Transcatheter pulmonary valve implantation (TPVI), may be considered **medically necessary** for patients with congenital heart disease and current right ventricular outflow tract obstruction (RVOT) or regurgitation including one or more of the following indications:

- Individuals with right ventricle-to-pulmonary artery conduit with or without bioprosthetic valve with at least moderate pulmonic regurgitation
- Individuals with native or patched RVOT with at least moderate pulmonic regurgitation
- Individuals with right ventricle-to-pulmonary artery conduit with or without bioprosthetic valve with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg)
- Individuals with native or patched RVOT with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg)

Transcatheter pulmonary valve implantation is considered **investigational** for all other indications.

**Policy Guidelines**

**Coding**
The following category I CPT code for this procedure:

- **33477**: Transcatheter pulmonary valve implantation, percutaneous approach, including pre-stenting of the valve delivery site, when performed

**Description**

Transcatheter pulmonary valve implantation (TPVI) is a less invasive alternative to open surgical pulmonary valve replacement or reconstruction for right ventricular outflow tract (RVOT) obstruction. Percutaneous pulmonary valve replacement may be indicated for congenital pulmonary stenosis. Pulmonary stenosis or regurgitation in a patient with congenital heart disease (CHD) who has previously undergone RVOT surgery are additional indications. Patients with prior CHD repair are at risk of needing repeated reconstruction procedures.

**Related Policies**

- N/A

**Benefit Application**

Benefit determinations should be based in all cases on the applicable contract language. To the extent there are any conflicts between these guidelines and the contract language, the contract language will control. Please refer to the member’s contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

Some state or federal mandates (e.g., Federal Employee Program [FEP]) prohibits plans from denying Food and Drug Administration (FDA)-approved technologies as investigational. In these instances, plans may have to consider the coverage eligibility of FDA-approved technologies on the basis of medical necessity alone.
Regulatory Status

Devices for transcatheter pulmonary valve implantation were initially cleared from marketing by the U.S. Food and Drug Administration (FDA) through the humanitarian device exemption (HDE) process or used off-label until approved by the FDA through the premarket approval (PMA) process between 2015 and 2016 (see Table 1).

<table>
<thead>
<tr>
<th>Device</th>
<th>Manufacturer</th>
<th>Date Approved</th>
<th>PMA No.</th>
<th>Indications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Melody® Transcatheter Pulmonary Valve (TPV)</td>
<td>Medtronic</td>
<td>Jan 2010</td>
<td>H080002 (HDE)</td>
<td>Pulmonary valve replacement for pediatric and adult patients with a dysfunctional, noncompliant RVOT conduit</td>
</tr>
<tr>
<td>Melody TPV</td>
<td>Medtronic</td>
<td>Jan 2015</td>
<td>P140017</td>
<td>Pulmonary valve replacement for pediatric and adult patients with a dysfunctional, noncompliant RVOT conduit</td>
</tr>
<tr>
<td>Melody TPV</td>
<td>Medtronic</td>
<td>Feb 2017</td>
<td>P140017/S005</td>
<td>Valve-in-valve for patients with a dysfunctional surgical bioprosthetic pulmonary valve</td>
</tr>
<tr>
<td>SAPIEN XT™ Transcatheter Heart Valve (pulmonic)</td>
<td>Edwards Lifesciences</td>
<td>Feb 2016</td>
<td>P130009/S037</td>
<td>Pulmonary valve replacement for pediatric and adult patients with a dysfunctional, noncompliant RVOT conduit</td>
</tr>
</tbody>
</table>

HDE: humanitarian device exemption; PMA: premarket approval; RVOT: right ventricular outflow tract.

The Melody® Transcatheter Pulmonary Valve (TPV) and the Ensemble® Transcatheter Valve Delivery System are used together for percutaneous replacement of a dysfunctional pulmonary valve. The Melody® valve consists of a section of bovine jugular vein with an intact native venous valve. The valve and surrounding tissue are sutured within a platinum-iridium stent scaffolding. The transcatheter delivery system consists of a balloon-in-balloon catheter with a retractable sheath and distal cup into which the valve is placed. The procedure is performed on a beating heart without the use of cardiopulmonary bypass.

The Melody® valve is first crimped to fit into the delivery system. It is introduced through the femoral vein and advanced into the right side of the heart and put into place at the site of the pulmonary valve. The inner balloon is inflated to open the artificial valve, and then the outer balloon is inflated to position the valve into place.

In January 2010, the Melody® TPV and the Ensemble® Transcatheter Valve Delivery System (Medtronic) were approved by the FDA under the HDE program for use as an adjunct to surgery in the management of pediatric and adult patients with the following clinical conditions:
- Existence of a full (circumferential) RVOT conduit that is 16 mm or greater in diameter when originally implanted, and
- Dysfunctional RVOT conduits with clinical indication for intervention, and either:
  - regurgitation: moderate-to-severe regurgitation, or
  - stenosis: mean RVOT gradient ≥35 mm Hg.

On January 27, 2015, approval of the Melody® system was amended to a PMA because the FDA determined that the device represented a breakthrough technology. The PMA was based, in part, on 2 prospective clinical studies, the Melody TPV Long-term Follow-up Post Approval Study and the Melody TPV New Enrollment Post Approval Study.

On February 24, 2017, approval of the Melody® system was expanded to include patients with a dysfunctional surgical bioprosthetic valve (valve-in-valve).
The Edwards SAPIEN XT™ Transcatheter Heart Valve (Pulmonic) (Edwards Lifesciences) is composed of a stainless steel frame with bovine pericardial tissue leaflets and available in 23- and 26-mm sizes. It includes a delivery accessories system. On February 29, 2016, it was approved by the FDA as a supplement “for use in pediatric and adult patients with a dysfunctional, noncompliant Right Ventricular Outflow Tract (RVOT) conduit with a clinical indication for intervention and:

- pulmonary regurgitation ≥ moderate and/or
- mean RVOT gradient ≥ 35 mmHg.”

The approval for the pulmonic valve indication is a supplement to the 2014 PMA for use of the Edwards SAPIEN XT™ Transcatheter Heart Valve System for relief of aortic stenosis in patients with symptomatic heart disease due to severe native calcific aortic stenosis and who are judged by a heart team, including a cardiac surgeon, to be at high or greater risk for open surgical therapy (i.e., Society of Thoracic Surgeons operative risk score ≥8% or at a ≥15% risk of mortality at 30 days).

FDA product code: NPV.

**Rationale**

**Background**

**Congenital Heart Disease**

Congenital heart disease, including tetralogy of Fallot, pulmonary atresia, and transposition of the great arteries, is generally treated by surgical repair at an early age. This involves reconstruction of the right ventricular outflow tract (RVOT) and pulmonary valve using a surgical homograft or a bovine-derived valved conduit. These repairs are prone to development of pulmonary stenosis or regurgitation over long periods of follow-up.

Because individuals with surgically corrected congenital heart disease repair are living into adulthood, RVOT dysfunction following initial repair has become more common. Calcification of the RVOT conduit can lead to pulmonary stenosis, while aneurysmal dilatation can result in pulmonary regurgitation. RVOT dysfunction can lead to decreased exercise tolerance, potentially fatal arrhythmias, and/or irreversible right ventricular dysfunction.

**Treatment**

Interventions for RVOT dysfunction often require numerous repeat open heart procedures for patients who live into adulthood. Treatment options for pulmonary stenosis are open surgery with valve replacement, balloon dilatation, or percutaneous stenting. Interventions for pulmonary regurgitation are primarily surgical, either reconstruction of the RVOT conduit or replacement of the pulmonary valve. The optimal timing of these interventions is not well understood.

Transcatheter pulmonary valve replacement offers a less invasive treatment option for patients with prior surgery for congenital heart disease and RVOT dysfunction. It is possible that a less invasive valve replacement technique could spare patients from multiple repeat open heart procedures over long periods of follow-up.

**Literature Review**

Evidence reviews assess the clinical evidence to determine whether the use of a technology improves the net health outcome. Broadly defined, health outcomes are length of life, quality of life, and ability to function— including benefits and harms. Every clinical condition has specific outcomes that are important to patients and to managing the course of that condition. Validated outcome measures are necessary to ascertain whether a condition improves or worsens; and whether the magnitude of that change is clinically significant. The net health outcome is a balance of benefits and harms.
To assess whether the evidence is sufficient to draw conclusions about the net health outcome of a technology, 2 domains are examined: the relevance and the quality and credibility. To be relevant, studies must represent one or more intended clinical use of the technology in the intended population and compare an effective and appropriate alternative at a comparable intensity. For some conditions, the alternative will be supportive care or surveillance. The quality and credibility of the evidence depend on study design and conduct, minimizing bias and confounding that can generate incorrect findings. The randomized controlled trial is preferred to assess efficacy; however, in some circumstances, nonrandomized studies may be adequate. Randomized controlled trials are rarely large enough or long enough to capture less common adverse events and long-term effects. Other types of studies can be used for these purposes and to assess generalizability to broader clinical populations and settings of clinical practice.

The published literature on transcatheter pulmonary valve implantation (TPVI) consists of the registration trials for U.S. Food and Drug Administration (FDA) approval, small case series, and cohort studies.

**Transcatheter Pulmonary Valve Implantation Devices and Uses Approved by the FDA**

**Review of Evidence**

**Melody Transcatheter Pulmonary Valve**

The multicenter U.S. Melody TPV trial was a prospective uncontrolled trial designed to assess the safety, procedural success, and short-term effectiveness of the Melody Transcatheter Pulmonary Valve (TPV). The Summary of Safety and Probable Benefit (SSPB) to support the approval of a humanitarian device exemption to market the Melody TPV was based on clinical data from 99 subjects who were catheterized for potential implantation with the TPV from January 2007, through December 2008, with expected follow-up and adverse event data on these subjects current through March 2009.3 Approved indications included right ventricular outflow tract (RVOT) dysfunction, defined as pulmonic regurgitation (moderate or greater) or pulmonic stenosis (mean gradient, ≥35 mm Hg). Also, a circumferential RVOT conduit should exist that is 16 mm or greater in diameter when originally implanted.

The investigators planned to follow 150 patients over 5 years. Eligibility criteria included a dysfunctional RVOT conduit or a dysfunctional bioprosthetic pulmonary valve, plus evidence of heart failure. For patients with New York Heart Association (NYHA) class I heart failure, a Doppler mean gradient of 40 mm Hg or greater or severe pulmonary regurgitation was required; for patients with NYHA class II to IV heart failure, a mean gradient of 35 mm Hg or greater or moderate pulmonary regurgitation was required. These inclusion criteria generally are indications for pulmonary valve replacement. The primary outcomes were defined as procedural success, adverse events from the procedure, and effectiveness, as measured by the proportion of patients with acceptable valve function at 6 months.

Trial results have been published in several reports.2,4,5 Short- and medium-term outcomes for 136 patients who underwent attempted TPVI were reported by McElhinney et al (2010).2 A total of 124 (91.2%) of 136 patients had successful implantation. In 12 patients, implantation was not possible due to anatomic or other intraprocedural findings. One (0.7%) death occurred as a result of the procedure, and serious adverse events occurred in 8 (6%) of 136 patients. Adverse events included coronary artery dissection, conduit rupture/tear, wide complex tachycardia, respiratory failure, femoral vein thrombosis, and perforation of the pulmonary artery.

Ninety-four patients with successful implantation had reached the 6-month follow-up at the time of publication. Acceptable valve function, defined as mild pulmonary regurgitation or less on echocardiography, was present in more than 90% of patients. Right ventricular (RV) pressure and RVOT gradient improved following the procedure, and 71 (75.5%) of 94 were in NYHA class I heart failure at 6 months. During follow-up, stent fractures were diagnosed in 25 (20.2%) of 124 patients, and 9 (7.3%) of 124 required implantation of a second valve.
Cheatham et al (2015) reported on outcomes up to 7 years following TPVI for the 148 patients who received and were discharged with a TPV in the U.S. Melody TPV trial (of 171 patients enrolled). Of the 171 patients enrolled, 167 underwent catheterization, 150 had a Melody valve implanted, and 148 of those survived to discharge with the Melody valve in place. On echocardiogram at discharge, pulmonary regurgitation was absent/trivial or mild in 140 patients and 5 patients, respectively, which represented a significant improvement from baseline. Over a median follow-up of 4.5 years (range, 0.4-7.0 years), 4 deaths occurred. During the follow-up period, 32 patients required a reintervention on RVOT, 25 of which were TPV reinterventions. A total of 11 patients required Melody valve explantation. Among the 113 patients who were alive and free from reintervention at a median of 4.5 years postimplantation, the most recent RVOT gradient was unchanged from early after valve implantation. Functional outcomes generally improved during the study: before TPVI, 14% of patients were in NYHA class I and 17% were in class III or IV. At every postimplantation annual evaluation, at least 74% of patients were in class I and no more than 1% to 2% were in class III or IV.

A secondary publication (2012) from the U.S. Melody TPV trial focused on the change in exercise function following TPVI. Patients completed a standardized cardiopulmonary regimen 2 months before and 6 months after TPVI. Results of pre- and postexercise parameters were available for 94 to 114 patients, depending on the specific outcome. Numerous physiologic outcome measures were reported, with some showing a statistically significant change between the 2 time points; and others not. For example, there was a significant increase in the percent predicted maximal workload from 65.0% at baseline to 68.3% at follow-up (p<0.001) and a significant decrease in the ratio of minute ventilation to CO2 production from 30.8 at baseline to 29.1 at follow-up (p<0.001). In contrast, there were no significant changes in peak oxygen consumption or spirometric measures of pulmonary function. This trial reported modest benefits in exercise parameters for patients treated with TPVI. The results were limited by the lack of a control group and by a large number of patients who did not have completed exercise results available (approximately one-third of total).

The 2015 premarket approval (PMA) of the Melody TPV was based on the interim analysis and a retrospective pooling analysis of the 2 postapproval studies conditioned by the prior humanitarian device exemption. An additional supplemental dataset from the Melody TPV European and Canadian Post-Market Surveillance Study (PMSS) was included in the PMA.

Armstrong et al (2014) published 1-year follow-up results of the Melody TPV Long-term Follow-up Post Approval Study (PAS), a prospective study designed to evaluate the short-term hemodynamic changes following device implantation. The study used historical controls from the Melody pivotal investigational device exemption (IDE) trial described above to investigate whether the short-term effectiveness of the device was noninferior to results shown in the IDE trial. PAS enrolled 120 subjects, 101 of whom underwent attempted TPVI. Patient selection was based on the criteria used in the IDE trial but did not include the age (≥5 years of age) and weight (≥30 kg) limitations. Procedure-related significant adverse events occurred in 16 patients (13.3% of total cohort; 15.8% of those who had an attempted TPVI), the most common of which was a confined conduit tear. Procedural success occurred in 99 subjects (98% of those with an attempted TPVI). At 1-year follow-up, the proportion of patients in NYHA class I heart failure increased from 35% at baseline to 89%. Of the 99 patients implanted for at least 24 hours, 87 had acceptable TPV hemodynamic function confirmed at 6 months (96.7% of those with evaluable echocardiographic data, 87.9% of entire cohort) and 82 had acceptable TPV hemodynamic function at 1 year (94.3% of those with evaluable echocardiographic data, 82.8% of the entire cohort). Following the procedural period, serious device-related adverse events occurred in 8%, most commonly endocarditis (n=3 patients).

Gillespie et al (2015) evaluated results of TPVI after a Ross procedure in a retrospective review of pooled findings from the U.S. Melody TPV trial and PAS and an additional European registry, the manufacturer-sponsored Melody TPV PMSS conducted in Canada and Europe (NCT00688571). In the pooled sample (total N=358 patients), 67 (19%) had a prior Ross procedure. A Melody
valve was successfully implanted in 56 (84%) of 67 Ross patients who underwent catheterization with intent for TPVI. Six (9%) patients had symptomatic coronary artery compression after TPVI or did not undergo implantation due to the risk of compression. RV hemodynamics generally improved after TPVI, but RVOT reinterventions were required in 12 of 55 patients discharged from the implant hospitalization with the Melody valve in place.

The Melody TPV New Enrollment Study was intended to roll in the new patient enrollment study specified as a condition of approval for the Melody TPV HDE on January 25, 2010.9 This study used the protocol dated September 24, 2013, Version 2, included in H080002/S015. The study is a prospective, nonrandomized, multicenter, historically controlled clinical trial, designed to assess the postmarket performance of the Melody TPV in a representative population of providers and patients, with 5-year follow-up. The primary endpoint is freedom from TPV dysfunction, with a performance goal of 75% or greater at 6 months. Secondary end points include procedural success, serious procedural- and device-related adverse events, stent fracture, reintervention on the TPV, surgical replacement of the RVOT conduit, death (all-cause, procedure-related, and device-related), and NYHA classification.

The February 2017 approval of the Melody system expanded to include patients with a dysfunctional surgical bioprosthetic valve (valve-in-valve) is based on data pooled from 3 sources:

- Melody TPV Long-term Follow-up PAS: 8 patients
- Melody TPV New Enrollment PAS: 17 patients
- Real-World Data: 100 patients.

Of 125 patients pooled from the 3 studies listed above, 56.8% (71) patients were available for analysis at study completion, the 1-year postimplant visit. Baseline pooled subject median age was 22.0 years (range, 5.0-79 years), with 45.6% female and 54.4% male. Tetralogy of Fallot was the most common congenital heart disease diagnosis recorded in 72.8% of subjects, 66.4% of whom had pulmonary stenosis or atresia. There was no mortality for any cause, major stent fracture, occurrence of endocarditis, RVOT reoperation, or catheter reintervention among available patients at 1 year. Procedural failure as defined by more than trivial pulmonary regurgitation by angiography postimplant occurred in 10.1% (12/119) subjects. There were no device explants within 24 hours of implantation. The mean RVOT gradient was reduced from 29.5 mm Hg at baseline to 14.3 mm Hg at 1 year postimplantation. In this PMA, existing clinical data were not leveraged to support approval of a pediatric patient population. This submission included pediatric data to support the pediatric indication and no extrapolation was necessary.

Additional Noncomparative Studies
A number of publications have reported on series of patients treated with TPVI. Some of the larger series are detailed next.

Lurz et al (2008) reported on 163 patients who underwent attempted TPVI from 4 clinical centers in Europe.11 Eligibility for the procedure included elevated RV systolic pressure, increased RVOT dimensions, and either symptoms or evidence of severe RV dysfunction. Procedural success was achieved in 155 (95.1%) of 163 patients. Procedural complications occurred in 12 (7.4%) of 163, 8 of which were considered serious and 5 of which required open surgery. Median follow-up was 28.4 months. During follow-up, 4 (2.6%) of 155 patients died, and an additional 5 (3.2%) developed infective endocarditis. At 12-month follow-up, more than 90% of patients had absent or mild valve dysfunction as measured by echocardiography.

Eicken et al (2011) reported on 102 consecutive patients (mean age, 21.5 years) undergoing TPVI at 2 centers in Germany.12 Eligibility for the procedure included RVOT dysfunction with evidence of RV compromise or increased RV pressure. One (1.0%) death occurred as a result of compression of the left coronary artery. Two (2.0%) patients had evidence of stent fracture immediately postprocedure and 1 (1.0%) other patient developed infective endocarditis at 6-month follow-up. At a median follow-up of 357 days, there was a significant decrease in the
RVOT gradient, from a median of 36 to 15 mm Hg (p<0.001). However, there was no significant change in exercise capacity as measured by maximal oxygen uptake.

Other case series have reported on smaller numbers of patients, ranging from 7 to 64 patients. These series have generally reported similar results as the larger series, with high procedural success and relatively low rates of serious complications. The longest follow-up, reported by Borik et al (2015), evaluated 51 patients who underwent TPVI with the Melody valve at a single institution. Over a mean follow-up of 4.5 years (range, 0.9-6.9 years), freedom from any reintervention was 87% and 68% at 3 and 5 years, respectively, and freedom from surgery was 90% at 5 years. Overall, RV functional parameters did not change with longer follow-up.

An Australian prospective observational registry (2017) reported information accumulated on 17 patients implanted with Melody device between 2009 and 2016. Mixed valvular dysfunction was present in 7 (41%) patients and 11 (59%) had corrected tetralogy of Fallot. Device implantation was successful in all patients. Peak RVOT gradient was significantly reduced, and there was no significant regurgitation postprocedure. There was 1 (6%) major procedural adverse event and 2 (12%) major adverse events at last recorded follow-up. There were no patient deaths. Follow-up cardiac magnetic resonance imaging revealed a significant reduction in indexed RV end diastolic volume.

**Edwards Sapien XT Transcatheter Heart Valve (Pulmonic)**

Edwards Lifesciences, manufacturer of the SAPIEN THV, performed a clinical study to establish the safety and efficacy of its pulmonic implantation in patients with dysfunctional RVOT conduits in the United States under IDE G060242 (the COngenital Multicenter trial of Pulmonic vAlve regurgitation Studying the SAPIEN InterventIO NaL THV, COMPASSION trial). Data from this clinical study were the basis for the PMA decision for the pulmonary valve implantation indication.

Patients were treated between April 2008 and November 2014. The database supplement reflects data collected through March 2015 and includes 81 patients. There were 7 investigational sites.

This prospective, nonrandomized, multicenter clinical study assessed the safety and effectiveness of pulmonic implantation of the SAPIEN THV in patients with dysfunctional RVOT conduits requiring treatment for moderate or severe pulmonary regurgitation by transthoracic echocardiography (TTE) and/or RVOT conduit obstruction with a mean gradient of 35 mm Hg or higher by TTE. The SAPIEN THV, the first-generation valve of the SAPIEN device line, is no longer available for distribution. The valve sizes used in the COMPASSION trial included the 23- and 26-mm sizes, which were the only sizes available for the SAPIEN THV. The 29-mm valve size was not evaluated in the COMPASSION trial. Most data derived from patients who received the 23-mm THV size. Aortic experience with the 29-mm SAPIEN XT THV showed no significant difference in the long-term performance compared with the 23- and 26-mm sizes. Furthermore, no observed results suggested that the 29-mm valve size would perform worse than other available sizes in the pulmonic location.

Enrollment in the COMPASSION study was limited to patients who met the following inclusion criteria:

- **Weight ≥35 kg.**
- **In situ conduit size of 20 to 26 mm in diameter.**
- **Moderate or severe pulmonary regurgitation defined as ≥3+ pulmonary regurgitation by TTE or**
- **RVOT conduit obstruction with a mean gradient of ≥35 mm Hg by TTE.**
- **Symptomatic as evidenced by cardiopulmonary exercise testing.**
- **Catheterization was determined to be feasible by the treating physician.**
All patients were scheduled to return for follow-ups at day 1 postprocedure, discharge, 30 days, 6 months, 12 months, and annually after that for 5 years postoperatively. Baseline evaluation included TTE, x-ray, magnetic resonance imaging, or computed tomography, angiogram, and electroencephalograph. Assessment of NYHA class, magnetic resonance imaging or computed tomography, and angiogram were part of the 6-month evaluation.

The primary end point was freedom from the device- or procedure-related death and/or reintervention at 1 year. The secondary end points were:

- Freedom from major adverse cardiac and cerebrovascular events at 6 months. Major adverse cardiac and cerebrovascular events was defined as all-cause mortality, myocardial infarction, reintervention, vascular injury resulting in the need for an unplanned vascular intervention, stroke, and pulmonary embolism.
- Functional improvement at 6 months as defined by:
  - Improved valve hemodynamics as demonstrated via TTE:
    - Decrease in pulmonary regurgitation to mild or less for regurgitant lesions
    - Decrease in the mean pulmonary gradient to less than 30 mm Hg for stenotic lesions
    - Improvement in both pulmonary regurgitation and gradient (above) for mixed lesions.
  - Improvement of 1 or more NYHA functional classes from baseline for patients in NYHA functional classes > 2 at baseline.
  - Freedom from recurrent pulmonary stenosis.

Of 81 patients enrolled in the PMA study, 2 patients were screening failures, 9 patients did not receive the valve, another received the valve in a nontarget location. Therefore, 69 patients were available for analysis in the valve implant population at study completion.

The median duration of follow-up for the safety population was 3.04 years (range, 0-5.31 years). Males were 65.8% of the population, and 63.3% were at least 22 years of age. The primary indications for valve implantation were pulmonary stenosis (8.9%), pulmonary regurgitation (12.7%), and both stenosis and regurgitation (78.5%). The primary etiology requiring reconstruction of the RVOT and placement of a pulmonary conduit for the safety population was tetralogy of Fallot (42%).

The prespecified performance goal for the primary end point was 75%. The primary outcome was met by 100% of patients; there were 3 reintervention events. At 5 years, the primary outcome using a Kaplan-Meier estimate was 77.1%. Because there was no device- or procedure-related patient deaths at 5 years, the incidence of reinterventions solely contributed to the estimate.

Freedom from reintervention to 5 years for the valve implant population using a Kaplan-Meier estimate was reported by type of reintervention: (a) freedom from surgical pulmonic valve repair was 98.3% at 1 year and 91.8% at 5 years; (b) freedom from TPVI was 97.1% at 1 year and 85.8% at 5 years; (c) freedom from balloon valvuloplasty was 100% at 1 year and 93.7% at 5 years; and (d) freedom from other types of reintervention was 100% at 1 year and 97.9% at 5 years.

For secondary outcomes, freedom from major adverse cardiac and cerebrovascular events at 6 months in the valve implant population was 94.1%. Because 2 (2.5%) of 79 patients experienced a device migration early in the trial, the instructions for use were modified. No other device migrations subsequently occurred in the trial. Serious adverse events for RVOT conduit ruptures occurred in 5 (6.3%) of 79 patients. These 5 ruptures were related to balloon valvuloplasty or placement of a pre-stent; no ruptures occurred during placement of the SAPIEN THV. There was 1 neurologic event (not stroke), 1 thromboembolism, and 4 endocarditis events at the 1-year follow-up.
Adjunctive analyses of safety and effectiveness stratified by patients ages 21 years or younger at baseline vs patients ages 22 years or older at baseline were conducted. The COMPASSION study was not designed to investigate the differences in outcomes between age groups and, therefore, no statistical inferences can be made. The analysis of functional improvement outcomes by age group is summarized in Table 2.

Table 2. Edwards Sapien XT Transcatheter Heart Valve (Pulmonic) PMA Approval Study: Overall Functional Improvement by Age Group for Valve Implanted Population

<table>
<thead>
<tr>
<th>End Points</th>
<th>Age 21 or Younger (n=27)</th>
<th>Age 22 or Older (n=42)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall functional improvement</td>
<td>18/22 (85.7) 3/7 (42.9)</td>
<td>29/33 (87.9) 5/8 (62.5)</td>
</tr>
<tr>
<td>Improved valve function</td>
<td>19/19 (100.0) 5/5 (100.0)</td>
<td>28/30 (93.3) 6/6 (100.0)</td>
</tr>
<tr>
<td>Functional improvement in NYHA class</td>
<td>13/14 (92.9) 4/4 (100.0)</td>
<td>32/33 (97.0) 8/8 (100.0)</td>
</tr>
<tr>
<td>Freedom from recurrent pulmonary stenosis</td>
<td>18/19 (94.7) 5/9 (55.6)</td>
<td>31/31 (100.0) 7/10 (70.0)</td>
</tr>
<tr>
<td>Improved gradient</td>
<td>7/8 (87.5) 3/3 (100.0)</td>
<td>7/8 (87.5) 2/2 (100.0)</td>
</tr>
</tbody>
</table>


As listed in Table 2, overall functional improvement was defined by the following 4 categories: (a) improved valve function demonstrated by a decrease in pulmonary regurgitation to mild or less per TTE at visit for patients with moderate or more (>2) pulmonary regurgitation at baseline, (b) functional improvement from baseline of 1 or more NYHA functional classes at visit for patients with baseline NYHA functional class of 2 or higher, (c) freedom from recurrent pulmonary stenosis at visit, and (d) improved valve function demonstrated by a decrease in pulmonary stenosis mean gradient to less than 30 mm Hg for patients with pulmonary stenosis mean gradient greater than 30 mm Hg at baseline. Patients with mild or less (≤2+) pulmonary regurgitation at baseline only use categories b, c, and d to determine overall functional improvement. Patients with an NYHA functional class of less than 2 at baseline only use categories a, c, and d to determine overall functional improvement. Patients treated for indications other than pulmonary stenosis only use categories a, b, and d for overall functional improvement. Patients with pulmonary stenosis mean gradient less than 30 mm Hg at baseline only use categories a, b, and c for overall functional improvement.

A small number of retrospective, comparative studies have compared outcomes of the Edwards SAPIEN and the Melody valves. Boshoff et al (2013) described the off-label uses in 21 patients treated with the Melody valve and 2 patients treated with the Edwards SAPIEN pulmonic valve. Use has included native RVOT obstruction, in conduits smaller than the FDA-labeled indications, and large RVOT with a dynamic outflow aneurysm. No deaths or major procedural complications were reported for these patients. Clinical outcomes data were lacking or very limited in this publication.

Faza et al (2013) reported on 20 patients who underwent successful implantation of the Edwards SAPIEN pulmonic valve at a single clinical center. There were no periprocedural deaths, and all but 1 patient had no or trivial pulmonary regurgitation on latest follow-up. A comparison of hemodynamic parameters in these 20 patients was made with 13 patients treated with the Melody valve. Immediately postprocedure, transvalvular gradients were similar between groups. At last follow-up, mean residual transvalvular gradient was higher for patients receiving the SAPIEN valve (18.4 mm Hg vs 11.2 mm Hg, p=0.016), but this difference disappeared when patients were matched for length of follow-up.

A few other small case series has assessed on the use of the Edwards SAPIEN pulmonic valve for RVOT obstruction. For example, Kenny et al (2011) reported on a phase 1 multicenter study of the SAPIEN valve in 36 patients from 4 clinical centers. Procedural success was reported in 97% of patients. Procedural complications occurred in 19% (7/36) of patients, including valve migration (n=3), pulmonary hemorrhage (n=2), ventricular fibrillation (n=1), and stent migration (n=1). At 6-month follow-up, there were no deaths, and 75% (27/36) of patients
were in NYHA class I, compared with 14% at baseline. Freedom from reintervention at 6 months was 97%.

Section Summary: Approved Transcatheter Pulmonary Valve Implantation Devices and Uses
The evidence for the use of TPVI with the Melody valve or the SAPIEN XT systems consists of the prospective, interventional, noncomparative pivotal studies on which each device’s FDA approval was based, along with postapproval registry studies and additional case series. Overall, the evidence would suggest that TPVI is associated with high rates of short-term technical success and improvements in heart failure-related symptoms and hemodynamic parameters. Studies with postprocedure follow-up extending to a maximum of 7 years have suggested that the functional and hemodynamic improvements are durable, with a number (20%-30%) requiring reintervention on the pulmonary valve.

Non-FDA-Approved Transcatheter Pulmonary Valve Implantation Devices and Uses
A variety of potential off-label uses of TPVI have been reported in the literature, including the use of approved devices for non-FDA-approved indications.

Data from the Valve-in-Valve International Database multicenter registry have been evaluated for the off-label use of transcatheter aortic and TPVI prostheses for tricuspid valve-in-valve implantation.30 One hundred fifty of 156 patients in the registry had successful tricuspid valve-in-valve with a Melody (n=93) or a SAPIEN (n=57) valve. During a median 13.3-month follow-up, 22 (15%) patients died, all with NYHA class III or IV. There were 10 (6.6%) tricuspid valve re-interventions and 3 (2%) other patients who had significant recurrent dysfunction of the valve. Preintervention, 71% of patients were in NYHA class III or IV; at follow-up, 77% of surviving patients were in NYHA class I or II (p<0.001).

A few case series have been on use of the Melody valve in patients with clinical characteristics not corresponding to FDA-approved indications.24,31 These indications have included the use of valves in positions other than pulmonic, patients with conduit sizes inconsistent with FDA indications, and patients with prior congenital heart repair surgery not involving the construction of an RVOT conduit. In general, these case series have reported high rates of procedural success with low rates of periprocedural complications, but longer term outcomes are lacking.

Although most studies have evaluated the use of TPVI in patients with a constructed RVOT conduit, a few have evaluated TPVI with either the Melody valve or the Edwards SAPIEN THV in a native RVOT or RVOT without a circumferential conduit. Meadows et al (2014) reported on results from a retrospective, 5-center review of patients who underwent TVP placement in a non-conduit RVOT, with the native tissue comprising at least part of the circumference.32 Thirty-one patients were included, with indications for RVOT intervention including primarily valvular insufficiency in 14 (45%), obstruction in 3 (10%), and mixed obstruction and insufficiency in 14 (45%). TPVI was successful in all patients, with serious complications in 2 (6%). At a median follow-up of 15 months (range, 1 month to 3.8 years), all patients were alive, and none reporting greater than mild pulmonary regurgitation. Among the 19 patients with adequate imaging at follow-up, 6 (32%) had evidence of stent fracture. Three patients were treated for endocarditis or bloodstream infection. Malekzadeh-Milani (2014) reported on outcomes for 34 patients with a native or patched noncircular RVOT who underwent Melody TPV insertion at a single center.33 The procedure was technically successful in all patients, with early complications occurring in 8.8% At a mean follow-up of 2.6 years, no patients had stent fracture or migration, and 32 (94.1%) of 34 had no or trivial pulmonary regurgitation.

Several other small case series, by Demkow et al (2014; N=10) and Odemis et al (2013; N=7), have reported on the use of the Edwards SAPIEN pulmonary valve for noncircumferential RVOT patch and large-diameter conduits, respectively. The authors reported high rates of successful valve implantation, but no long-term follow-up.
Adverse Events

In addition to the adverse events reported in the case series, several publications have focused on adverse events following TPVI. The FDA reviewed results from the U.S. Melody TPV trial as part of its approval process and reported data on complications from the procedure. At that time, data were available for 99 patients enrolled between January 2007 and December 2008. Ninety patients were deemed suitable for implantation following catheterization, and 87 of them had successful implantation. There was 1 (1.1%) procedure-related death. Table 3 is adapted from the FDA summary of safety and probable benefit.

Table 3. Device-Related Adverse Events (N=89 Subjects implanted > 24 hours)

<table>
<thead>
<tr>
<th>Adverse Events</th>
<th>Subjects With Event, n (%)</th>
<th>Freedom From Event at 12 Months (SE), %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stent fracture (all)</td>
<td>16 (18)</td>
<td>77.1 (7.5)</td>
</tr>
<tr>
<td>Minora</td>
<td>11 (12)</td>
<td>84.1 (6.7)</td>
</tr>
<tr>
<td>Majora</td>
<td>5 (6)</td>
<td>90.6 (5.2)</td>
</tr>
<tr>
<td>Valve stenosis</td>
<td>6 (7)</td>
<td>90.5 (4.8)</td>
</tr>
<tr>
<td>Worsening tricuspid regurgitation</td>
<td>1 (1)</td>
<td>100 (-)</td>
</tr>
<tr>
<td>Reinterventionb</td>
<td>6 (7)</td>
<td>93.5 (4.3)</td>
</tr>
<tr>
<td>Reoperation</td>
<td>1 (1)</td>
<td>98.6 (2.2)</td>
</tr>
</tbody>
</table>

a Stent fractures that did not require intervention were defined as minor; those that required reintervention were defined as major.

b Reinterventions were balloon angioplasty in 1 patient; repeat implantation of a second transcatheter pulmonary valve in 5 patients.

Sixty-four patients in the FDA analysis reached the 6-month follow-up. Of them, 56 (87.5%) had acceptable hemodynamic valve function by Doppler echocardiography. At 6 months, approximately 75% of patients were in NYHA class I and 25% were in NYHA class II. Pulmonary regurgitation that was mild or worse was present in 6.2% of patients.

Another publication focusing on adverse events from the U.S. Melody TPV trial was published in 2011.35 This report assessed adverse events at a median follow-up of 30 months in 150 patients. Stent fracture occurred in 26% (39/150) of patients. The estimated freedom from stent fracture was 77% at 14 months and 60% at 39 months. Freedom from reinterventions for all patients was estimated to be 86% at 27 months, and freedom from reinterventions for patients with stent fracture was estimated at 49% at 2 years.

McElhinney et al (2013) reported on rates of infective endocarditis from 3 prospective case series enrolling 311 patients followed for a median of 2.5 years.36 Sixteen (5.1%) patients were diagnosed with endocarditis at any location, and 6 (1.9%) patients had endocarditis at the pulmonic valve location. This corresponded to an annualized rate of pulmonic valve endocarditis of 0.88% per patient-year. Malekzadeh-Milani et al (2014) evaluated patients with right-sided infective endocarditis at a single center to compare endocarditis rates in patients who had TPVs with those who had surgically paced pulmonary valves.37 Thirty-one patients with right-sided endocarditis and pulmonary valve implantation for congenital heart disease were included. Rates of endocarditis were 1.2 and 3.9 cases/100 person-years in patients with surgically implanted valves and TPVs, respectively (p=0.03).

Boudjemline et al (2016) conducted a prospective observational study to evaluate predictors of conduit rupture during the preparation of the RVOT for TPVI in a cohort of patients older than age 5 years with RVOT obstruction, pulmonary regurgitation, or mixed lesions, who underwent transcatheter therapies, including balloon dilatation, bare metal stent placement, or TPV placement.38 Ninety-nine patients were included, 56 of whom were adults. Of the total cohort, 83.8% underwent Melody TPV. Conduit rupture occurred in 9 (9.09%) patients. In 2 of the 9 patients, conduit rupture was angiographically obvious and severe with extension, causing hemodynamic instability. All conduit ruptures occurred during balloon dilatation and occurred in patients with RVOT obstruction. Heavy calcification and the presence of a homograft were associated with conduit rupture risk.
Coronary artery compression during balloon angioplasty or stent placement in the RVOT conduit is considered a relative contraindication to TPV placement. Several studies have evaluated the incidence of coronary artery compression. Morray et al (2013) reported on the incidence of coronary artery compression in a 4-center series of 404 patients who underwent attempted TPVI. Three hundred forty-three (85%) patients underwent TPVI, and 21 (5%) patients had evidence of coronary artery compression. Most (n=19) patients with coronary artery compression did not undergo TPV placement. Using the same cohort reported in the Boudjemline study, Fraisse et al (2014) reported on the incidence, diagnosis, and outcome of coronary compression among patients treated with transcatheter RVOT interventions for RVOT obstruction, pulmonary regurgitation, or mixed lesions. All patients underwent balloon dilatation and coronary assessment with angiography, which was followed by TPV placement if RVOT dysfunction was ongoing. Of 100 patients evaluated, 83% had implantation of a Melody TPV. Coronary artery compression occurred in 6 cases, all of which could be diagnosed by selective coronary angiogram and/or aortic root angiogram during balloon dilation of the RVOT. No specific risk factors for coronary artery compression were identified.

Van Dijck et al (2015) compared rates of infective endocarditis between TPVs and surgically implanted pulmonary valves in a retrospective, single-center study that included 677 patients (738 conduits). Patients who underwent procedures from 1989 to 2013 were included. A total of 107 Melody conduits were implanted in 107 patients. A total of 577 pulmonary valves cryopreserved homografts were implanted in 517 patients, and 54 Contegra grafts were implanted in 53 patients. Freedom from infective endocarditis at 5 years using Kaplan-Meier analysis was 85%, 88%, and 99% for patients with Melody conduits, Contegra grafts, or cryopreserved homografts, respectively.

Malekzadeh-Milani et al (2015) reported on the incidence of infective endocarditis among 86 prospectively enrolled consecutive patients who underwent TPVI with the Melody valve. Over a mean follow-up of 23.6 months (range, 2.6-28.3 months) after Melody implantation, 5 patients developed infective endocarditis (5.8%; 95% confidence interval, 0.9% to 10.7%). Factors related to demographics, conduit type, procedural success, residual gradient, and duration of Melody valve implantation did not differ significantly between patients who did or did not develop infective endocarditis. Patients with infective endocarditis were more likely to have undergone invasive procedures after TPVI without antibiotic prophylaxis (odds ratio, 13.69; 95% confidence interval, 1.98 to 94.52; p=0.014), and aspirin use was preventive for infective endocarditis (relative risk, 20.1; 95% confidence interval, 3.34 to 120.9; p=0.001), although confidence intervals around risk estimates for both factors were wide.

A 2017 retrospective review of 25 patients undergoing Melody TPVI and 178 surgical pulmonic valve surgeries (bioprostheses and homografts) was reported from New Zealand from October 2009 to May 2015. Four (16%) implant patients experienced endocarditis. Two patients presented with life-threatening endocarditis and obstructive vegetation at 14 and 26 months postimplant, respectively. Two additional patients presented with subacute endocarditis at 5.5 years postimplant. At a median follow-up of 2.9 years, 4 (2%) patients had developed endocarditis in the surgical group.

**Summary of Evidence**

For individuals who have a history of CHD and current RVOT obstruction who receive TPVI with a U.S. Food and Drug Administration (FDA) approved device and indication, the evidence includes prospective, interventionai, noncomparative studies, and multiple prospective and retrospective case or cohort series. Relevant outcomes are overall survival, symptoms, functional outcomes, quality of life, hospitalizations, and treatment-related mortality and morbidity. Results of the case series have indicated that there is a high rate of procedural success and low procedural mortality, although the rates of serious procedural adverse events reported ranged from 3.0% to 7.4%. Most valves have demonstrated competent functioning by Doppler echocardiography at 6- to 12-month follow-ups, but complications (e.g., stent fractures, need for...
reinterventions) were reported in an FDA analysis at rates of 18% and 7%, respectively. Other publications with longer follow-up have reported stent fractures in up to 26% of patients; however, most stent fractures did not require reintervention. Studies with follow-up extending to a maximum of 7 years postprocedure have suggested that the functional and hemodynamic improvements are durable, but a relatively high proportion of patients (20%-30%) have required reintervention on the pulmonary valve. No comparative studies were identified, and there is no direct evidence that TPVI reduces future open heart procedures. The evidence is insufficient to determine the effects of the technology on health outcomes.

For individuals who have a history of CHD and current RVOT obstruction who receive TPVI with a non-FDA-approved device or indication, the evidence includes case series. Relevant outcomes are overall survival, symptoms, functional outcomes, quality of life, hospitalizations, and treatment-related mortality and morbidity. There is limited evidence on the off-label use of TPVI including the use of a non-FDA-approved valve or use of an approved valve for a non-FDA-approved indication. The published case series enrolled relatively few patients and are heterogeneous regarding devices used and indications for TPVI. The evidence is insufficient to determine the effects of the technology on health outcomes.

**Clinical Input**

**Objective**

In 2018, clinical input was sought to help determine whether the use of transcatheter pulmonary valve implantation for individuals with congenital heart disease and current right ventricular outflow tract (RVOT) obstruction or regurgitation would provide a clinically meaningful improvement in the net health outcome and whether its use is consistent with generally accepted medical practice.

**Respondents**

Clinical input was provided by the following specialty societies:

- American College of Cardiology (ACC) and Society for Cardiovascular Angiography and Interventions (SCAI)

\(^a\) Indicates that conflicts of interest related to the topic where clinical input is being sought were identified by this respondent (see Appendix).

Clinical input provided by the specialty society at an aggregate level is attributed to the specialty society. Clinical input provided by a physician member designated by the specialty society or health system is attributed to the individual physician and is not a statement from the specialty society or health system. Specialty society and physician respondents participating in the Evidence Street® clinical input process provide a review, input, and feedback on topics being evaluated by Evidence Street. However, participation in the clinical input process by a special society and/or physician member designated by the specialty society or health system does not imply an endorsement or explicit agreement with the Evidence Opinion published by BCBSA or any Blue Plan.
Clinical Input Responses

Figure 1:


<table>
<thead>
<tr>
<th>Clinical Indication</th>
<th>Respondent</th>
<th>Fac or No</th>
<th>YES</th>
<th>Intermediate</th>
<th>NO</th>
</tr>
</thead>
<tbody>
<tr>
<td>TPVI using Melody® Transcatheter Pulmonary Valve (TPVI)</td>
<td>ACC / SCAI</td>
<td>YES</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RV-PA conduit with or without bioprosthetic valve with at least moderate pulmonary regurgitation</td>
<td>ACC / SCAI</td>
<td>YES</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Native or patched RVOT with at least moderate pulmonary regurgitation</td>
<td>ACC / SCAI</td>
<td>YES</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glue: RV-PA conduit with or without bioprosthetic valve with pulmonary stenosis (mean RVOT gradient at least 30 mm Hg)</td>
<td>ACC / SCAI</td>
<td>YES</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Native or patched RVOT with pulmonary stenosis (mean RVOT gradient at least 35 mm Hg)</td>
<td>ACC / SCAI</td>
<td>YES</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>


Table 1: Clinical Input Responses

- Indicates that conflicts of interest related to the topic where clinical input is being sought were identified by this respondent (see Appendix).
- The choice of SAPIEN transcatheter heart valve may be based on clinical judgment such as size needed for patient anatomy.

Additional Comments

- "TPVI is a well accepted and FDA-approved therapy for RV-PA conduit and bioprosthetic valve (BPV) regurgitation, regardless of the surgical risk, because of 5-7 year follow-up data from multiple prospective studies showing excellent results and durability." (ACC / SCAI)
- "While there are no transcatheter valves FDA-approved for use in a native or patched RVOT, this therapy is considered a reasonable alternative to surgical pulmonary valve replacement (SPVR), regardless of surgical risk." (ACC / SCAI)
- "Transcatheter pulmonary valve placement in the setting of stenosis has been reported to have excellent procedural success and has been noted to have durable improvements in patient status. Perhaps one argument against transcatheter pulmonary valve placement when compared to surgical intervention is the published rate of reintervention with TPVR. This is an effect of the pre-stent era and in this population of stenotic conduits, reinterventions were often a result of stent fracture. Additionally, previously reported risk in this population of a stenotic conduit includes conduit rupture, now able to be adequately treated with covered stents now available following the COAST and PARCS trials in nearly all cases. Eliminating at least one stenotomy, often more, in patients that will require several reentry stenotomies most arguably can be lifesaving." (ACC / SCAI)
- "The stenotic native RVOT carries fewer risks for transcatheter valve placement compared to dilated RVOT and severely stenotic conduit cohort. This group additionally has highly variable and dynamic RVOT anatomy which can make patient selection key in success." (ACC / SCAI)

See Appendix 1 and 2 for details.
Supplemental Information

Clinical Input From Physician Specialty Societies and Academic Medical Centers

While the various physician specialty societies and academic medical centers may collaborate with and make recommendations during this process, through the provision of appropriate reviewers, input received does not represent an endorsement or position statement by the physician specialty societies or academic medical centers, unless otherwise noted.

2018 Input

In response to requests from Blue Cross Blue Shield Association, clinical input on the use of transcatheater pulmonary valve implantation (TPVI) was received from 2 specialty society-level respondents in 2018. The combined clinical input response incorporated input from a panel including physicians affiliated with academic medical centers.

Based on the evidence and independent clinical input, the clinical input supports that the following indications provide a clinically meaningful improvement in the net health outcome and are consistent with generally accepted medical practice:

- Use of TPVI for individuals with right ventricle-to-pulmonary artery conduit with or without bioprosthetic valve with at least moderate pulmonic regurgitation;
- Use of TPVI for individuals with native or patched right ventricular outflow tract (RVOT) with at least moderate pulmonic regurgitation;
- Use of TPVI for individuals with right ventricle-to-pulmonary artery conduit with or without bioprosthetic valve with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg); or
- Use of TPVI for individuals with native or patched RVOT with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg).

2011 Input

In response to requests from Blue Cross Blue Shield Association, input was received from 6 academic medical centers in 2011. Overall response to whether TPVI was investigational was mixed, with 2 of 5 reviewers agreeing with the investigational status, and 3 reviewers indicating partial support. Most reviewers (4/5) indicated that there is a subpopulation of patients who are high risk for surgery or who are not candidates for surgery, for whom there are no other available options. These reviewers suggested TPVI was a viable alternative that offered potential benefit for these patients.

Practice Guidelines and Position Statements

Society for Cardiovascular Angiography and Interventions et al

The Society for Cardiovascular Angiography and Interventions, American Association for Thoracic Surgery, American College of Cardiology (ACC), and the Society of Thoracic Surgeons (2015) published a consensus-based report on operator and institutional requirements for transcatheter pulmonary valve implantation (TPVI). Recommendations to qualify for a TPVI program included 150 catheterizations per year, association with a surgical program, submission of all cases to a national registry, and, for patients, 80% freedom from reintervention at 1 year.

American Heart Association and American College of Cardiology

The 2014 American Heart Association (AHA) and ACC guidelines on the management of patients with valvular disease and the 2017 AHA and ACC focused update do not make specific recommendations on the treatment of primary pulmonary valve disease (stenosis or regurgitation), but instead referred to the 2008 guidelines for the management of adults with congenital heart disease.

In 2008, the AHA and ACC (in collaboration with other medical societies) issued guidelines for the management of adults with congenital heart disease. For patients with isolated valvular pulmonary stenosis, the guidelines make recommendations on balloon valvulotomy or surgical intervention; however, TPVI was not addressed. In 2015, an AHA scientific statement on congenital heart disease in older adults was published and meant to complement the 2008 ACC and AHA guidelines for adults with congestive heart disease. The intent was to outline the
natural history, ramifications of childhood repair, and late initial diagnosis of congestive heart disease in older adults. The statement commented on the emerging use of the currently available transcatheter valve repair devices for both pulmonary stenosis and pulmonary valve regurgitation primarily after repair of tetralogy of Fallot. There was a specific comment that contemporary morbidity, mortality, and durability of surgical pulmonary valve regurgitation are unknown and, therefore, no contemporaneous benchmark against which to compare transcatheter valve implantation.

U.S. Preventive Services Task Force Recommendations
Not applicable.

Medicare National Coverage
State or federal mandates (e.g., Federal Employee Program) may dictate that certain U.S. Food and Drug Administration approved devices, drugs, or biologics may not be considered investigational, and thus these devices may be assessed only by their medical necessity.

Ongoing and Unpublished Clinical Trials
Some currently unpublished trials that might influence this review are listed in Table 4.

Table 4. Summary of Key Trials

<table>
<thead>
<tr>
<th>NCT No.</th>
<th>Trial Name</th>
<th>Planned Enrollment</th>
<th>Completion Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ongoing</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>NCT00740870a</td>
<td>Implantation of the Medtronic Melody Transcatheter Pulmonary Valve in Patients With Dysfunctional RVOT Conduits: A Feasibility Study</td>
<td>171</td>
<td>Jul 2020</td>
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<tr>
<td>NCT02744677a</td>
<td>Congenital Multicenter Trial of Pulmonic Valve Dysfunction Studying the SAPIEN 3 InterventIONal THV</td>
<td>108</td>
<td>Dec 2027</td>
</tr>
<tr>
<td>NCT02979587</td>
<td>The Medtronic Harmony™ Transcatheter Pulmonary Valve Clinical Study</td>
<td>184</td>
<td>Dec 2024</td>
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<tr>
<td>NCT02987387a</td>
<td>New Enrollment SAPIEN XT Post-Approval Study</td>
<td>191</td>
<td>Aug 2027</td>
</tr>
</tbody>
</table>

NCT: national clinical trial.

* Denotes industry-sponsored or cosponsored trial.

Appendix

Appendix 1. Clinical Input Responses
Appendix Table 1. Respondent Profile

<table>
<thead>
<tr>
<th>Specialty Society</th>
<th>No.</th>
<th>Name of Organization</th>
<th>Clinical Specialty</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
<td>American College of Cardiology and Society for Cardiovascular Angiography and Interventions</td>
<td>Cardiology and Cardiovascular Angiography and Interventions</td>
</tr>
</tbody>
</table>

Appendix Table 2. Respondent Conflict of Interest Disclosure

<table>
<thead>
<tr>
<th>No.</th>
<th>Conflict of Interest Policy Statement</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>The ACC and SCAI typically strive to have less than 50% of participants in clinical writing workgroups with conflicts of interest. For this project, four of the five (80%) physicians comprising the ACC-SCAI workgroup completing this document report that they have no conflicts of interest. The one member of the workgroup that does have a COI serves as a proctor/consultant for both Medtronic and Edwards Life Sciences, the two companies with TPVI devices.</td>
</tr>
</tbody>
</table>

Individual physician respondents answered at individual level. Specialty Society respondents provided aggregate information that may be relevant to the group of clinicians who provided input to the Society-level response.

ACC: American College of Cardiology; SCAI: Society for Cardiovascular Angiography and Interventions; TPVI: transcatheter pulmonary valve implantation.
Appendix 2. Clinical Input Responses

CI - Objective

Transcatheter pulmonary valve implantation (TPVI) (also referred to as transcatheter pulmonary valve replacement [TPVR]) is intended for rehabilitation of dysfunctional right ventricular outflow tract (RVOT) conduits that have been surgically placed. There is also one TPV approved for use in dysfunctional bioprosthetic pulmonary valves in the pulmonary position. TPVI is being performed off-label in native RVOTs and surgically patched RVOTs for stenosis and/or regurgitation in an attempt to provide a less invasive option than surgery and to decrease the number of sternotomy procedures needed over a lifetime. Patients with congenital heart disease involving the RVOT often require multiple open heart surgeries for RVOT reconstruction and repair and delaying the need for surgery may be of meaningful clinical benefit.

The following PICO applies to this indication.

<table>
<thead>
<tr>
<th>Populations</th>
<th>Interventions</th>
<th>Comparators</th>
<th>Outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Individuals:</td>
<td>Interventions of interest are:</td>
<td>Comparators of interest are:</td>
<td>Relevant outcomes include:</td>
</tr>
<tr>
<td>• With congenital heart disease and current right ventricular outflow tract obstruction or regurgitation</td>
<td>• Transcatheter pulmonary valve implantation</td>
<td>• Open surgical pulmonary valve replacement or reconstruction</td>
<td>• Overall survival</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Symptoms</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Functional outcomes</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Quality of life</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Hospitalizations</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Treatment-related mortality</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>• Treatment-related morbidity</td>
</tr>
</tbody>
</table>

Clinical input is sought to help determine whether the use of a particular technology for a population would provide a clinically meaningful improvement in net health outcome and whether the use is consistent with generally accepted medical practice.

Responses

- Based on the evidence and your clinical experience, describe for each clinical indication listed below the narrative rationale that includes: (1) relevant authoritative scientific evidence and/or relevant clinical scenarios (e.g., a chain of evidence) supporting that use of the technology provides clinical meaningful improvement in net health outcome; and (2) any relevant patient inclusion/exclusion criteria or clinical context important to achieve a clinically meaningful improvement in net health outcome. Please include the PMID for any relevant references. As applicable, please discuss the role of TPVI compared with open surgical valve replacement or reconstruction, particularly when patients are not considered good candidates for open valve surgery due to one or more of the following conditions: (a) high risk for surgery due to concomitant medical comorbidities; or (b) poor surgical candidate due to multiple prior sternotomy procedures for open heart surgery.

  - Right ventricle-to-pulmonary artery connection (RV-PA conduit) with or without bioprosthetic valve with at least moderate pulmonic regurgitation: This includes individuals with at least moderate pulmonary regurgitation and an RV-PA conduit that supports placement of the TPV device in the pulmonary position.

<table>
<thead>
<tr>
<th>No.</th>
<th>Rationale</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>TPVI is a well accepted and FDA-approved therapy for RV-PA conduit and bioprosthetic valve (BPV) regurgitation, regardless of the surgical risk, because of 5-7 year follow-up data from multiple prospective studies showing excellent results and durability.</td>
</tr>
<tr>
<td>No.</td>
<td>Rationale</td>
</tr>
<tr>
<td>-----</td>
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</tr>
</tbody>
</table>

Freedom from TPVI valve dysfunction is comparable to that for surgical conduits and BPVs, particularly when comparing age-matched studies, as valve dysfunction is age-related ().


Comparing valve dysfunction is more useful than comparing reintervention rates, as many surgical patients were left with dysfunction without undergoing reoperation, prior to the availability of TPVI therapy. One of the great benefits of TPVI therapy is treatment of conduit and BPV dysfunction without open heart surgery and redo sternotomy operations. STS-CHSD data show that unadjusted mortality and other complications increase with the increasing number of prior cardiopulmonary bypass (CPB) operations.


Therefore, it is beneficial to decrease the number of CPB operations over a patient's lifetime. In addition, TPVI has fewer complications of death/stroke/early reoperation than surgical valve replacement (0.8% vs. 4.3%).


Furthermore, hospital length of stay averages 1 day with TPVI compared to 4-7 days with surgical conduit or valve replacement. Cost effectiveness studies have shown TPVI cost to be comparable to or less than surgical pulmonary valve replacement.


- Native or patched RVOT with at least moderate pulmonic regurgitation: This includes individuals with at least moderate pulmonary regurgitation and native or patched RVOT that supports placement of the TPV device in the pulmonary position.
<table>
<thead>
<tr>
<th>No.</th>
<th>Rationale</th>
</tr>
</thead>
</table>

At this time, the literature is limited to case reports and case series, but larger retrospective and prospective studies are forthcoming. This practice is currently considered standard-of-care in many centers, when a landing zone is adequate, in order to decrease the number of cardiopulmonary bypass operations, hospital length of stay, and time off of work for patients.

- Right ventricle-to-pulmonary artery connection (RV-PA conduit) with or without bioprosthetic valve with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg): This includes individuals with pulmonic stenosis with a mean RVOT gradient greater or equal to 35 mm Hg and RV-PA conduit that supports placement of the TPV device.

<table>
<thead>
<tr>
<th>No.</th>
<th>Rationale</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Transcatheter pulmonary valve placement in the setting of stenosis has been reported to have excellent procedural success and has been noted to have durable improvements in patient status. Perhaps one argument against transcatheter pulmonary valve placement when compared to surgical intervention is the published rate of reintervention with TPVR. This is an effect of the pre-stent era and in this population of stenotic conduits, reinterventions were often a result of stent fracture. Additionally, previously reported risk in this population of a stenotic conduit includes conduit rupture, now able to be adequately treated with covered stents now available following the COAST and PARCS trials in nearly all cases. Eliminating at least one sternotomy, often more, in patients that will require several reentry stenotomies most arguably can be lifesaving.</td>
</tr>
</tbody>
</table>

A recent paper by Calbalka et al describes outcomes in prospective North American and European Melody valve trials. In this study, freedom from reintervention in 358 patients was 85% at 3 years. This improved rate of freedom from reintervention was attributed to the adopted practice of prestenting, thus reducing Melody frame fracture. In this study, 162 patients had stenosis or mixed disease (stenosis and regurgitation) with good results.

- Native or patched RVOT with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg): This includes individuals with pulmonic stenosis with a mean RVOT gradient greater or equal to 35 mm Hg and native or patched RVOT that supports placement of the TPV device in the pulmonary position.

<table>
<thead>
<tr>
<th>No.</th>
<th>Rationale</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>The stenotic native RVOT carries fewer risks for transcatheter valve placement compared to dilated RVOT and severely stenotic conduit cohort. This group additionally has highly variable and dynamic RVOT anatomy which can make patient selection key in success. This was outlined in a recent article by Meadows et al (2014). This study highlights the importance of selection in this native RVOT population. In this population of patients from 5 centers, 3 of the 31 patients had stenosis as the primary indication and were mixed (stenosis and regurgitation) in an additional 14 patients. All implants were successful without risk of stent migration in the stenotic group. Follow up results were good with median gradient across the RVOT of 23 mmHg and mild TPV insufficiency. Stent</td>
</tr>
</tbody>
</table>
Rationale

- Based on the evidence and your clinical experience for each of the clinical indications described in Question 1a-1d when performing TPVI using the Melody Transcatheter Pulmonary Valve (TPV):
  - Clinically Meaningful Improvement in Net Health Outcome
    - Respond YES or NO for each clinical indication whether the use of TPVI would be expected to provide a clinically meaningful improvement in net health outcome; AND
    - Rate your level of confidence in your YES or NO response using the 1 to 5 scale outlined below.

<table>
<thead>
<tr>
<th>No.</th>
<th>Indications</th>
<th>Yes/No</th>
<th>Low Confidence</th>
<th>Intermediate Confidence</th>
<th>High Confidence</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Q1a: RV-PA conduit with or without bioprosthetic valve with at least moderate pulmonic regurgitation</td>
<td>Yes</td>
<td></td>
<td></td>
<td>X</td>
</tr>
<tr>
<td></td>
<td>Q1b: Native or patched RVOT with at least moderate pulmonic regurgitation</td>
<td>Yes</td>
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<td>X</td>
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<td></td>
<td>Q1c: RV-PA conduit with or without bioprosthetic valve with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg)</td>
<td>Yes</td>
<td></td>
<td></td>
<td>X</td>
</tr>
<tr>
<td></td>
<td>Q1d: Native or patched RVOT with pulmonic stenosis (mean RVOT gradient at least 35 mm Hg)</td>
<td>Yes</td>
<td></td>
<td></td>
<td>X</td>
</tr>
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- Generally Accepted Medical Practice
  - Respond YES or NO for each clinical indication whether the use of TPVI is consistent with generally accepted medical practice; AND
  - Rate your level of confidence in your YES or NO response using the 1 to 5 scale outlined below.
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</table>

Additional narrative rationale or comments and/or any relevant scientific citations (including the PMID) supporting your clinical input on this topic.

<table>
<thead>
<tr>
<th>No.</th>
<th>Additional Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Not applicable to TPVI</td>
</tr>
</tbody>
</table>

Is there any evidence missing from the attached draft review of evidence that demonstrates clinically meaningful improvement in net health outcome?

<table>
<thead>
<tr>
<th>No.</th>
<th>Yes/No</th>
<th>Citations of Missing Evidence</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>No</td>
<td></td>
</tr>
</tbody>
</table>

[a] The choice of SAPIEN transcatheter heart valve may be based on clinical judgment such as size needed for patient anatomy.
References


Documentation for Clinical Review

Please provide the following documentation:
- History and physical and/or consultation notes including:
  - All previous surgeries, treatments, and responses pertaining to request
  - New York Heart Association Classification of symptoms
  - Pulmonary valve stenosis severity description
  - Reason for procedure
- Echocardiogram within last six months

Post Service (in addition to the above, please include the following):
- Operative report(s)

Coding

This Policy relates only to the services or supplies described herein. Benefits may vary according to product design; therefore, contract language should be reviewed before applying the terms of the Policy. Inclusion or exclusion of codes does not constitute or imply member coverage or provider reimbursement.

MN/IE
The following services may be considered medically necessary in certain instances and investigational in others. Services may be considered medically necessary when policy criteria are met. Services may be considered investigational when the policy criteria are not met or when the code describes application of a product in the position statement that is investigational.

<table>
<thead>
<tr>
<th>Type</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CPT®</td>
<td>33477</td>
<td>Transcatheter pulmonary valve implantation, percutaneous approach, including pre-stenting of the valve delivery site, when performed</td>
</tr>
<tr>
<td>HCPCS</td>
<td>None</td>
<td>None</td>
</tr>
</tbody>
</table>

Policy History

This section provides a chronological history of the activities, updates and changes that have occurred with this Medical Policy.

<table>
<thead>
<tr>
<th>Effective Date</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>03/30/2012</td>
<td>New Policy Adoption</td>
</tr>
<tr>
<td>06/30/2015</td>
<td>Coding update</td>
</tr>
<tr>
<td>02/01/2016</td>
<td>Policy revision without position change Coding update</td>
</tr>
<tr>
<td>08/01/2016</td>
<td>Policy revision without position change</td>
</tr>
<tr>
<td>09/01/2017</td>
<td>Policy revision without position change</td>
</tr>
<tr>
<td>07/01/2018</td>
<td>Policy revision without position change</td>
</tr>
<tr>
<td>09/01/2019</td>
<td>Policy revision without position change</td>
</tr>
<tr>
<td>08/01/2020</td>
<td>Annual review. No change to policy statement. Literature review updated.</td>
</tr>
</tbody>
</table>
Definitions of Decision Determinations

Medically Necessary: Services that are Medically Necessary include only those which have been established as safe and effective, are furnished under generally accepted professional standards to treat illness, injury or medical condition, and which, as determined by Blue Shield, are: (a) consistent with Blue Shield medical policy; (b) consistent with the symptoms or diagnosis; (c) not furnished primarily for the convenience of the patient, the attending Physician or other provider; (d) furnished at the most appropriate level which can be provided safely and effectively to the patient; and (e) not more costly than an alternative service or sequence of services at least as likely to produce equivalent therapeutic or diagnostic results as to the diagnosis or treatment of the Member’s illness, injury, or disease.

Investigational/Experimental: A treatment, procedure, or drug is investigational when it has not been recognized as safe and effective for use in treating the particular condition in accordance with generally accepted professional medical standards. This includes services where approval by the federal or state governmental is required prior to use, but has not yet been granted.

Split Evaluation: Blue Shield of California/Blue Shield of California Life & Health Insurance Company (Blue Shield) policy review can result in a split evaluation, where a treatment, procedure, or drug will be considered to be investigational for certain indications or conditions, but will be deemed safe and effective for other indications or conditions, and therefore potentially medically necessary in those instances.

Prior Authorization Requirements (as applicable to your plan)

Within five days before the actual date of service, the provider must confirm with Blue Shield that the member's health plan coverage is still in effect. Blue Shield reserves the right to revoke an authorization prior to services being rendered based on cancellation of the member's eligibility. Final determination of benefits will be made after review of the claim for limitations or exclusions.

Questions regarding the applicability of this policy should be directed to the Prior Authorization Department at (800) 541-6652, or the Transplant Case Management Department at (800) 637-2066 ext. 3507708 or visit the provider portal at www.blueshieldca.com/provider.

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. Blue Shield of California may consider published peer-reviewed scientific literature, national guidelines, and local standards of practice in developing its medical policy. Federal and state law, as well as contract language, including definitions and specific contract provisions/exclusions, take precedence over medical policy and must be considered first in determining covered services. Member contracts may differ in their benefits. Blue Shield reserves the right to review and update policies as appropriate.