

*National Imaging Associates, Inc.	
Clinical guideline	Original Date: October 2009
TRANSESOPHAGEAL (TEE) ECHO	
CPT codes: 93312, 93313, 93314, 93315, 93316,	Last Revised Date: April 2023
93317, 93318, +93320, +93321, +93325	
Guideline Number: NIA_CG_066	Implementation Date: January 2024

#### **GENERAL INFORMATION**

- It is an expectation that all patients receive care/services from a licensed clinician. All appropriate supporting documentation, including recent pertinent office visit notes, laboratory data, and results of any special testing must be provided. If applicable: All prior relevant imaging results and the reason that alternative imaging cannot be performed must be included in the documentation submitted.
- Where a specific clinical indication is not directly addressed in this guideline, medical necessity
  determination will be made based on widely accepted standard of care criteria. These criteria are
  supported by evidence-based or peer-reviewed sources such as medical literature, societal
  guidelines and state/national recommendations.

#### INDICATIONS FOR TRANSESOPHAGEAL ECHOCARDIOGRAPHY (TEE)

#### General Criteria<sup>1-5</sup>

 TEE may be performed after a nondiagnostic transthoracic echocardiogram (TTE) due to inadequate visualization of relevant structures, or if there is a high likelihood of a nondiagnostic TTE

### **Aortic Pathology**

- Suspected acute aortic pathology, such as aortic dissection<sup>1,6</sup>
- Dilated aortic sinuses or ascending aorta on TTE
- Evaluation of aortic sinuses, sinotubular junction, or ascending aorta in patients with bicuspid aortic valve when morphology cannot be assessed by TTE, and other imaging including CT or MRI (Magnetic Resonance Imaging) have not been done

#### Valvular Disease<sup>1,7</sup>

 Discordance between clinical assessment and TTE assessment of the severity of mitral regurgitation (MR)

- Evaluation of mitral stenosis, when there is a discrepancy between clinical signs or symptoms, and TTE is inadequate
- Discordance between clinical assessment and TTE assessment of the severity of aortic regurgitation (AR)
- Evaluation of native or prosthetic valves with clinical signs or symptoms suggesting valve dysfunction, when TTE is inadequate
- Re-evaluation of known prosthetic valve dysfunction when it would change management or guide therapy, (and TTE is inadequate)

### Infective Endocarditis<sup>1,8,9</sup>

- Suspected infective endocarditis (IE) of native valve, prosthetic valve, or endocardial lead with positive blood culture or new murmur
- Moderate to high pretest probability of IE (i.e., staph bacteremia, fungemia, prosthetic heart valve, or intracardiac device) when TTE is negative
- Re-evaluation of IE in a patient with a change in clinical status or cardiac examination (e.g., new murmur, embolism, persistent fever, heart failure (HF), abscess, or atrioventricular block)
- Re-evaluation of IE if the patient is at elevated risk for progression/complications or when the findings alter therapy, when TTE is inadequate

#### Cardiac Mass or Source of Emboli

- Initial evaluation of patient to exclude cardiac origin of TIA or ischemic stroke<sup>1</sup>
- Evaluation of cardiac mass, suspected tumor, or thrombus<sup>1,9</sup>
- Re-evaluation of prior TEE finding for interval change (e.g., resolution of thrombus after anticoagulation), when the findings would change therapy

### Atrial Fibrillation/Flutter1

 Evaluation for clinical decision-making regarding anticoagulation, cardioversion, and/or radiofrequency ablation

## TAVR (Transcatheter Aortic Valve Replacement/Repair)<sup>1,10</sup>

- Pre-procedural assessment of annular size and shape, number of cusps, and degree of calcification, when computed tomography (CT) or CMR (Cardiovascular Magnetic Resonance) cannot be performed
- Post-procedural assessment of degree of aortic regurgitation (including valvular and paravalvular) with suspicion of valve dysfunction, if TTE is inadequate

# Patent Foramen Ovale or Atrial Septal Defect<sup>1,11</sup>



- Evaluation for anatomy, potential cardiac source of emboli, and suitability for percutaneous device closure
- Evaluation post device closure with clinical concern for infection, malposition, embolization, or persistent shunt

### Left Atrial Appendage Occlusion<sup>12</sup>

- Evaluation of anatomy, potential cardiac source of emboli, and suitability for percutaneous occlusion device placement
- Surveillance at 45 days and 1 year or FDA (U.S. Food and Drug Administration) guidance/guidelines for follow-up to assess device stability and device leak, and exclude migration, displacement, or erosion<sup>13,14</sup>
  - Reassessment at 6 months if 45-day TEE shows incomplete closure of left atrial appendage<sup>13,14</sup>

### Percutaneous Mitral Valve Repair<sup>1</sup>

- Determination of patient eligibility for percutaneous mitral valve procedures
- Pre-procedural evaluation for percutaneous mitral valve procedures may be performed in addition to CT imaging<sup>15</sup>
- To exclude the presence of intracardiac mass, thrombus, or vegetation prior to (within 3 days of) the procedure

# Hypertrophic Cardiomyopathy<sup>16</sup>

 When TTE is inconclusive in planning for myectomy,<sup>17</sup> to exclude subaortic membrane or mitral regurgitation, or to assess need for septal ablation

## Adult Congenital Heart Disease<sup>11,18</sup>

- Imaging with provocative maneuvers (Valsalva, cough) to assess the presence of rightto-left cardiac shunt
- Evaluation prior to planned repair of the following lesions when TTE, CMR, or CT are not adequate:
  - Isolated secundum atrial septal defect
  - Sinus venosus defect and/or partial anomalous pulmonary venous connection
  - Congenital mitral stenosis or mitral regurgitation
  - Subvalvular aortic stenosis
  - Transposition of the Great Arteries
- Evaluation postoperative or post catheter-based repair due to change in clinical status and/or new concerning signs or symptoms when TTE, CMR, or CT are not adequate

## Ventricular Assist Devices 1,19



- Preoperative evaluation of suitability for ventricular assist device (VAD)
- Re-evaluation of VAD-related complication or suspected infection

### **BACKGROUND**

Transesophageal echocardiography (TEE) enables cardiac ultrasound imaging from within the esophagus, which provides a window for enhanced quality images as well as additional views, beyond that acquired by standard transthoracic echocardiography (TTE).



#### **Abbreviations**

AR aortic regurgitation

CMR cardiac magnetic resonance

CT(A) computed tomography (angiography)

HF heart failure

IE infective endocarditis MR mitral regurgitation

MRI magnetic resonance imaging

TAVR transcatheter aortic valve replacement/repair

TEE transesophageal echocardiography

TIA transient ischemia attack

TTE transthoracic echocardiography

VAD ventricular assist device



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# **POLICY HISTORY**

Date	Summary	
April 2023	<ul> <li>Added statement on clinical indications not addressed in this guideline</li> </ul>	
June 2022	<ul> <li>Updated surveillance protocol of left atrial appendage occlusion device based on FDA guidance</li> </ul>	
February 2022	No significant changes	

### Reviewed / Approved by NIA Clinical Guideline Committee

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