# ConnectiCare

POLICY NUMBER	LAST REVIEW DATE	APPROVED BY
MG.MM.SU.54fC2	12/8/2023	MPC (Medical Policy Committee)

#### IMPORTANT NOTE ABOUT THIS MEDICAL POLICY:

Property of ConnectiCare, Inc. All rights reserved. The treating physician or primary care provider must submit to ConnectiCare, Inc. the clinical evidence that the patient meets the criteria for the treatment or surgical procedure. Without this documentation and information, ConnectiCare will not be able to properly review the request for prior authorization. This clinical policy is not intended to pre-empt the judgment of the reviewing medical director or dictate to health care providers how to practice medicine. Health care providers are expected to exercise their medical judgment in rendering appropriate care. The clinical review criteria expressed below reflects how ConnectiCare determines whether certain services or supplies are medically necessary. ConnectiCare established the clinical review criteria based upon a review of currently available clinical information (including clinical outcome studies in the peer-reviewed published medical literature, regulatory status of the technology, evidence-based guidelines of public health and health research agencies, evidence-based guidelines and positions of leading national health professional organizations, views of physicians practicing in relevant clinical areas, and other relevant factors). ConnectiCare, Inc. expressly reserves the right to revise these conclusions as clinical information changes, and welcomes further relevant information. Identification of selected brand names of devices, tests and procedures in a medical coverage policy is for reference only and is not an endorsement of any one device, test or procedure over another. Each benefit plan defines which services are covered. The conclusion that a particular service or supply is medically necessary does not constitute a representation or warranty that this service or supply is covered and/or paid for by ConnectiCare, as some plans exclude coverage for services or supplies that ConnectiCare considers medically necessary. If there is a discrepancy between this guideline and a member's benefits plan, the benefits plan will govern. In addition, coverage may be mandated by applicable legal requirements of the State of CT and/or the Federal Government. Coverage may also differ for our Medicare members based on any applicable Centers for Medicare & Medicaid Services (CMS) coverage statements including including National Coverage Determinations (NCD), Local Coverage Determinations (LCD) and/or Local Medical Review Policies(LMRP). All coding and web site links are accurate at time of publication.

### Definitions

Transcatheter aortic	Transcatheter aortic valve replacement (TAVR), also known as
valve replacement	transcatheter aortic valve implantation [TAVI) is a minimally invasive
(TAVR)	procedure for the treatment of aortic stenosis. A bioprosthetic valve is
	implanted percutaneously in the orifice of the native aortic valve. There
	are two access routes for TAVI— transfemoral and transapical (involving
	thoracotomy).

### Guideline

Members are eligible for TAVR coverage when the method of insertion and clinical indication are commensurate with the FDA's approval of the device and when the following criteria are met:

- 1. Severe native valve a rtic stenosis or failure defined by  $\geq 1$  of the following:
  - a. Mean aortic valve gradient  $\geq$  40mmHg
  - b. Peak jet velocity  $\geq$  4.0 m/s
  - c. Aortic valve area (AVA) < 0.8 cm2
  - d. AVA Index  $< 0.6 \text{ cm}^2/\text{m}^2$
- 2. Presence of symptomatic aortic stenosis

TAVR, as a repair to a previously implanted bio-prosthetic valve ("valve-in-valve") that has degenerated, is considered medically necessary for members at high or greater risk for open



surgery (i.e., <u>Society of Thoracic Surgeons</u> operative risk score of  $\geq 8$  % or  $\geq 15$  % risk of mortality for surgical replacement).

#### Limitations/Exclusions

TAVR is not considered medically necessary for members with existing co-morbidities that would preclude the expected benefit from correction of the aortic stenosis.

### **Applicable Procedure Codes**

33361	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; percutaneous femoral artery approach
33362	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; open femoral artery approach
33363	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; open axillary artery approach
33364	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; open iliac artery approach
33365	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; transaortic approach (eg, median sternotomy, mediastinotomy)
33366	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; transapical exposure (eg, left thoracotomy)
33367	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; cardiopulmonary bypass support with percutaneous peripheral arterial and venous cannulation (eg, femoral vessels) (List separately in addition to code for primary procedure)
33368	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; cardiopulmonary bypass support with open peripheral arterial and venous cannulation (eg, femoral, iliac, axillary vessels) (List separately in addition to code for primary procedure)
33369	Transcatheter aortic valve replacement (TAVR/TAVI) with prosthetic valve; cardiopulmonary bypass support with central arterial and venous cannulation (eg, aorta, right atrium, pulmonary artery) (List separately in addition to code for primary procedure)
33370	Transcatheter placement and subsequent removal of cerebral embolic protection device(s), including arterial access, catheterization, imaging, and radiological supervision and interpretation, percutaneous (List separately in addition to code for primary procedure)

### Applicable ICD-10 Diagnosis Codes

I06.0	Rheumatic aortic stenosis
I06.2	Rheumatic aortic stenosis with insufficiency
I06.8	Other rheumatic aortic valve diseases



I06.9	Rheumatic aortic valve disease, unspecified
I08.0	Rheumatic disorders of both mitral and aortic valves
I08.8	Other rheumatic multiple valve diseases
I08.9	Rheumatic multiple valve disease, unspecified
I35.0	Nonrheumatic aortic (valve) stenosis
I35.2	Nonrheumatic aortic (valve) stenosis with insufficiency
Q23.0	Congenital stenosis of aortic valve

### References

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Saurabh Sanon, MD et al. Transcatheter Tricuspid Valve-in-Valve and Valve-in-Ring Implantation for Degenerated Surgical Prosthesis. JACC: Cardiovascular Interventions. August 2019. Volume 12, Issue 15. DOI: 10.1016/j.jcin.2019.05.029.

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Feldman T, Saibal K, Elmariah S, et al. Randomized comparison of percutaneous repair of surgery for mitral regurgitation. 5-year results of EVERST II. JACC. 2015; 66(25):2844-285.

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Specialty matched clinical peer review.

#### **Revision history**

DATE	REVISION
10/08/2021	<ul> <li>Changed prerequisite language to "Presence of symptomatic aortic stenosis" instead of "Presence of New York Heart Association (NYHA) symptoms ≥ class II symptomatic aortic stenosis"</li> </ul>
11/11/2019	<ul> <li>Removed surgical risk prerequisite</li> <li>Reformatted and reorganized policy, transferred content to new template</li> </ul>
05/09/19	Added positive coverage for valve-in-valve repair to a previously implanted valve



• Removed provider and facility credentialing prerequisite