Transvenous Implantable Cardioverter Defibrillator

MCG Health Ambulatory Care 26th Edition

AUTH: Surgical 133 (AC)

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Coverage

See the appropriate benefit document for specific coverage determination. Member specific benefits take precedence over medical policy.

Application to Products

· Policy is applicable to all products.

Authorization Requirements

Pre-certification by the Plan is required.

Description of Item or Service

Transvenous Implantable cardioverter defibrillator (ICD) is a battery-powered electronic device with a generator which is implanted under the skin with attached lead wires inserted through the vessels to the inside of the heart, which deliver an electric pulse or shock to help restore a normal heartbeat.

Exceptions and Limitations

There is insufficient scientific evidence to support the medical necessity of Transvenous Implantable Cardioverter Defibrillator for uses other than those listed in the clinical indications for procedure section.

Clinical Indications for Procedure

- Transvenous Implantable Cardioverter Defibrillator is considered medically necessary with ALL of the following
 - Individual has cardiac condition that requires Implantable Cardioverter-Defibrillator (ICD) placement as indicated by 1 or more of the following:
 - Ischemic cardiomyopathy (known coronary artery disease) and 1 or more of the following:
 - New York Heart Association class I heart failure with left ventricular ejection fraction less than or equal to 30%
 - New York Heart Association class II or III heart failure with left ventricular ejection fraction less than or equal to 35%
 - Syncope presumed to be due to ventricular arrhythmia[A] and inducible sustained (lasting more than 30 seconds) ventricular tachycardia
 - Left ventricular ejection fraction less than 40% and **ALL** of the following:

- Nonsustained ventricular tachycardia
- Inducible sustained ventricular tachycardia
- Sustained (lasting more than 30 seconds) or hemodynamically significant (eg, Hypotension)
 [C] ventricular tachycardia not believed to be due to reversible etiology (eg, myocardial ischemia, severe electrolyte abnormality)
- Nonischemic cardiomyopathy and 1 or more of the following :
 - · Syncope presumed to be due to ventricular tachycardia
 - Stable ventricular tachycardia not due to reversible causes
 - Induced or spontaneous ventricular tachycardia that is hemodynamically significant (eg, Hypotension) or sustained (lasting 30 seconds or longer)
 - New York Heart Association class I to III heart failure and ALL of the following:
 - Left ventricular ejection fraction less than or equal to 35%
 - Individual treated for at least 3 months with guideline-directed medical therapy
 - Survival is reasonably expected to be greater than 1 year
 - New York Heart Association class IV heart failure and 1 or more of the following:
 - Heart transplant candidate
 - · Left ventricular assist device candidate or implanted
 - Candidate for cardiac resynchronization therapy (ie, device will incorporate both pacing and defibrillation capabilities)
 - History of Chagas disease
 - History of myotonic dystrophy
 - · History of heart failure due to amyloidosis
 - · History of giant cell myocarditis
 - Peripartum cardiomyopathy persisting more than 3 months post partum
- Cardiac arrest due to ventricular fibrillation or ventricular tachycardia without known treatable precipitating cause (eg, myocardial ischemia, electrolyte disorder, myocarditits)
- Sustained (lasting 30 seconds or longer) ventricular tachycardia and left ventricular ejection fraction less than or equal to 35%
- Child with sustained ventricular tachycardia inadequately controlled with medication or catheter ablation, and with no evidence of incessant ventricular tachyarrhythmias
- Ventricular fibrillation or polymorphic ventricular tachycardia within 48 hours of MI and 1 or more of the following:
 - · Revascularization of infarct vessel not feasible
 - Inducible sustained ventricular tachycardia or ventricular fibrillation at electrophysiologic study performed 4 or more days after revascularization
- Individual within 40 days of MI and 1 or more of the following:
 - Syncope presumed to be due to ventricular arrhythmia[A] and inducible sustained (lasting more than 30 seconds) ventricular tachycardia
 - New York Heart Association class I heart failure with ALL of the following:
 - Left ventricular ejection fraction less than or equal to 30%
 - Recovery of left ventricular function not expected
 - Individual having permanent pacemaker placed
 - New York Heart Association class II or III heart failure with ALL of the following:
 - Left ventricular ejection fraction less than or equal to 35%
 - · Recovery of left ventricular function not expected
 - Individual having permanent pacemaker placed
 - Left ventricular ejection fraction less than 40% and ALL of the following:
 - Nonsustained ventricular tachycardia
 - Inducible sustained ventricular tachycardia
 - New York Heart Association class IV heart failure with ALL of the following:
 - Individual is ambulatory (not bed bound)
 - Individual is a candidate for cardiac transplant, left ventricular assist device, or cardiac resynchronization therapy
 - Sustained (lasting more than 30 seconds) or hemodynamically significant (eg, Hypotension) ventricular tachycardia not believed to be due to reversible etiology (eg, myocardial ischemia, severe electrolyte abnormality)
- Individual who has been revascularized (with bypass surgery or percutaneous intervention) within last 90 days and is not within 40 days of acute MI with 1 or more of the following:
 - Individual requires permanent pacemaker and ALL of the following:

- Left ventricular ejection fraction less than or equal to 35%
- Recovery of left ventricular ejection fraction not expected
- Syncope presumed to be due to ventricular arrhythmia[A] and inducible sustained (lasting more than 30 seconds) ventricular tachycardia
- Left ventricular ejection fraction less than 40% and ALL of the following:
 - Nonsustained ventricular tachycardia
 - · Inducible sustained ventricular tachycardia
- Sustained (lasting more than 30 seconds) or hemodynamically significant (eg, Hypotension) ventricular tachycardia not believed to be due to reversible etiology (eg, myocardial ischemia, severe electrolyte abnormality)
- Individual with history of MI (more than 40 days ago, not revascularized in last 90 days) 1 or more of the following:
 - Sustained (lasting more than 30 seconds) or hemodynamically significant (eg, Hypotension) ventricular tachycardia not believed to be due to reversible etiology (eg, myocardial ischemia, severe electrolyte abnormality)
 - Syncope presumed to be due to ventricular arrhythmia[A] and inducible sustained (lasting more than 30 seconds) ventricular tachycardia
 - New York Heart Association class I heart failure with left ventricular ejection fraction less than or equal to 30%
 - New York Heart Association class II or III heart failure with left ventricular ejection fraction less than or equal to 35%
 - Left ventricular ejection fraction less than 40% and ALL of the following:
 - Nonsustained ventricular tachycardia
 - Inducible sustained ventricular tachvcardia
 - Left ventricular ejection fraction less than or equal to 40%, and individual having permanent pacemaker placed
- Syncope presumed to be due to ventricular arrhythmia with 1 or more of the following:
 - · Inducible sustained ventricular tachycardia
 - Left ventricular ejection fraction less than or equal to 35%
 - Cardiac amyloidosis
 - · Arrhythmogenic right ventricular dysplasia or cardiomyopathy
 - Hypertrophic obstructive cardiomyopathy
 - Advanced structural heart disease (eq. valvular heart disease, congenital heart disease)
- Individual has genetic condition that increases risk of sudden cardiac death as indicated by 1 or more of the following:
 - Long QT syndrome and 1 or more of the following:
 - History of cardiac arrest (eg, resuscitated ventricular fibrillation)
 - Corrected QT interval greater than 500 milliseconds while receiving beta-blocker
 - Adult in whom beta-blocker therapy is ineffective (eg, persistent symptoms) or not tolerated
 - Child with symptomatic ventricular arrhythmia in whom beta-blocker therapy is ineffective or not tolerated and alternative therapies (eg, cardiac sympathetic denervation, alternative antiarrhythmic agents) are ineffective or inappropriate
 - Syncope presumed to be due to ventricular arrhythmia
 - Genotypes LQT2 or LQT3
 - Female with genotype LQT2
 - Arrhythmogenic right ventricular dysplasia or cardiomyopathy and 1 or more of the following
 - Sustained (lasting 30 seconds or longer) or hemodynamically significant (eg, Hypotension) ventricular tachycardia
 - Right or left ventricular ejection fraction less than or equal to 35%
 - Syncope presumed to be due to ventricular arrhythmia
 - Risk factors for sudden cardiac death indicated by 1 or more of the following
 - Frequent premature ventricular contractions
 - Family history of premature sudden death
 - Inducible ventricular tachycardia
 - Brugada syndrome and 1 or more of the following
 - Individual has spontaneous type 1 Brugada syndrome ECG pattern and 1 or more of the following

- Sustained (lasting 30 seconds or longer) or hemodynamically significant (eg, Hypotension) ventricular tachycardia
- Inducible sustained (lasting longer than 30 seconds) ventricular tachycardia
- History of cardiac arrest (eg, resuscitated ventricular fibrillation)
- Syncope presumed to be due to ventricular arrhythmia
- Individual with other than spontaneous type 1 Brugada syndrome ECG pattern with response to pharmacologic challenge (eg, procainamide, flecainide, ajmaline) of 1 or more of the following
 - Ventricular arrhythmia
 - Marked QRS widening
 - Type 1 Brugada syndrome ECG pattern
- Catecholaminergic polymorphic ventricular tachycardia and 1 or more of the following
 - Adult with sustained (lasting 30 seconds or more) ventricular tachycardia while receiving beta-blocker therapy
 - Adult with syncope presumed to be due to a ventricular arrhythmia[B] while receiving beta-blocker therapy
 - Child with arrhythmic syncope, or polymorphic/bidirectional ventricular tachycardia despite alternative treatment (eg, beta-blocker, flecainide, cardiac sympathetic denervation)
- Hypertrophic cardiomyopathy and 1 or more of the following
 - Previous documented cardiac arrest
 - Sustained ventricular tachycardia (lasting longer than 30 seconds)
 - History of sudden death attributed to hypertrophic cardiomyopathy in first-degree or second-degree relative 50 years of age or younger
 - Left ventricular hypertrophy 30 mm or greater in any segment
 - Nonsustained ventricular tachycardia (eg, identified on ambulatory monitoring)
 - Syncope suspected to be due to arrhythmia
 - Left ventricular ejection fraction less than 50%
 - · Late gadolinium enhancement on cardiac MRI
 - Adult individual with left ventricular apical aneurysm
- · Short QT syndrome
- · Noncompaction of left ventricle
- Lamin A/C mutation and 2 or more of the following
 - Male sex
 - Left ventricular ejection fraction less than or equal to 45%
 - Nonsustained ventricular tachycardia
- Lamin A/C mutation and indication for permanent pacemaker is present
- Phospholamban cardiomyopathy and 1 or more of the following
 - Left ventricular ejection fraction of less than 45%
 - Nonsustained ventricular tachycardia
- Filamin-C cardiomyopathy and left ventricular ejection fraction of less than 45%
- · Other familial cardiomyopathy associated with sudden death
- Cardiac sarcoidosis and 1 or more of the following
 - Ventricular tachycardia that is hemodynamically significant (eg, Hypotension) or sustained (lasting 30 seconds or longer)
 - Left ventricular ejection fraction less than or equal to 35% after treatment with optimal medical therapy for heart failure and immunosuppression (if active inflammation is present)
 - Left ventricular ejection fraction of 36% to 49% and right ventricular ejection fraction less than 40% despite treatment with optimal medical therapy for heart failure and immunosuppression (if active inflammation is present)
 - Evidence of myocardial scar by cardiac MRI or PET scan
 - · Permanent pacemaker required
 - · Syncope presumably due to ventricular arrhythmia
 - Inducible sustained ventricular arrhythmia
- Congenital heart disease and 1 or more of the following
 - Sustained (lasting 30 seconds or longer) or hemodynamically significant[C] ventricular tachycardia without identified reversible etiology (eg, myocardial ischemia, severe electrolyte abnormality)

- Inducible sustained (lasting 30 seconds or longer) ventricular tachycardia
- Single ventricle or systemic right ventricle with ejection fraction less than or equal to 35%
- Left ventricular ejection fraction 35% or less
- Awaiting heart transplant
- · Syncope presumably due to ventricular arrhythmia
- Tetralogy of Fallot and 1 or more of the following
 - Left ventricular systolic or diastolic dysfunction
 - QRS duration of 180 milliseconds or more
 - QRS fragmentation (eg, multiple notched R waves)
 - · Right ventricular scarring
 - · Inducible sustained ventricular tachycardia
 - Other risk for sudden cardiac death (eg, impaired systolic or diastolic function, nonsustained ventricular tachycardia)
- Left ventricular assist device with ventricular tachycardia that is sustained (lasting longer than 30 seconds)
- Post heart transplant with ALL of the following:
 - · Left ventricular dysfunction
 - Severe allograft vasculopathy
- Individual without contraindication to ICD placement as indicated by ALL of the following:
 - No condition limiting life expectancy to less than 1 year (eg, advanced malignancy)
 - No treatment-refractory class IV heart failure in individual who is not candidate for cardiac transplant or left ventricular assist device
 - No significant psychiatric illness that may be aggravated by device implantation or that may preclude regular follow-up
 - No ongoing IV drug abuse
 - No unresolved infection associated with risk for hematogenous seeding
 - No history of significant nonadherence with medical therapy and follow-up

Document History

- · Revised Dates:
- · Reviewed Dates:
- Effective Date: July 2023

Coding Information

- CPT/HCPCS codes covered if policy criteria is met:
 - CPT 33216 Insertion of a single transvenous electrode, permanent pacemaker or implantable defibrillator
 - CPT 33217 Insertion of 2 transvenous electrodes, permanent pacemaker or implantable defibrillator
 - CPT 33249 Insertion or replacement of permanent implantable defibrillator system, with transvenous lead(s), single or dual chamber
- CPT/HCPCS codes considered not medically necessary per this Policy:
 - None

Codes

CPT®: 33216, 33217, 33249

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