
- Septic shock.
- Active extrapulmonary or disseminated infection.
- Active tuberculosis infection.
- Human immunodeficiency virus infection with detectable viral load.
- Limited functional status (e.g., non-ambulatory) with poor potential for post-transplant rehabilitation.
- Progressive cognitive impairment.
- Repeated episodes of non-adherence without evidence of improvement (Note: This is not an absolute contraindication for pediatric members. Ongoing assessment of non-adherence should occur as they progress through different developmental stages.)
- Active substance use or dependence including current tobacco use, vaping, marijuana smoking, or intravenous drug use.
- Other severe uncontrolled medical conditions expected to limit survival after transplant.

Relative contraindications are listed in the Appendix.

Alternative covered services

Guideline-directed maximum medical management of underlying disease.

Background

Lung transplantation, or pulmonary transplantation, is a surgical procedure in which a patient's diseased lung(s) are partially or totally replaced by healthy lungs from a donor. Donor lungs can be retrieved from a living donor or a deceased donor. A living donor can only donate one lung lobe. With some lung diseases, a recipient may only need to receive a single lung. With other lung diseases, such as cystic fibrosis, it is imperative that a recipient receive two lungs. While lung transplants carry certain associated risks, such as life-threatening complications and infections, especially in the first year after surgery, they can also extend life expectancy and enhance the quality of life for patients with end-stage pulmonary disease (U.S. National Heart, Lung, and Blood Institute, 2022).

The number of Americans undergoing a lung transplant was 2,562 in 2018, an increase of 31% from five years earlier (Valapour, 2020). As of August 23, 2022, 1,011 registered Americans were awaiting a lung transplant, plus an additional 36 for heart/lung transplants (U.S. Health Resources and Services Administration, 2023b).

The United Network for Organ Sharing maintains a national U.S. registry for organ matching. Its purposes are to operate and monitor a system for allocation of organs donated for transplantation, maintain a waiting list of potential recipients, and match potential recipients with organ donors according to established medical criteria

(U.S. Health Resources and Services Administration, 2023a). Over time, the system for lung allocation has evolved to increase the availability of organs for transplant and their success through better donor matching.

The expanding waiting list for lung transplants prompted a 2005 change in the United States to assign priority to candidates based on a greater predicted survival benefit from transplantation and waitlist urgency, calculated as the Lung Allocation Score, instead of waiting time. These changes impacted patients aged 12 years and older. Subsequent changes included exceptions for younger patients and geographic distribution. On December 6, 2021, the Continuous Distribution system was approved to replace the Lung Allocation Scoring system and is anticipated to take effect early in 2023 (Benvenuto, 2021).

The composite allocation score for lung transplants was officially approved in December 2021, with phased implementation beginning in November 2022. This new system, replacing the Lung Allocation Score (LAS), considers a score of 25 or higher as the threshold for listing a candidate. It prioritizes patients based on five factors: medical urgency (25%), post-transplant survival (25%), patient access (25%), biological disadvantages (15%), and placement efficiency (10%). Developed by the Organ Procurement and Transplantation Network (OPTN) Lung Transplantation Committee, this system arose from a process involving community feedback, simulations, and analysis. It aims to balance urgency, survival probability, and logistical factors, marking a significant improvement in lung allocation policy (Organ Procurement and Transplantation Network, 2023).

A deceased donor, also known as cadaveric donor, is the most common donor source used for lung transplantation. Use of a live donor as a source for lung transplantation was initiated in 1993 due to the higher demand than supply for patients waiting for lung transplantation. Deceased donor transplantation is preferred, and the proportion of transplants from live donors is falling due to the change in allocating lungs from deceased donors (Date, 2017). In 2022, there were 1,918 deceased donor lungs and no living donor lungs transplanted in U.S. recipients (U.S. Health Resources and Services Administration, 2023c).

Findings

A number of guidelines affecting lung transplants have been published. The definitive work addressing criteria on selecting lung transplant patients, and contraindications against performing surgery, were written by the Pulmonary Scientific Council of the International Society for Heart and Lung Transplantation, which published versions in 1998, 2006, and 2015 (Weill, 2015, 2018). The same group of international experts that wrote the 2006 guidelines recently developed a consensus opinion on timing of referral and listing for lung transplantation. The panel included lung allocation scores and expanded indications for and contraindications to lung transplant (Weill, 2015).

In 2021, the International Society for Heart and Lung Transplantation updated its consensus guidance on selection of lung transplant candidates (Leard, 2021). This guidance differs from prior versions by creating categories for risk factors, considering the candidate as a whole and the risk tolerance and expertise of transplant centers. They stressed optimizing all potentially modifiable risk factors prior to lung transplant to improve long-term outcomes.

Two consensus papers from the International Society for Heart and Lung Transplantation provide additional guidance on cardiothoracic transplantation in patients with connective tissue disease (Bermudez, 2021; Crespo, 2021). Patients with connective tissue disease and advanced lung disease are medically complex and often regarded as poor surgical risks, but research demonstrates comparable survival and allograft dysfunction outcomes for carefully selected patients undergoing lung transplantation for other indications.

The Society defined the disease states considered connective tissue diseases and described their unique extrapulmonary manifestations that require special management before, during, and after lung or heart-lung transplantation. They recommend considering lung transplantation when the clinical status has progressively

declined despite maximal medical therapy. Finally, the Society listed absolute and relative contraindications for several connective tissue diseases to improve risk stratification and selection of candidates for lung transplantation based on consensus, recognizing the paucity of evidence on specific predictors of prognosis after cardiothoracic transplantation in this population (Crespo, 2021).

Bilateral lung transplantation is generally preferred over single lung transplantation for its theoretical survival and functional advantages over time, but single lung transplantation can be offered in the absence of secondary pulmonary hypertension when the clinical condition requires a shorter time on the waiting list. Heart-lung transplantation is reserved for patients with intrinsic lung disease or severe pulmonary artery hypertension and severe structural or cardiac dysfunction unlikely to improve with normalization of pulmonary pressures. Patients with moderate/severe left ventricular dysfunction or profound right ventricular dilation and dysfunction on high doses of inotropic support should be considered at high risk for lung transplantation alone (Bermudez, 2021).

Lung re-transplants are now being performed in increasing numbers. The median survival rate for those with re-transplants is lower than that of transplants (2.6 years versus 5.6 years), and the five-year survival rate is lower as well, at 34.5% versus 53.3%. Lower survival was associated with single lung transplants ($P = .021$), transplantations done between 2009 and 2013 ($P = .041$), multiple retransplantations ($P = .023$), and recipients requiring pre-transplantation ventilator support. Careful selection of candidates for re-transplant is advised (Thomas, 2015).

A systematic review/meta-analysis of 54 studies of over 5,000 subjects determined that *B. cepacia* complex significantly increased the risk of mortality after lung transplantation. Other factors that did not affect mortality included *P. aeruginosa* colonization, forced expiratory volume in one second, pulmonary hypertension, body mass index, pancreatic insufficiency, and cystic fibrosis-related diabetes (Koutsokera, 2019).

One study of 8,778 lung transplant patients showed a significantly lower survival ($P < .001$) for single lung transplants. Nearly 92% of the patients had a lung allocation score over 75, representing patients more likely to benefit from a lung transplant (Black, 2014). A study of 580 Canadian lung transplant patients with cystic fibrosis found the five- and 10-year survival rates to be 67% and 50%, with significantly higher death rates for young and old patients, plus those with pancreatic sufficiency or *B. cepacia* infection (Stephenson, 2015).

A systematic review/meta-analysis of 30 studies ($n = 4,092$) concluded that bilateral (versus unilateral) lung transplants had better long-term survival, better postoperative lung function and less bronchiolitis obliterans syndrome. No differences existed for in-hospital mortality and postoperative complications (Yu, 2019).

Another systematic review of 5,601 lung transplant recipients showed a clear pattern of greater survival among double transplant patients, namely 57% versus 50% after five years, 35.3% versus 27.8% after 10 years, and 24% versus 13.9% after 15 years (Wilson-Smith, 2020).

An exception to the pattern of lower mortality for double lung transplants was found in a recent systematic review/meta-analysis of 16 studies ($n = 17,872$) that addressed outcomes of lung transplant patients with idiopathic pulmonary fibrosis. Those with single transplants had a lower post-treatment percentage of Forced Exhaled Volume in one second than those with double transplants ($P < .001$). Survival outcomes were not significantly different at $P < .39$, but after single transplants, significantly fewer deaths were due to primary graft dysfunction and more due to malignancy, both $P < .001$ (Li, 2020).

A systematic review of six studies ($n = 1,305$) showed enhancement in quality of life for lung transplant candidates in five of the studies, using the SF-36 questionnaire and the six-minute walk test (Hoffman, 2017). A large systematic review of 73 studies showed that quality of life after lung transplants is enhanced, especially during the first year after the procedure in physical health and functioning domains, and may improve more after bilateral transplants and heart and liver transplants than after single lung transplants (Singer, 2013).

Some studies have identified factors that increase the risk of mortality in lung transplant patients. A review of 13 cohort studies ($n = 40,742$) reviewed weight at surgery and found significantly elevated relative risks compared to normal weight for underweight (1.36), obese (1.90), and overweight plus underweight (1.36) (Upala, 2016). Clinical risk factors were identified in 13 studies ($n = 10,042$) of primary graft dysfunction; the main causes of elevated morbidity and mortality include female gender, African American race, idiopathic pulmonary fibrosis, sarcoidosis, primary pulmonary hypertension, elevated body mass index, and use of cardiopulmonary bypass (Liu, 2014).

Non-adherence to medical regimens after transplants was the focus of a systematic review. While non-adherence rates varied across risk factors, they were not a significant factor in mortality (Hu, 2017).

Lung transplantations from deceased donors remain a minority of all such procedures; one meta-analysis of nine studies ($n = 301$) compared the mortality of patients who received a graft from deceased and live (conventional) donors. One year survival ranged from 50% to 100% among transplants from deceased donors, compared to 72% to 88% from live donors. One of the studies included nearly half of the transplants (138 of 301); one-year survival was significantly lower among deceased donor transplants (65.1% versus 84.1% for live donor transplants, $P < .001$) (Eberlein, 2017).

A comparison of lungs transplanted after > 12 hours ex-vivo preservation by splitting one cold ischemic time into two shorter ones ($n = 97$, average 875.7 minutes) with lungs transplanted with single waits < 12 hours ($n = 809$, average 400.8 minutes) was made. Median length of stay for both hospital and intensive-care unit lengths of stay did not differ significantly between groups (23.0 days versus 25.5 days, $P = .60$, and 4.0 days versus 4.0 days, $P = .53$). Primary graft dysfunction grade was not significantly different between the groups 72 hours after transplantation ($P = .85$), and there was no significant difference in survival ($P = .61$) (Yeung, 2017).

A systematic review of 13 studies found the ability of cardiopulmonary exercise testing to assist in making decisions in optimal timing for lung transplantation was limited due to problems of retrospective studies, patient selection, insufficient adjustment for confounders, and inadequate statistical analyses (Barratt, 2020).

An analysis of 65,265 lung transplant patients (median age 50.3 years) followed an average of 5.2 years found 17.3% died of cancer after surgery, a rate 4.28-fold higher than the general population. The most common types of malignancies causing death were lymphatic/hematological, integumentary, respiratory, digestive, and reproductive/urinary. Use of immunosuppressive therapy contributed in part to high rates. Authors state findings can improve individualized guidance for transplant patients (Ge, 2020).

In 2022, we added three updated consensus statements (Bermudez, 2021; Crespo, 2021; Leard, 2021), described earlier, and three new systematic reviews and meta-analyses. A meta-analysis of 72 eligible studies found no donor variables and only post-transplantation need for extra-corporeal membrane oxygenation that predicted one-year mortality with high certainty (hazard ratio 1.91, 95% confidence interval 1.79 to 2.04) in adult recipients. The authors stated lack of prognostic significance for some widely accepted factors (e.g., donor smoking, age) may relate to limits in the range of these variables found in selected donors and recipients (Foroutan, 2022).

A systematic review of 27 low-quality studies found 95% early survival rates in recipients with end-stage lung disease from COVID-19 infection. The data suggest bilateral lung transplantation is an effective option with reasonable short-term outcomes for these patients (Hawkins, 2021).

A systematic review of 51 prospective studies examined primary or secondary patient-important outcomes. The outcomes considered were mortality, pain, physical function, pulmonary, gastrointestinal, neuropsychological, cardiac, sleep or sexual symptoms, and quality of life. Mortality was the most frequently reported patient-important outcome (29.4% of studies), and quality of life was studied in 12%. The authors concluded outcomes other than mortality were insufficiently considered in lung transplantation studies.

We updated the references and modified the coverage based on new guidance and added an appendix of risk factor considerations from an updated consensus statement by the International Society for Heart and Lung Transplantation (Leard, 2021).

In 2023, we added four systematic reviews and meta-analyses to the policy with no policy changes warranted.

Two analyses provided mixed results regarding the outcomes of coronary revascularization performed before or during lung transplantation. In one analysis of 12 studies, the pooled mortality rates at one, three, and five years suggested significantly inferior survival in recipients with a prior history of coronary artery bypass grafting ($P < .00001$, $P = .0003$, $P = .008$, respectively) (Fialka, 2022). Another analysis of seven studies found no difference in mortality at one, three, and five years (overall hazard ratio = 1.02, 95% confidence interval = 0.80 to 1.31, $P = .99$) or hospital length of stay (standardized mean difference = 0.32, 95% confidence interval = -0.91 to 1.55) among participants who underwent lung transplantation with or without concomitant cardiac surgery. The group undergoing concomitant surgery experienced higher postoperative complication rates (Meng, 2022). For lung transplantation candidates who present with coronary artery disease, the optimum revascularization strategy has not been determined. Coronary artery disease should be regarded as a relative, not absolute, contraindication.

Two systematic reviews provide evidence of improved health-related quality of life outcomes following lung transplantation. In participants with advanced-stage cystic fibrosis, one systematic review of ten generally low-quality studies ($n = 1,494$) found lung transplantation improved health-related quality of life for up to five years, and to levels comparable to the general population and non-waitlisted candidates, even as medical management continues to evolve (Raguragavan, 2023a). The second analysis of ten studies ($n = 1,916$) found bilateral lung transplantation recipients reported significantly greater scores in both the physical and mental health domains of health-related quality of life beyond one-year post-lung transplantation, and recipients maintained these gains over the long-term compared to single lung transplantation recipients (Raguragavan, 2023b).

In 2024, we added some detail from the Organ Procurement and Transplantation Network about a new patient selection criterion for lung transplant patients in the background section (Organ Procurement and Transplantation Network, 2023). No other relevant literature was found and no policy changes were warranted.

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On October 11, 2024, we searched PubMed and the databases of the Cochrane Library, the U.K. National Health Services Centre for Reviews and Dissemination, the Agency for Healthcare Research and Quality, and the Centers for Medicare & Medicaid Services. Search terms were “lung transplantation (MeSH),” “heart lung transplantation (MeSH),” “lung transplantation,” and “pulmonary arterial hypertension.” We included the best available evidence according to established evidence hierarchies (typically systematic reviews, meta-analyses, and full economic analyses, where available) and professional guidelines based on such evidence and clinical expertise.

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Policy updates

10/2015: initial review date and clinical policy effective date: 1/2016

10/2016: Policy references updated.

10/2017: Policy references updated.

10/2018: Policy references updated.

10/2019: Policy references updated. Policy ID changed to CCP.1202.

10/2020: Policy references updated.

11/2021: Policy references updated.

11/2022: Policy references updated. Coverage modified.

11/2023: Policy references updated.

11/2024: Policy references updated.

Appendix

Table 1. International Society for Heart and Lung Transplantation factors associated with high or substantially increased risk.

Members with these conditions may be considered in centers with expertise specific to the condition. There may be insufficient data supporting transplantation in members with these risk factors, or currently available data suggest substantially increased risk, and further research is needed to better inform future recommendations. Presence of more than one of these risk factors may be multiplicative in terms of increasing risk of adverse outcomes. Modifiable conditions should be optimized when possible.

1. Age > 70 years.
2. Severe coronary artery disease that requires coronary artery bypass grafting at transplant.
3. Reduced left ventricular ejection fraction < 40%.
4. Significant cerebrovascular disease.
5. Severe esophageal dysmotility.
6. Untreatable hematologic disorders including bleeding diathesis, thrombophilia, or severe bone marrow dysfunction.
7. Body mass index > 35 kg/m².
8. Body mass index < 16 kg/m².
9. Limited functional status with potential for post-transplant rehabilitation.
10. Psychiatric, psychological, or cognitive conditions with potential to interfere with medical adherence without sufficient support systems.
11. Unreliable support system or caregiving plan.

12. Lack of understanding of disease and / or transplant despite teaching.
13. *Mycobacterium abscessus* infection.
14. *Lomentospora prolificans* infection.
15. *Burkholderia cenocepacia* or *gladioli* infection.
16. Hepatitis B or C infection with detectable viral load and liver fibrosis.
17. Chest wall or spinal deformity expected to cause restriction after transplant.
18. Extracorporeal life support.
19. Retransplant < one year following initial lung transplant.
20. Retransplant for restrictive chronic lung allograft dysfunction.
21. Retransplant for antibody mediated rejection as etiology for chronic lung allograft dysfunction.

Source: Leard (2021).

Table 2. Additional International Society for Heart and Lung Transplantation risk factors.

These risk factors have unfavorable implications for short- or long-term outcomes after lung transplantation. While members with these risk factors may be acceptable for lung transplantation programs to consider, multiple risk factors together may increase risk for adverse lung transplant outcomes.

1. Age 65-70 years.
2. Glomerular filtration rate 40-60 mL/min/1.73m².
3. Mild to moderate coronary artery disease.
4. Severe coronary artery disease that can be revascularized via percutaneous coronary intervention prior to transplant.
5. Member with prior coronary artery bypass grafting.
6. Reduced left ventricular ejection fraction 40-50%.
7. Peripheral vascular disease.
8. Connective tissue diseases (scleroderma, lupus, inflammatory myopathies).
9. Severe gastroesophageal reflux disease.
10. Esophageal dysmotility.
11. Thrombocytopenia, leukopenia, or anemia with high likelihood of persistence after transplant.
12. Osteoporosis.
13. Body mass index 30-34.9 kg/m².
14. Body mass index 16-17 kg/m².
15. Frailty.
16. Hypoalbuminemia.
17. Poorly controlled diabetes.
18. Edible marijuana use.

19. *Scedosporium apiospermum* infection.
20. Human immunodeficiency virus infection with undetectable viral load.
21. Previous thoracic surgery.
22. Prior pleurodesis.
23. Mechanical ventilation.
24. Retransplant > one year for obstructive chronic lung allograft dysfunction.

Source: Leard (2021).