2014 Billing and Coding Guide

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Dear GuidePoint User,

The Boston Scientific Health Economics and Reimbursement team is pleased to bring you the 2014 GuidePoint materials. GuidePoint is our suite of health economics and reimbursement resources for hospitals, physicians, clinicians, and reimbursement professionals. GuidePoint members have access to the following resources:

- **Billing and Coding Guide** — Quickly find coding and billing information, including common scenarios relevant to your medical practice.
- **Procedural Payment Guide**—Locate facility and physician payment information for cardiology, rhythm, and intervention procedures in conveniently organized summaries.
- **Webcasts** — Hear from nationally acclaimed experts addressing basic and advanced CRM and EP reimbursement topics.
- **Physician Website** — Keep current with the latest reimbursement news and find other reimbursement education resources.

For over 35 years, Boston Scientific CRM and EP have been committed to making more possible through innovation, clinical science, and collaboration. We’re dedicated to providing physicians and allied health professionals with world class programs and services to help advance the standard of patient care. We are proud to continue this spirit of partnership with GuidePoint.

We welcome your feedback. Please send comments to crm.reimbursement@bsci.com. If you have questions about GuidePoint resources or would like additional guides, contact Boston Scientific at 1.800.CARDIAC (227.3422). To access additional reimbursement resources, visit our website at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement).

Boston Scientific
Health Economics & Reimbursement
The information in this guide is current as of January 1, 2014. The Centers for Medicare and Medicaid Services (CMS) may initiate changes to coverage, coding, or payment guidelines at any time. Check the CMS website (http://www.cms.gov) for current information.

A Word to Our Customers

Boston Scientific is pleased you have chosen to partner with us to help you save and improve patients’ lives. We are committed to working directly with you to ensure timely patient access to innovative medical solutions. As part of this commitment, we also work with the Centers for Medicare and Medicaid Services (CMS), private insurers, and other industry stakeholders to ensure appropriate reimbursement for physicians and hospitals.

Explanation of Contents

This document contains commonly used billing codes for physicians and hospitals related to Boston Scientific devices and procedures.

Disclaimer

Please note: this coding information may include codes for procedures for which Boston Scientific currently offers no cleared or approved products. In those instances, such codes have been included solely in the interest of providing users with comprehensive coding information and are not intended to promote the use of any Boston Scientific products for which they are not cleared or approved.

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Payer policies will vary and should be verified prior to treatment for limitations on diagnosis, coding or site of service requirements. The coding options listed within this guide are commonly used codes and are not intended to be an all-inclusive list. We recommend consulting your relevant manuals for appropriate coding options.

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Introduction

GuidePoint Reimbursement Resources at a Glance

REIMBURSEMENT CUSTOMER SUPPORT LINE
Certified reimbursement professionals answer reimbursement questions related to Boston Scientific products and procedures.

- Call 1.800.CARDIAC (227.3422). Ask to be connected with the Reimbursement Customer Support Line available Monday through Friday, 9 am to 4 pm Central.

BILLING AND CODING GUIDE
The 2014 Billing and Coding Guide is a useful tool for hospital and physician billers and coders. The guide includes practical coverage and coding reference materials for Boston Scientific products and procedures.

PROCEDURAL PAYMENT GUIDE
The 2014 Procedural Payment Guide provides facility and physician payment information for cardiology, rhythm, and intervention procedures in convenient summaries.

WEBCAST PROGRAMS
Attend a live webcast or view on-demand topics related to coverage, coding, and payment. Webcast registration will open approximately three weeks before the live event. The webcasts are approximately one hour in length and will be available on the website for future viewing. On-demand courses are made available for you to access at your viewing convenience.

Our webcast programs are intended for hospitals, physicians, clinicians, and reimbursement professionals seeking a better understanding of reimbursement for Boston Scientific products and procedures.

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PHYSICIAN WEBSITE
Dedicated to topics associated with reimbursement, the website provides resources for those seeking a better understanding of reimbursement for Boston Scientific products and procedures.

- Make the website your first stop for all your Boston Scientific reimbursement needs; access http://www.bostonscientific.com/reimbursement.
**Medicare Payment Overview**

**OVERVIEW OF MEDICARE PAYMENT SYSTEMS**

Medicare is a federally-funded, national health insurance program providing coverage to Americans who are 65 years of age or older, certain younger people with disabilities, and individuals with end-stage renal disease (ESRD). Payment by Medicare is predicated on Medical Necessity.

**Note:** Medical Necessity is defined by CMS as services or supplies that are: proper and needed for the diagnosis or treatment of the patient’s medical condition; are provided for the diagnosis, direct care, and treatment of the patient’s medical condition; meet the standards of good medical practice in the local area; and are not mainly for the convenience of the patient’s doctor. CMS’s definition of Medical Necessity can be found at: [https://www.cms.gov/apps/glossary/default.asp?Letter=M&Language=English](https://www.cms.gov/apps/glossary/default.asp?Letter=M&Language=English)

There are several payment systems within the Medicare program, including payment for inpatient hospital services, outpatient hospital services, ambulatory surgery centers, home health, physicians, and skilled nursing. In this guide, you will find information specific to facility and physician payment systems.

**Hospital Inpatient Payment**

The hospital inpatient payment system is a prospective payment system (PPS) that classifies patients according to diagnosis, type of treatment, age, and other relevant criteria using the ICD-9-CM coding system. Under this system, hospitals typically receive a predefined payment for treating patients within a particular category or Medicare Severity Diagnosis Related Group (MS-DRG).

**Note:** Medicare’s hospital inpatient payment information in this document is effective for Fiscal Year (FY) 2014 (October 1, 2013 – September 30, 2014).

**Note:** Maryland hospitals are paid under a program waiver (section 1814(b)(3) of the Social Security Act) in which the state establishes hospital inpatient and outpatient payment rates for Medicare, Medicaid, and private payers.¹²

**Hospital Outpatient Payment**

The hospital outpatient payment system, OPPS, is also a prospective payment system. In this system, hospitals receive a fixed payment, called an Ambulatory Payment Classification (APC), for a specific procedure. Each procedure described by a CPT® (Current Procedural Terminology) code is assigned directly to an APC. Unlike the inpatient (MS-DRG) payment system, if multiple procedures are performed, the hospital may be eligible to receive more than one APC payment per outpatient admission.

**Note:** Medicare’s hospital outpatient payment information in this document is effective for Calendar Year (CY) 2014 (January 1, 2014 – December 31, 2014).

**Ambulatory Surgery Center (ASC) Payment**

The Medicare ASC payment system, effective January 1, 2014, is a prospective payment system based on Medicare’s OPPS using APCs. The new ASC payment rates for most surgical procedures are set at ~ 65% of the APC payment rate for OPPS. Device intensive procedures (such as pacemakers and defibrillators) will be paid at a higher rate (~86–96%) of the OPPS rate. ASCs should bill Medicare using a CMS-1500 claim form and use CPT® codes to describe procedures performed.

**Note:** Medicare’s ASC payment information in this document is effective for Calendar Year (CY) 2014 (January 1, 2014 – December 31, 2014).
**Physician Payment**

Physicians receive payment for each CPT® procedure code based on a fee schedule called the Physician Fee Schedule. The Physician Fee Schedule is based on a scale of national uniform values for all physician services, commonly referred to as the Resource-Based Relative Value Scale (RBRVS).

*Note*: Medicare’s physician payment information in this document is effective for Calendar Year (CY) 2014 (January 1, 2014 – December 31, 2014).

**OVERVIEW OF MEDICARE PAYMENT PROCESS**

*All Medicare payment processes include these common steps:*

1. **Step 1**: Physician documentation in patient medical record
2. **Step 2**: Transfer of information to billing/coding department
3. **Step 3**: Selection of appropriate diagnosis and procedure codes
4. **Step 4**: Submission of billing form to Medicare Administrative Contractor (MAC)
5. **Step 5**: Review of coding and physician documentation for medical necessity
6. **Step 6**: Payment from MAC to hospital or physician (if deemed medically necessary)

**Payer Coverage + Correct Coding + Compliance = Payment**

![Diagram of Medicare payment process]

**Coverage**
Reasonable and necessary
National coverage determinations (NCDs)
Local coverage determinations (LCDs)

**ICD-9-CM diagnosis codes**
Why patient received treatment

**CPT® HCPCS procedure codes**
Treatment patient received

**ICD-9-CM procedure codes**
Treatment patient received

**Fee schedule payment**
(physician)

**APC payment**
(hospital outpatient)

**MS-DRG payment**
(hospital inpatient)

*Note*: ICD-9-CM codes and HCPCS/CPT® codes are also recognized by non-Medicare payers.
**Medicare National Coverage Determination (NCD) Policies**

**MEDICARE NCD FOR CARDIAC PACEMAKERS**

**Pacemaker – Implantable**
- **Effective date of this version:** April 30, 2004
- **Implementation date:** April 30, 2004

**Benefit Category**
- Prosthetic Devices

**Note:** This may not be an exhaustive list of all applicable Medicare benefit categories for this item or service.

**Coverage Topic**
- Prosthetic Devices

**Item/Service Description**
Cardiac pacemakers are self-contained, battery-operated units that send electrical stimulation to the heart. They are generally implanted to alleviate symptoms of decreased cardiac output related to abnormal heart rate and/or rhythm. Pacemakers are generally used for persistent, symptomatic second- or third-degree atrioventricular (AV) block, and symptomatic sinus bradycardia.

**Indications and Limitations of Coverage**
Cardiac pacemakers are covered as prosthetic devices under the Medicare program, subject to the following conditions and limitations. While cardiac pacemakers have been covered under Medicare for many years, there were no specific guidelines for their use other than the general Medicare requirement that covered services be reasonable and necessary for the treatment of the condition. Services rendered for cardiac pacing on or after the effective dates of this instruction are subject to these guidelines, which are based on certain assumptions regarding the clinical goals of cardiac pacing. While some uses of pacemakers are relatively certain or unambiguous, many other uses require considerable expertise and judgment.

Consequently, the medical necessity for permanent cardiac pacing must be viewed in the context of overall patient management. The appropriateness of such pacing may be conditional on other diagnostic or therapeutic modalities having been undertaken. Although significant complications and adverse side effects of pacemaker use are relatively rare, they cannot be ignored when considering the use of pacemakers for dubious medical conditions, or marginal clinical benefit.

These guidelines represent current concepts regarding medical circumstances in which permanent cardiac pacing may be appropriate or necessary. As with other areas of medicine, advances in knowledge and techniques in cardiology are expected. Consequently, judgments about the medical necessity and acceptability of new uses for cardiac pacing in new classes of patients may change as more conclusive evidence becomes available. This instruction applies only to permanent cardiac pacemakers, and does not address the use of temporary, non-implanted pacemakers.

The two groups of conditions outlined below deal with the necessity for cardiac pacing for patients in general. These are intended as guidelines in assessing the medical necessity for pacing therapies, taking into account the particular circumstances in each case. However, as a general rule, the two groups of current medical concepts may be viewed as representing:

- **Group I: Single-chamber Cardiac Pacemakers** – a) conditions under which single-chamber pacemaker claims may be considered covered without further claims development; and b) conditions under which single-chamber pacemaker claims would be denied unless further claims development shows that they fall into the covered category, or special medical circumstances sufficient to convince the contractor that the claim should be paid.
• **Group II: Dual-chamber Cardiac Pacemakers** – a) conditions under which dual-chamber pacemaker claims may be considered covered without further claims development, and b) conditions under which dual-chamber pacemaker claims would be denied unless further claims development shows that they fall into the covered categories for single- and dual-chamber pacemakers, or special medical circumstances exist sufficient to convince the contractor that the claim should be paid.

CMS opened the NCD on Cardiac Pacemakers to afford the public an opportunity to comment on the proposal to revise the language contained in the instruction. The revisions transfer the focus of the NCD from the actual pacemaker implantation procedure itself to the reasonable and necessary medical indications that justify cardiac pacing. This is consistent with our findings that pacemaker implantation is no longer considered routinely harmful or an experimental procedure.

**GROUP I SINGLE-CHAMBER CARDIAC PACEMAKERS**

- **Effective: March 16, 1983**

A. **Nationally Covered Indications**

Conditions under which cardiac pacing is generally considered acceptable or necessary, provided that the conditions are chronic or recurrent and not due to transient causes such as acute myocardial infarction, drug toxicity, or electrolyte imbalance. (In cases where there is a rhythm disturbance, if the rhythm disturbance is chronic or recurrent, a single episode of a symptom such as syncope or seizure is adequate to establish medical necessity).

1. Acquired complete (also referred to as third-degree) AV heart block.
2. Congenital complete heart block with severe bradycardia (in relation to age), or significant physiological deficits or significant symptoms due to the bradycardia.
3. Second-degree AV heart block of Type II (i.e., no progressive prolongation of P-R interval prior to each blocked beat. P-R interval indicates the time taken for an impulse to travel from the atria to the ventricles on an electrocardiogram).
4. Second-degree AV heart block of Type I (i.e., progressive prolongation of P-R interval prior to each blocked beat) with significant symptoms due to hemodynamic instability associated with the heart block.
5. Sinus bradycardia associated with major symptoms (e.g., syncope, seizures, congestive heart failure); or substantial sinus bradycardia (heart rate less than 50) associated with dizziness or confusion. The correlation between symptoms and bradycardia must be documented, or the symptoms must be clearly attributable to the bradycardia rather than to some other cause.
6. In selected and few patients, sinus bradycardia of lesser severity (heart rate 50-59) with dizziness or confusion. The correlation between symptoms and bradycardia must be documented, or the symptoms must be clearly attributable to the bradycardia rather than to some other cause.
7. Sinus bradycardia is the consequence of long-term necessary drug treatment for which there is no acceptable alternative when accompanied by significant symptoms (e.g., syncope, seizures, congestive heart failure, dizziness, or confusion). The correlation between symptoms and bradycardia must be documented, or the symptoms must be clearly attributable to the bradycardia rather than to some other cause.
8. Sinus node dysfunction with or without tachyarrhythmias or AV conduction block (i.e., the bradycardia-tachycardia syndrome, sino-atrial block, sinus arrest) when accompanied by significant symptoms (e.g., syncope, seizures, congestive heart failure, dizziness, or confusion).
9. Sinus node dysfunction with or without symptoms when there are potentially life-threatening ventricular arrhythmias or tachycardia secondary to the bradycardia (e.g., numerous premature ventricular contractions, couplets, runs of premature ventricular contractions, or ventricular tachycardia).
10. Bradycardia associated with supraventricular tachycardia (e.g., atrial fibrillation, atrial flutter, or paroxysmal atrial tachycardia) with high-degree AV block which is unresponsive to appropriate
pharmacological management and when the bradycardia is associated with significant symptoms (e.g., syncope, seizures, congestive heart failure, dizziness, or confusion).

11. The occasional patient with hypersensitive carotid sinus syndrome with syncope due to bradycardia and unresponsive to prophylactic medical measures.

12. Bifascicular or trifascicular block accompanied by syncope which is attributed to transient complete heart block after other plausible causes of syncope have been reasonably excluded.

13. Prophylactic pacemaker use following recovery from acute myocardial infarction during which there was temporary complete (third-degree) and/or Mobitz Type II second-degree AV block in association with bundle branch block.

14. In patients with recurrent and refractory ventricular tachycardia, “overdrive pacing” (pacing above the basal rate) to prevent ventricular tachycardia.

- Effective: May 9, 1985

15. Second-degree AV heart block of Type I with the QRS complexes prolonged.

B. Nationally Non-covered Indications

Conditions which, although used by some physicians as a basis for permanent cardiac pacing, are considered unsupported by adequate evidence of benefit and therefore should not generally be considered appropriate uses for single-chamber pacemakers in the absence of the above indications. Contractors should review claims for pacemakers with these indications to determine the need for further claims development prior to denying the claim, since additional claims development may be required. The object of such further development is to establish whether the particular claim actually meets the conditions in A above. In claims where this is not the case or where such an event appears unlikely, the contractor may deny the claim.

1. Syncope of undetermined cause.
2. Sinus bradycardia without significant symptoms.
3. Sino-atrial block or sinus arrest without significant symptoms.
4. Prolonged P-R intervals with atrial fibrillation (without third-degree AV block) or with other causes of transient ventricular pause.
5. Bradycardia during sleep.
6. Right bundle branch block with left axis deviation (and other forms of fascicular or bundle branch block) without syncope or other symptoms of intermittent AV block.
7. Asymptomatic second-degree AV block of Type I unless the QRS complexes are prolonged or electrophysiological studies have demonstrated that the block is at or beyond the level of the His bundle (a component of the electrical conduction system of the heart).

- Effective: October 1, 2001
8. Asymptomatic bradycardia in post-myocardial infarction patients about to initiate long-term beta-blocker drug therapy.

GROUP II DUAL-CHAMBER CARDIAC PACEMAKERS

- Effective: May 9, 1985

A. Nationally Covered Indications

Conditions under dual-chamber cardiac pacing are considered acceptable or necessary in the general medical community unless conditions 1 and 2 under Group II B are present:
1. Patients in whom single-chamber pacing (ventricular pacing) at the time of pacemaker insertion elicits a definite drop in blood pressure, retrograde conduction, or discomfort.

2. Patients in whom the pacemaker syndrome (atrial ventricular asynchrony), with significant symptoms, has already been experienced with a pacemaker that is being replaced.

3. Patients in whom even a relatively small increase in cardiac efficiency will importantly improve the quality of life, e.g., patients with congestive heart failure despite adequate other medical measures.

4. Patients in whom the pacemaker syndrome can be anticipated, e.g., in young and active people.

Dual-chamber pacemakers also may be covered for the conditions, as listed in Group IA if the medical necessity is sufficiently justified through adequate claims development. Expert physicians differ in their judgments about what constitutes appropriate criteria for dual-chamber pacemaker use. The judgment that such a pacemaker is warranted in the patient meeting accepted criteria must be based upon the individual needs and characteristics of that patient, weighing the magnitude and likelihood of anticipated benefits against the magnitude and likelihood of disadvantages to the patient.

B. Nationally Non-covered Indications
Whenever the following conditions (which represent overriding contraindications) are present, dual-chamber pacemakers are not covered:

1. Ineffective atrial contractions (e.g., chronic atrial fibrillation or flutter, or giant left atrium).

2. Frequent or persistent supraventricular tachycardias, except where the pacemaker is specifically for the control of the tachycardia.

3. A clinical condition in which pacing takes place only intermittently and briefly, and which is not associated with a reasonable likelihood that pacing needs will become prolonged, (e.g., the occasional patient with hypersensitive carotid sinus syndrome with syncope due to bradycardia and unresponsive to prophylactic medical measures).

4. Prophylactic pacemaker use following recovery from acute myocardial infarction during which there was temporary complete (third-degree) and/or Type II second-degree AV block in association with bundle branch block.

C. Other
All other indications for dual-chamber cardiac pacing for which CMS has not specifically indicated coverage remain nationally non-covered, except for Category B IDE clinical trials, or as routine costs of dual-chamber cardiac pacing associated with clinical trials, in accordance with section 310.1 of the NCD Manual. (This NCD last reviewed June 2004).

MEDICARE NCD FOR CARDIAC PACEMAKER – EVALUATION SERVICES

- **Effective date of this version:** October 1, 1984

**Benefit Category**
- Diagnostic Services in Outpatient Hospital
- Diagnostic Tests (other)

**Note:** This may not be an exhaustive list of all applicable Medicare benefit categories for this item or service.

**Coverage Topic**
- Diagnostic Tests, X-rays, and Lab Services
Item/Service Description

There are two general types of pacemakers in current use—single-chamber pacemakers which sense and pace the ventricles of the heart, and dual-chamber pacemakers which sense and pace both the atria and the ventricles. These differences require different monitoring patterns over the expected life of the units involved.

Indications and Limitations of Coverage

Medicare covers a variety of services for the post-implant follow-up and evaluation of implanted cardiac pacemakers. The following guidelines are designed to assist contractors in identifying and processing claims for such services.

Note: These new guidelines are limited to lithium battery-powered pacemakers, because mercury-zinc battery-powered pacemakers are no longer being manufactured and virtually all have been replaced by lithium units. Contractors still receiving claims for monitoring such units should continue to apply the guidelines published in 1980 to those units until they are replaced.

One fact of which contractors should be aware is that many dual-chamber units may be programmed to pace only the ventricles; this may be done either at the time the pacemaker is implanted or at some time afterward. In such cases, a dual-chamber unit, when programmed or reprogrammed for ventricular pacing, should be treated as a single-chamber pacemaker in applying screening guidelines.

The decision as to how often any patient’s pacemaker should be monitored is the responsibility of the patient’s physician, who is best able to take into account the condition and circumstances of the individual patient. These may vary over time, requiring modifications of the frequency with which the patient should be monitored. In cases where monitoring is done by some entity other than the patient’s physician, such as a commercial monitoring service or hospital outpatient department, the physician’s prescription for monitoring is required and should be periodically renewed (at least annually) to assure that the frequency of monitoring is proper for the patient.

Where a patient is monitored both during clinic visits and transtelephonically, the contractor should be sure to include frequency data on both types of monitoring in evaluating the reasonableness of the frequency of monitoring services received by the patient.

Since there are more than 200 pacemaker models in service at any given point, and a variety of patient conditions that give rise to the need for pacemakers, the question of the appropriate frequency of monitorings is a complex one. Nevertheless, it is possible to develop guidelines within which the vast majority of pacemaker monitorings will fall, and contractors should do this, using their own data and experience, as well as the frequency guidelines that follow, in order to limit extensive claims development to those cases requiring special attention.

PACEMAKER – TRANSTELEPHONIC MONITORING5

Benefit Category

• Outpatient Hospital Services Incident to a Physician’s Service

Note: This may not be an exhaustive list of all applicable Medicare benefit categories for this item or service.

A. General

Transtelephonic monitoring of pacemakers is furnished by commercial suppliers, hospital outpatient departments, and physicians’ offices.

Telephone monitoring of cardiac pacemakers as described below is medically efficacious in identifying early signs of possible pacemaker failure, thus reducing the number of sudden pacemaker failures requiring emergency replacement. All systems that monitor the pacemaker rate (bpm) in both the free-running and/or magnetic mode are effective in detecting subclinical pacemaker failure due to battery depletion. More sophisticated systems are also capable of detecting internal electronic problems within the pulse generator itself and other potential problems. In the case of dual-chamber pacemakers in particular, such monitoring may detect failure of synchronization of the atria and ventricles, and the need for adjustment and reprogramming of the device.
Note: The transmitting device furnished to the patient is simply one component of the diagnostic system, and is not covered as durable medical equipment. Those engaged in transtelephonic pacemaker monitoring should reflect the costs of the transmitters in setting their charges for monitoring.

B. Definition of Transtelephonic Monitoring

In order for transtelephonic monitoring services to be covered, the services must consist of the following elements:

- A minimum 30-second readable strip of the pacemaker in the free-running mode
- Unless contraindicated, a minimum 30-second readable strip of the pacemaker in the magnetic mode
- A minimum 30 seconds of readable ECG strip

Indications and Limitations of Coverage

C. Frequency Guidelines for Transtelephonic Monitoring

The guidelines below constitute a system that contractors should use, in conjunction with their knowledge of local medical practices, to screen claims for transtelephonic monitoring prior to payment. It is important to note that they are not recommendations with respect to a minimum frequency for such monitorings, but rather a maximum frequency (within which payment may be made without further claims development). As with previous guidelines, more frequent monitorings may be covered in cases where contractors are satisfied that such monitorings are medically necessary; e.g., based on the condition of the patient, or with respect to pacemakers exhibiting unexpected defects or premature failure. Contractors should seek written justification for more frequent monitorings from the patient’s physician and/or any monitoring service involved.

These guidelines are divided into two broad categories—Guideline I, which will apply to the majority of pacemakers now in use, and Guideline II, which will apply only to pacemaker systems (pacemaker and leads) for which sufficient long-term clinical information exists to assure that they meet the standards of the Inter-Society Commission for Heart Disease Resources (ICHD) for longevity and end-of-life decay. (The ICHD standards are: (1) 90% cumulative survival at five years following implant; and (2) an end-of-life decay of less than a 50% drop of output voltage and less than 20% deviation of magnet rate, or a drop of five beats per minute or less, over a period of three months or more). Contractors should consult with their medical advisers and other appropriate individuals and organizations (such as the North American Society of Pacing and Electrophysiology, which publishes product reliability information) should questions arise over whether a pacemaker system meets the ICHD standards.

The two groups of guidelines are then further broken down into two general categories – single-chamber and dual-chamber pacemakers. Contractors should be aware that the frequency with which a patient is monitored may be changed from time to time for a number of reasons, such as a change in the patient’s overall condition, a reprogramming of the patient’s pacemaker, the development of better information on the pacemaker’s longevity or failure mode, etc. Consequently, changes in the proper set of guidelines may be required. Contractors should inform physicians and monitoring services to alert contractors to any changes in the patient’s monitoring prescription that might necessitate changes in the screening guidelines applied to that patient. (Of particular importance is the reprogramming of a dual-chamber pacemaker to a single-chamber mode of operation. Such reprogramming would shift the patient from the appropriate dual-chamber guideline to the appropriate single chamber guideline).

MEDICARE’S FREQUENCY GUIDELINES FOR TRANSTELEPHONIC MONITORING OF CARDIAC PACEMAKERS

Guideline I

1. Single-chamber pacemakers:
   1st month: every 2 weeks
   2nd through 36th month: every 8 weeks
   37th month to failure: every 4 weeks
2. Dual-chamber pacemakers:
   1st month: every 2 weeks
   2nd through 6th month: every 4 weeks
   7th through 36th month: every 8 weeks
   37th month to failure: every 4 weeks

Guideline II

1. Single-chamber pacemakers:
   1st month: every 2 weeks
   2nd through 48th month: every 12 weeks
   49th through 72nd month: every 8 weeks
   Thereafter: every 4 weeks

2. Dual-chamber pacemakers:
   1st month: every 2 weeks
   2nd through 30th month: every 12 weeks
   31st through 48th month: every 8 weeks
   Thereafter: every 4 weeks

D. Pacemaker Clinic Services

1. General
   Pacemaker monitoring is also covered when done by pacemaker clinics. Clinic visits may be done in
   conjunction with transtelephonic monitoring or as a separate service; however, the services rendered by a
   pacemaker clinic are more extensive than those currently possible by telephone. They include, for
   example, physical examination of patients and reprogramming of pacemakers. Thus, the use of one of
   these types of monitoring does not preclude concurrent use of the other.

2. Frequency Guidelines
   As with transtelephonic pacemaker monitoring, the frequency of clinic visits is the decision of the patient's
   physician taking into account, among other things, the medical condition of the patient. However,
   contractors can develop monitoring guidelines that will prove useful in screening claims. The following are
   recommendations for monitoring guidelines on lithium-battery pacemakers:

MEDICARE'S FREQUENCY GUIDELINES FOR PACEMAKER CLINIC SERVICES

- For single-chamber pacemakers: twice in the first 6 months following implant, then once every 12 months
- For dual-chamber pacemakers: twice in the first 6 months, then once every 6 months

Pacemaker – Temporary

At this time there is no specific Medicare National Coverage Determination (NCD) for temporary pacemakers. It is
important for medical providers to check with their local MAC or non-Medicare payer(s) to determine patient
coverage and coding/billing guidelines.

Note: Search the Medicare Coverage Database on the CMS website
(http://www.cms.hhs.gov/mcd/search.asp) for coverage descriptions and updates.

MEDICARE NCD FOR IMPLANTABLE CARDIOVERTER-DEFIBRILLATORS® (ICDS)

- Effective date of this version: January 27, 2005
- Implementation date: January 27, 2005

Benefit Category

- Prosthetic Devices

Note: This may not be an exhaustive list of all applicable Medicare benefit categories for this item or service.
**Item/Service Description**

**A. General**

The implantable automatic defibrillator is an electronic device designed to detect and treat life-threatening tachyarrhythmias. The device consists of a pulse generator and electrodes for sensing and defibrillating.

**Indications and Limitations of Coverage**

**B. Covered Indications**

1. Documented episode of cardiac arrest due to ventricular fibrillation (VF), not due to a transient or reversible cause (effective July 1, 1991).

2. Documented sustained ventricular tachyarrhythmia (VT), either spontaneous or induced by an electrophysiology (EP) study, not associated with an acute myocardial infarction (MI) and not due to a transient or reversible cause (effective July 1, 1999).

3. Documented familial or inherited conditions with a high risk of life-threatening VT, such as long QT syndrome or hypertrophic cardiomyopathy (effective July 1, 1999).

**Additional indications effective for services performed on or after October 1, 2003:**

4. Coronary artery disease with a documented prior MI, a measured left ventricular ejection fraction (LVEF) < 0.35, and inducible, sustained VT or VF at EP study. (The MI must have occurred more than 40 days prior to defibrillator insertion. The EP test must be performed more than four weeks after the qualifying MI.)

5. Documented prior MI and a measured LVEF / < 0.30 and a QRS duration of > 120 milliseconds (the QRS restriction does not apply to services performed on or after January 27, 2005). Patients must not have:
   a) New York Heart Association (NYHC) classification IV;
   b) Cardiogenic shock or symptomatic hypotension while in a stable baseline rhythm;
   c) Had a coronary artery bypass graft (CABG) or percutaneous transluminal coronary angioplasty (PTCA) within past three months;
   d) Had an enzyme positive MI within the past month (Effective for services on or after January 27, 2005, patients must not have had an acute MI in the past 40 days);
   e) Clinical symptoms or findings that would make them a candidate for coronary revascularization; or
   f) Any disease, other than cardiac disease (e.g., cancer, uremia, liver failure), associated with a likelihood of survival less than 1 year.

**Additional indications effective for services performed on or after January 27, 2005:**

6. Patients with ischemic dilated cardiomyopathy (IDCM), documented prior MI, NYHA Class II and III heart failure, and measured LVEF < 35%;

7. Patients with non-ischemic dilated cardiomyopathy (NIDCM) > 9 months, NYHA Class II and III heart failure, and measured LVEF < 35%;

8. Patients who meet all current Centers for Medicare and Medicaid Services (CMS) coverage requirements for a cardiac resynchronization therapy (CRT) device and have NYHA Class IV heart failure.

**All indications must meet the following criteria:**

a) Patients must not have irreversible brain damage from preexisting cerebral disease;

b) MIs must be documented and defined according to the consensus document of the Joint European Society of Cardiology/American College of Cardiology Committee for the Redefinition of Myocardial Infarction.
Indications 3–8 (primary prevention of sudden cardiac death) must also meet the following criteria:

a) Patients must be able to give informed consent;

b) Patients must not have:
   - Cardiogenic shock or symptomatic hypotension while in a stable baseline rhythm;
   - Had a CABG or PTCA within the past three months;
   - Had an acute MI within the past 40 days;
   - Clinical symptoms or findings that would make them a candidate for coronary revascularization;
   - Any disease, other than cardiac disease (e.g., cancer, uremia, liver failure), associated with a likelihood of survival less than one year.

c) Ejection fractions must be measured by angiography, radionuclide scanning, or echocardiography;

d) The beneficiary receiving the defibrillator implantation for primary prevention is enrolled in either a Food and Drug Administration (FDA)-approved category B investigational device exemption (IDE) clinical trial (42 CFR §405.201), a trial under the CMS Clinical Trial Policy (National Coverage Determination (NCD) Manual §310.1), or a qualifying data collection system including approved clinical trials and registries. Initially, an implantable cardiac defibrillator (ICD) database will be maintained using a data submission mechanism that is already in use by Medicare participating hospitals to submit data to the Iowa Foundation for Medical Care (IFMC), a Quality Improvement Organization (QIO) contractor, for determination of reasonable and necessary quality improvement. Initial hypothesis and data elements are specified in this decision (Appendix VI) and are the minimum necessary to ensure that the device is reasonable and necessary. Data collection will be completed using the ICDA (ICD Abstraction Tool) and transmitted via QNet (Quality Network Exchange) to the IFMC, who will collect and maintain the database. Additional stakeholder-developed data collection systems to augment or replace the initial QNet system, addressing at a minimum the hypotheses specified in this decision, must meet the following basic criteria:
   - Written protocol on file;
   - Institutional review board review and approval;
   - Scientific review and approval by two or more qualified individuals who are not part of the research team;
   - Certification that investigators have not been disqualified.

For purposes of this coverage decision, CMS will determine whether specific registries or clinical trials meet these criteria.

e) Providers must be able to justify the medical necessity of devices other than single lead devices. This justification should be available in the patient’s medical record.

9. Patients with NIDCM > 3 months, NYHA Class II or III heart failure, and measured LVEF < 35%, only if the following additional criteria are also met:

a) Patients must be able to give informed consent;

b) Patients must not have:
   - Cardiogenic shock or symptomatic hypotension while in a stable baseline rhythm;
   - Had a CABG or PTCA within the past three months;
   - Had an acute MI within the past 40 days;
   - Clinical symptoms or findings that would make them a candidate for coronary revascularization;
   - Irreversible brain damage from preexisting cerebral disease;
• Any disease, other than cardiac disease (e.g., cancer, uremia, liver failure), associated with a likelihood of survival less than one year;

  c) Ejection fractions must be measured by angiography, radionuclide scanning, or echocardiography;

  d) MIs must be documented and defined according to the consensus document of the Joint European Society of Cardiology/American College of Cardiology Committee for the Redefinition of Myocardial Infarction;

  e) The beneficiary receiving the defibrillator implantation for this indication is enrolled in either an FDA-approved category B IDE clinical trial (42 CFR §405.201), a trial under the CMS Clinical Trial Policy (NCD Manual §310.1), or a prospective data collection system meeting the following basic criteria:

    • Written protocol on file;
    • Institutional Review Board review and approval;
    • Scientific review and approval by two or more qualified individuals who are not part of the research team;
    • Certification that investigators have not been disqualified.

For purposes of this coverage decision, CMS will determine whether specific registries or clinical trials meet these criteria.

  f) Providers must be able to justify the medical necessity of devices other than single lead devices. This justification should be available in the patient’s medical record.

C. Other Indications

All other indications for implantable automatic defibrillators not currently covered in accordance with this decision will continue to be covered under Category B IDE trials (42 CFR §405.201) and the CMS routine clinical trials policy (NCD Manual §310.1). (This NCD last reviewed February 2005).

Alpert and Thygesen et al., 2000. *Criteria for acute, evolving or recent MI*. Either one of the following criteria satisfies the diagnosis for an acute, evolving or recent MI:

  1) Typical rise and gradual fall (troponin) or more rapid rise and fall (CK-MB) of biochemical markers of myocardial necrosis with at least one of the following:
     a) ischemic symptoms;
     b) development of pathologic Q waves on the ECG;
     c) ECG changes indicative of ischemia (ST segment elevation or depression); or
     d) coronary artery intervention (e.g., coronary angioplasty).

  2) Pathologic findings of an acute MI.

    *Criteria for established MI*. Any one of the following criteria satisfies the diagnosis for established MI:

    1) Development of new pathologic Q waves on serial ECGs. The patient may or may not remember previous symptoms. Biochemical markers of myocardial necrosis may have normalized, depending on the length of time that has passed since the infarct developed.

    2) Pathologic findings of a healed or healing MI.

MEDICARE NCD FOR CARDIAC RESYNCHRONIZATION THERAPY PACEMAKERS (CRT-PS)

A cardiac resynchronization therapy pacemaker (CRT-P) utilizes biventricular pacing to coordinate the contraction of the ventricles with the intent of improving the hemodynamic status of the patient. This technology utilizes both conventional pacing technology as well as the addition of a third electrode that provides sensing and pacing capabilities in the left ventricle.

At this time there is no specific NCD for CRT-Ps. However, some MACs have developed Local Coverage Determinations (LCDs) for CRT-P that apply to certain regions. *It is important for medical providers to check with their local MAC for non-Medicare payer(s) to determine patient coverage and coding/billing guidelines.*
MEDICARE NCD FOR CARDIAC RESYNCHRONIZATION THERAPY DEFIBRILLATORS (CRT-D)

A cardiac resynchronization therapy defibrillator (CRT-D) utilizes biventricular pacing to coordinate the contraction of the ventricles and ICD capabilities to prevent ventricular tachyarrhythmias and ultimately the prevention of sudden cardiac death.

At this time there is no specific NCD for CRT-Ds. However, some MACs have developed Local Coverage Determinations (LCDs) for CRT-D that apply to certain regions. It is important for medical providers to check with their local MAC or non-Medicare payer(s) to determine patient coverage and coding/billing guidelines.

MEDICARE NCD FOR INTRACARDIAC ELECTROPHYSIOLOGY AND RELATED PROCEDURES

Some cardiovascular procedures, such as pacemakers and cardioverter-defibrillators, contain very clear national coverage criteria as defined by CMS. Other procedures, such as electrophysiology studies (EPS), do not have clearly defined coverage criteria at the national level. Some MACs have developed Local Coverage Determinations (LCDs) for EPS that apply to certain regions. It is important for providers to check with their local MAC or non-Medicare payer(s) to determine patient coverage and coding/billing guidelines.

Note: Search the Medicare Coverage Database on the CMS website (http://www.cms.hhs.gov/ncd/search.asp) for coverage descriptions and updates.

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7 Alpert and Thygesen et al., 2000. Criteria for acute, evolving or recent MI. Either one of the following criteria satisfies the diagnosis for an acute, evolving or recent MI:
   1) Typical rise and gradual fall (troponin) or more rapid rise and fall (CK-MB) of biochemical markers of myocardial necrosis with at least one of the following:
      a) ischemic symptoms;
      b) development of pathologic Q waves on the ECG;
      c) ECG changes indicative of ischemia (ST segment elevation or depression); or
      d) coronary artery intervention (e.g., coronary angioplasty).
   2) Pathologic findings of an acute MI.

Criteria for established MI. Any one of the following criteria satisfies the diagnosis for established MI:
   1) Development of new pathologic Q waves on serial ECGs. The patient may or may not remember previous symptoms. Biochemical markers of myocardial necrosis may have normalized, depending on the length of time that has passed since the infarct developed.
   2) Pathologic findings of a healed or healing MI.
1 Pacemakers

Pacemaker Coding Overview 1-1

Commonly Billed Pacemaker Scenarios 1-2
Pacemaker Coding Overview

Pacemaker Implant Procedure
The implant of a permanent pacemaker system requires the use of a pacemaker pulse generator and one electrode or lead for a single chamber system, or two electrodes or leads for a dual chamber system. The leads monitor and deliver electrical stimulation to the right atrium or right ventricle for a single chamber system, or both the right atrium and right ventricle for a dual chamber system. The lead(s) are inserted through the subclavian vein and are positioned in the right atrium and/or right ventricle. In some cases, the cephalic or internal jugular vein may be used as an alternative to the subclavian vein.

A STEP-BY-STEP DESCRIPTION OF A TYPICAL INITIAL PACEMAKER SYSTEM IMPLANT PROCEDURE
1. The subclavian vein is accessed and a pulse generator pocket is formed.
2. Under fluoroscopy, the pacing lead(s) are inserted into the right atrium (33206) or right ventricle (33207) for a single chamber system, or into the right atrium and right ventricle for a dual chamber system (33208).
3. Lead measurement tests, including pacing and sensing thresholds and lead impedances, are performed.
4. The pacemaker pulse generator (included in 33206, 33207, and 33208) is connected to the lead(s) that are in place.
5. Additional testing of the lead(s) is completed.
6. The lead(s) and device are secured and the pulse generator pocket is closed.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Pacemaker Scenarios

Key:

- Moderate sedation (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)

| Physician CPT® Codes | Hospital Outpatient CPT® Codes | Hospital Inpatient ICD-9-CM Codes |

1.1 Initial single chamber rate-responsive pacemaker system implant with right atrial lead

**Scenario 1.1: Physician CPT® Codes**

- 33206 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial

**Scenario 1.1: Hospital Outpatient CPT® Codes**

- 33206 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial

**Scenario 1.1: Hospital Inpatient ICD-9-CM Codes**

- 37.73 Initial insertion of transvenous lead [electrode] into atrium
  - Excludes:
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)
- 37.82 Initial insertion of single chamber device, rate responsive
  - Excludes:
    - Replacement of existing pacemaker device (37.85–37.87)

1.2 Initial single chamber rate-responsive pacemaker system implant with right ventricular lead

**Scenario 1.2: Physician CPT® Codes**

- 33207 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); ventricular

**Scenario 1.2: Hospital Outpatient CPT® Codes**

- 33207 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); ventricular
1.2 **Hospital Inpatient ICD-9-CM Codes**

- **37.71** Initial insertion of transvenous lead [electrode] into ventricle
  - **Excludes:**
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)

- **37.82** Initial insertion of single chamber device, rate responsive
  - **Excludes:**
    - Rate responsive to physiologic stimuli other than atrial rate
    - Replacement of existing pacemaker device (37.85-37.87)

### 1.3 Initial dual chamber pacemaker system implantation

#### Scenario 1.3: Physician CPT® Codes
- **33208** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial and ventricular

#### Scenario 1.3: Hospital Outpatient CPT® Codes
- **33208** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial and ventricular

#### Scenario 1.3: Hospital Inpatient ICD-9-CM Codes
- **37.83** Initial insertion of dual chamber device
  - **Excludes:**
    - Atrial ventricular sequential device
- **37.72** Initial insertion of transvenous leads [electrodes] into atrium and ventricle
  - **Excludes:**
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)

### 1.4 Initial dual chamber pacemaker insertion with external cardioversion performed prior to device implant session for treatment of patient in atrial fibrillation

#### Scenario 1.4: Physician CPT® Codes
- **33208** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial and ventricular
- **92960-59**, **51** Cardioversion, elective, electrical conversion of arrhythmia; external

#### Scenario 1.4: Hospital Outpatient CPT® Codes
- **33208** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial and ventricular
- **92960-59** Cardioversion, elective, electrical conversion of arrhythmia; external
Scenario 1.4: Hospital Inpatient ICD-9-CM Codes

37.83 Initial insertion of dual chamber device
  • Atrial ventricular sequential device
Excludes:
  • Replacement of existing pacemaker device (37.85–37.87)

37.72 Initial insertion of transvenous leads [electrodes] into atrium and ventricle
Excludes:
  • Insertion of temporary transvenous pacemaker system (37.78)
  • Replacement of atrial and/or ventricular lead(s) (37.76)

99.62 Other electric countershock of heart
  • Cardioversion: NOS, external
  • Conversion to sinus rhythm
  • Defibrillation, external electrode stimulation

1.5 Replacement of single chamber rate-responsive pulse generator

Scenario 1.5: Physician CPT® Codes

○ 33227 Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; single lead system

Scenario 1.5: Hospital Outpatient CPT® Codes

○ 33227 Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; single lead system

Scenario 1.5: Hospital Inpatient ICD-9-CM Codes

37.86 Replacement of any type pacemaker device with single chamber device, rate responsive
  • Rate responsive to physiologic stimuli other than atrial rate

1.6 Replacement of dual chamber pacemaker, insertion of new atrial lead, capping of existing atrial lead

Scenario 1.6: Physician CPT® Codes

○ 33206 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial
○ 33233-51 Removal of permanent pacemaker pulse generator only

Scenario 1.6: Hospital Outpatient CPT® Codes

○ 33206 Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); atrial
○ 33233 Removal of permanent pacemaker pulse generator only
### Scenario 1.6: Hospital Inpatient ICD-9-CM Codes

**37.76** Replacement of transvenous atrial and/or ventricular lead(s) [electrode]
- Removal or abandonment of existing transvenous or epicardial lead(s) with transvenous lead(s) replacement

**Excludes:**
- Replacement of epicardial lead [electrode] (37.74)

**37.87** Replacement of any type pacemaker device with dual chamber device
- Atrial ventricular sequential device

### 1.7 Replacement of dual chamber pacemaker, insertion of new ventricular lead, capping of existing ventricular lead

**Scenario 1.7: Physician CPT® Codes**

- **33207** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); ventricular
- **33233** Removal of permanent pacemaker pulse generator only

**Scenario 1.7: Hospital Outpatient CPT® Codes**

- **33207** Insertion of new or replacement of permanent pacemaker with transvenous electrode(s); ventricular
- **33233** Removal of permanent pacemaker pulse generator only

**Scenario 1.7: Hospital Inpatient ICD-9-CM Codes**

**37.76** Replacement of transvenous atrial and/or ventricular lead(s) [electrode]
- Removal or abandonment of existing transvenous or epicardial lead(s) with transvenous lead(s) replacement

**Excludes:**
- Replacement of epicardial lead [electrode] (37.74)

**37.87** Replacement of any type pacemaker device with dual chamber device
- Atrial ventricular sequential device

### 1.8 Replacement of dual chamber pacemaker on a pacemaker-dependent patient with temporary pacemaker insertion

**Scenario 1.8: Physician CPT® Codes**

- **33228** Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; dual lead system

Effective 2013 the National Correct Coding Initiative Edits (NCCI) no longer allow temporary pacing codes 33210-33211 to be reported with open or percutaneous cardiac procedures performed at the same patient encounter.
**Scenario 1.8: Hospital Outpatient CPT® Codes**

- **33228** Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; dual lead system

  Effective 2013 the National Correct Coding Initiative Edits (NCCI) no longer allow temporary pacing codes 33210-33211 to be reported with open or percutaneous cardiac procedures performed at the same patient encounter.

  Exceptions to NCCI edits for Hospital Services Only: Since the hospital incurs the cost for the temporary pacemaker device, for hospital billing (not physician) the NCCI edits allow a -59 modifier based on medical necessity.

  - Inserted in an emergency setting and the patient is monitored until a decision is made for an appropriate definitive surgery. The insertion of the temporary pacemaker is at a separate session and requires routine care involving regular cardiovascular assessment, level of consciousness, heart rhythm, pacer activity and hemodynamic response. Following this period of monitoring, a subsequent procedure or surgery may be performed at a separate session from the temporary pacemaker insertion.

**Scenario 1.8: Hospital Inpatient ICD-9-CM Codes**

- **37.87** Replacement of any type pacemaker device with dual chamber device
  - Atrial ventricular sequential device

- **37.78** Insertion of temporary transvenous pacemaker system
  
  **Excludes:**
  - Intraoperative cardiac pacemaker (39.64)

### 1.9 Replacement of dual chamber pulse generator

**Scenario 1.9: Physician CPT® Codes**

- **33228** Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; dual lead system

**Scenario 1.9: Hospital Outpatient CPT® Codes**

- **33228** Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; dual lead system

**Scenario 1.9: Hospital Inpatient ICD-9-CM Codes**

- **37.87** Replacement of any type pacemaker device with dual chamber device
  - Atrial ventricular sequential device

### 1.10 Upgrade from single chamber pacemaker with a ventricular lead to a dual chamber pacemaker with the addition of the right atrial lead

**Scenario 1.10: Physician CPT® Codes**

- **33214** Upgrade of implanted pacemaker system, conversion of single chamber system to dual chamber system (includes removal of previously placed pulse generator, testing of existing lead, insertion of new lead, insertion of new pulse generator)
**Scenario 1.10: Hospital Outpatient CPT® Codes**

- 33214 Upgrade of implanted pacemaker system, conversion of single chamber system to dual chamber system (includes removal of previously placed pulse generator, testing of existing lead, insertion of new lead, insertion of new pulse generator)

**Scenario 1.10: Hospital Inpatient ICD-9-CM Codes**

- 37.87 Replacement of any type pacemaker device with dual chamber device
  - Atrial ventricular sequential device
- 37.73 Initial insertion of transvenous lead [electrode] into atrium
  - Excludes:
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)

### 1.11 Insertion of one permanent transvenous pacing electrode

**Scenario 1.11: Physician CPT® Codes**

- 33216 Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator

**Scenario 1.11: Hospital Outpatient CPT® Codes**

- 33216 Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator

**Scenario 1.11: Hospital Inpatient ICD-9-CM Codes**

ICD-9-CM procedural coding specifies lead placement into atrium or ventricle:

- 37.71 Initial insertion of transvenous lead [electrode] into ventricle
  - Excludes:
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)
  or
- 37.73 Initial insertion of transvenous lead [electrode] into atrium
  - Excludes:
    - Insertion of temporary transvenous pacemaker system (37.78)
    - Replacement of atrial and/or ventricular lead(s) (37.76)

### 1.12 Insertion of two permanent transvenous pacing electrodes

**Scenario 1.12: Physician CPT® Codes**

- 33217 Insertion of two transvenous electrodes, permanent pacemaker or cardioverter-defibrillator

**Scenario 1.12: Hospital Outpatient CPT® Codes**

- 33217 Insertion of two transvenous electrodes, permanent pacemaker or cardioverter-defibrillator

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See page ii for important information about the uses and limitations of this document. See page 1-14 for Sources and Footnotes pertaining to this section.

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### Scenario 1.12: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>37.72</td>
<td>Initial insertion of transvenous leads [electrodes] into atrium and ventricle</td>
</tr>
</tbody>
</table>

**Excludes:**
- Insertion of temporary transvenous pacemaker system (37.78)
- Replacement of atrial and/or ventricular lead(s) (37.76)

### 1.13 Single lead extraction from a single lead system; pacemaker electrode

#### Scenario 1.13: Physician CPT® Codes

- 33234 Removal of transvenous pacemaker electrode(s); single lead system, atrial or ventricular

#### Scenario 1.13: Hospital Outpatient CPT® Codes

- 33234 Removal of transvenous pacemaker electrode(s); single lead system, atrial or ventricular

#### Scenario 1.13: Hospital Inpatient ICD-9-CM Code

<table>
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<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>37.77</td>
<td>Removal of lead(s) [electrode] without replacement</td>
</tr>
</tbody>
</table>

**Removal:**
- Epicardial lead (transthoracic approach)
- Transvenous lead(s)

**Excludes:**
- Removal of temporary transvenous pacemaker system – omit code
- That with replacement of:
  - atrial and/or ventricular lead(s) [electrode] (37.76)
  - epicardial lead [electrode] (37.74)

### 1.14 Repositioning of right atrial or right ventricular electrode within 90 days of implant performed by the implanting physician

#### Scenario 1.14: Physician CPT® Codes

- 33215-78 Repositioning of previously implanted transvenous pacemaker or pacing cardioverter-defibrillator (right atrial or right ventricular) electrode

#### Scenario 1.14: Hospital Outpatient CPT® Codes

- 33215-78* Repositioning of previously implanted transvenous pacemaker or pacing cardioverter-defibrillator (right atrial or right ventricular) electrode

*78 Modifier for Hospitals only applies to the same day of the original procedure.
**Scenario 1.14: Hospital Inpatient ICD-9-CM Codes**

37.75 Revision of lead [electrode]
- Repair of electrode [removal with re-insertion]
- Repositioning of lead(s) (AICD) (cardiac device) (CRT-D) (CRT-P) (defibrillator) (pacemaker) (pacing) (sensing) [electrode]
- Revision of lead NOS

**Excludes:**
- Repositioning of temporary transvenous pacemaker system – omit code

### 1.15 Single chamber pacemaker follow-up (in person)

**Scenario 1.15: Physician CPT® Codes**

93288 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

or

93279 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

**Scenario 1.15: Hospital Outpatient CPT® Codes**

93288 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

or

93279 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

**Scenario 1.15: Hospital Inpatient ICD-9-CM Codes**

89.45 Artificial pacemaker rate check
- Artificial pacemaker function check NOS
- Bedside device check of pacemaker or cardiac resynchronization pacemaker [CRT-P]
- Interrogation only without arrhythmia induction

**Excludes:**
- Catheter based invasive electrophysiologic testing (37.26)
- Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
1.16 Dual chamber pacemaker follow-up (in person)

**Scenario 1.16: Physician CPT® Codes**¹

<table>
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<tr>
<td>93288</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system</td>
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<tr>
<td>93280</td>
<td>Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; dual lead pacemaker system</td>
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**Scenario 1.16: Hospital Outpatient CPT® Codes**²

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**Scenario 1.16: Hospital Inpatient ICD-9-CM Codes**³

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<td>Artificial pacemaker rate check</td>
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</table>

- Artificial pacemaker function check NOS
- Bedside device check of pacemaker or cardiac resynchronization pacemaker [CRT-P]
- Interrogation only without arrhythmia induction

*Excludes:*

- Catheter based invasive electrophysiologic testing (37.26)
- Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
1.17 Device programming evaluation dual chamber with wound check* performed by implanting physician 14 days post-op in clinic

**Scenario 1.17: Physician CPT® Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 93280 | Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; dual lead pacemaker system  
*Wound checks are included in the 90-day global surgical package and not separately billable* |

**Scenario 1.17: Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
<th>Code</th>
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</thead>
</table>
| 93280 | Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; dual lead pacemaker system  
*Wound checks are included in the 90-day global surgical package and not separately billable* |

**Scenario 1.17: Hospital Inpatient ICD-9-CM Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>N/A</td>
<td></td>
</tr>
</tbody>
</table>
1.18 Dual chamber device follow-up – device permanently programmed VVIR due to damaged atrial lead. At same office visit, patient seen by physician for medication adjustment

**Scenario 1.18: Physician CPT® Codes**

- **93288** Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

  *or*

- **93279** Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

- **99211-99215-25** Office or other outpatient visit for the evaluation and management of an established patient (The correct level of service will depend on the documented elements; please refer to the AMA’s 2014 Current Procedural Terminology manual). Definition of -25 Modifier: Significant, Separately Identifiable Evaluation and Management Service by the Same Physician or Other Qualified Health Care Professional on the Same Day of the Procedure or Other Service.

**Scenario 1.18: Hospital Outpatient CPT® Codes**

- **93288** Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

  *or*

- **93279** Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

**Scenario 1.18: Hospital Inpatient ICD-9-CM Codes**

- **N/A**
### 1.19  Single, dual or multi chamber pacemaker follow-up (remote)

<table>
<thead>
<tr>
<th>Scenario 1.19: Physician CPT® Codes¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>93294  Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296  Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead pacemaker system or implantable defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

<table>
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<tr>
<th>Scenario 1.19: Hospital Outpatient CPT® Codes²</th>
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<td>93294  Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
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</tr>
</thead>
<tbody>
<tr>
<td>N/A</td>
</tr>
</tbody>
</table>
1.20 **Single, dual, or multi chamber pacemaker follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM)**

### Scenario 1.20: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
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</tr>
</thead>
<tbody>
<tr>
<td>93294</td>
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</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead pacemaker system or implantable defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
</tbody>
</table>

### Scenario 1.20: Hospital Outpatient CPT® Codes

<table>
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<tbody>
<tr>
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<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
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<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead pacemaker system or implantable defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
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### Scenario 1.20: Hospital Inpatient ICD-9-CM Codes

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2. As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement). Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at [http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf](http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf).


4. Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA’s 2014 *Current Procedural Terminology* for complete descriptions. Always verify appropriate usage with payers.
Implantable Cardioverter Defibrillators (ICDs)

Implantable Cardioverter-Defibrillator (ICD) Coding Overview 2-1

Commonly Billed Cardioverter-Defibrillator (ICD) Scenarios 2-2
Implantable Cardioverter-Defibrillator (ICD) Coding Overview

**ICD Implant Procedure**

The implant of an ICD system requires the use of an ICD pulse generator and a defibrillation electrode, or lead, placed in the right ventricle for a single chamber system. If a dual chamber ICD system is required, a defibrillation lead is placed in the right ventricle and a pacing electrode or lead is placed in the right atrium. The defibrillation lead delivers electrical shock therapy if a lethal arrhythmia is detected. In addition, the lead system monitors and delivers electrical pacing stimulation if required. The leads are inserted through the subclavian vein. In some cases, the cephalic or internal jugular vein may be used as an alternative to the subclavian vein.

**A STEP-BY-STEP DESCRIPTION OF A TYPICAL INITIAL ICD SYSTEM IMPLANT PROCEDURE**

1. The subclavian vein is accessed and a pulse generator pocket is formed.
2. Using fluoroscopy, a defibrillation lead is inserted into the right ventricle.
3. If implanting a dual chamber system, a pacing lead is also inserted into the right atrium under fluoroscopy.
4. Lead measurement tests, including pacing and sensing thresholds and lead impedances, are performed.
5. The ICD pulse generator (33249 includes the generator and one or two leads) is connected to the lead(s).
6. Testing of defibrillation thresholds (93641), including arrhythmia induction, is performed.
7. Additional testing of the lead(s) is completed.
8. The lead(s) and device are secured and the pulse generator pocket is closed.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Implantable Cardioverter-Defibrillator (ICD) Scenarios

Key:

- **Moderate sedation** (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)
- **Add-on code**

<table>
<thead>
<tr>
<th>Physician CPT® Codes</th>
<th>Hospital Outpatient CPT® Codes</th>
<th>Hospital Inpatient ICD-9-CM Codes</th>
</tr>
</thead>
</table>

### 2.1 Initial single or dual chamber ICD system implant, with defibrillator threshold testing at time of implant

#### Scenario 2.1: Physician CPT® Codes

- 33249: Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- 93641–26/51: Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

#### Scenario 2.1: Hospital Outpatient CPT® Codes

- 33249: Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- 93641: Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

#### Scenario 2.1: Hospital Inpatient ICD-9-CM Codes

- 37.94: Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD]

**Note:** Device testing during procedure – omit code

- Implantation of defibrillator with leads (epicardial patches), formation of pocket (abdominal fascia) (subcutaneous), any transvenous leads, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
- Techniques: lateral thoracotomy, medial sternotomy, subxiphoid procedure

**Excludes:**

- Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)
2.2 Replacement of single chamber ICD pulse generator with defibrillator threshold testing at time of replacement

**Scenario 2.2: Physician CPT® Codes**

- 33262 Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; single lead system
- 93641-26/51 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.2: Hospital Outpatient CPT® Codes**

- 33262 Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; single lead system
- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.2: Hospital Inpatient ICD-9-CM Codes**

- 37.98 Replacement of automatic cardioverter-defibrillator pulse generator only

*Note: Device testing during procedure – omit code*

*Excludes:*
- Replacement of cardiac resynchronization defibrillator, pulse generator device only [CRT-D] (00.54)
2.3 Single chamber ICD upgrade to dual chamber ICD with retention of right ventricular ICD lead and insertion of new right atrial pacing lead, and defibrillator threshold testing at the time of replacement

**Scenario 2.3: Physician CPT® Codes**

- **33249** Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- **33241-51** Removal of pacing cardioverter-defibrillator pulse generator only
- **93641-26/51** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.3: Hospital Outpatient CPT® Codes**

- **33249** Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- **33241** Removal of pacing cardioverter-defibrillator pulse generator only
- **93641** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.3: Hospital Inpatient ICD-9-CM Codes**

- **37.94** Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD]

  **Note:** Device testing during procedure – omit code
  - Implantation of defibrillator with leads (epicardial patches), formation of pocket (abdominal fascia) (subcutaneous), any transvenous leads, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
  - Techniques: lateral thoracotomy, medial sternotomy, subxiphoid procedure

**Excludes:**
- Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)
2.4 **Dual chamber pacemaker upgrade to dual chamber ICD, with capping of pacemaker leads and insertion of new right atrial and right ventricular ICD leads, with defibrillator threshold testing at the time of implant**

**Scenario 2.4: Physician CPT® Codes**

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- 33233–51 Removal of permanent pacemaker pulse generator only
- 93641-26/51 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.4: Hospital Outpatient CPT® Codes**

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- 33233 Removal of permanent pacemaker pulse generator only
- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.4: Hospital Inpatient ICD-9-CM Codes**

- 37.94 Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD]

  **Note:** Device testing during procedure – omit code

  - Implantation of defibrillator with leads (epicardial patches), formation of pocket (abdominal fascia) (subcutaneous), any transvenous leads, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
  - Techniques: lateral thoracotomy, medial sternotomy, subxiphoid procedure

  **Excludes:**

  - Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)
2.5 **Replacement of single chamber cardioverter-defibrillator lead, extraction of existing lead(s), with defibrillator threshold testing of ICD system**

### Scenario 2.5: **Physician CPT® Codes**

- **33216** Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator
- **33244** Removal of single or dual chamber pacing cardioverter-defibrillator electrode(s); by transvenous extraction
- **93641-26/51** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

### Scenario 2.5: **Hospital Outpatient CPT® Codes**

- **33216** Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator
- **33244** Removal of single or dual chamber pacing cardioverter-defibrillator electrode(s); by transvenous extraction
- **93641** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

### Scenario 2.5: **Hospital Inpatient ICD-9-CM Codes**

- **37.97** Replacement of automatic cardioverter-defibrillator lead(s) only

  **Excludes:**
  - Replacement of epicardial lead [electrode] into epicardium (37.74)
  - Replacement of transvenous lead [electrode] into left ventricular coronary venous system (00.52)
2.6 **Removal of right atrial and right ventricular leads, insertion of new right atrial and ventricular leads, with defibrillator threshold testing of ICD system**

**Scenario 2.6: Physician CPT® Codes**
- 33217 Insertion of 2 transvenous electrodes, permanent pacemaker or cardioverter-defibrillator
- 33244-51 Removal of single or dual chamber pacing cardioverter-defibrillator electrode(s); by transvenous extraction
- 93641-26/51 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.6: Hospital Outpatient CPT® Codes**
- 33217 Insertion of 2 transvenous electrodes, permanent pacemaker or cardioverter-defibrillator
- 33244 Removal of single or dual chamber pacing cardioverter-defibrillator electrode(s); by transvenous extraction
- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 2.6: Hospital Inpatient ICD-9-CM Codes**
- 37.97 Replacement of automatic cardioverter-defibrillator lead(s) only

  **Excludes:**
  - Replacement of epicardial lead [electrode] into epicardium (37.74)
  - Replacement of transvenous lead [electrode] into left ventricular coronary venous system (00.52)
2.7 **Insertion of Sub-Q Array** with **defibrillator threshold testing of ICD system**

### Scenario 2.7: **Physician CPT® Codes**

- 33999 Unlisted procedure, cardiac surgery

- 33216 Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator

  The HRS Coding Guide indicates many carriers/payers will accept existing codes for Sub-Q Array; however some carriers/payers may request use of the unlisted code. HRS recommends confirming payers requirements prior to claim submission.\(^5\)

- 93641-26/51\(^4\) Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

### Scenario 2.7: **Hospital Outpatient CPT® Codes**

- 33999 Unlisted procedure, cardiac surgery

- 33216 Insertion of a single transvenous electrode, permanent pacemaker or cardioverter-defibrillator

  The HRS Coding Guide indicates many carriers/payers will accept existing codes for Sub-Q Array; however some carriers/payers may request use of the unlisted code. HRS recommends confirming payers requirements prior to claim submission.\(^5\)

- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

### Scenario 2.7: **Hospital Inpatient ICD-9-CM Codes**

37.70 Initial insertion of lead [electrode], not otherwise specified

**Excludes:**
- Insertion of temporary transvenous pacemaker system (37.78)
- Replacement of atrial and/or ventricular lead(s) (37.76)
2.8 **Single chamber ICD follow-up (in person) in clinic**

### Scenario 2.8: Physician CPT® Codes

93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

or

93282 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead implantable cardioverter-defibrillator system

### Scenario 2.8: Hospital Outpatient CPT® Codes

93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

or

93282 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead implantable cardioverter-defibrillator system

### Scenario 2.8: Hospital Inpatient ICD-9-CM Codes

N/A
2.9 Dual chamber ICD follow-up (in person) in clinic

Scenario 2.9: Physician CPT® Codes

93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

or

93283 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; dual lead implantable cardioverter-defibrillator system

Scenario 2.9: Hospital Outpatient CPT® Codes

93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

or

93283 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; dual lead implantable cardioverter-defibrillator system

Scenario 2.9: Hospital Inpatient ICD-9-CM Codes

N/A

2.10 Single, dual or multi chamber ICD follow-up (remote)

Scenario 2.10: Physician CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

Scenario 2.10: Hospital Outpatient CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results
Scenario 2.10: Hospital Inpatient ICD-9-CM Codes

N/A

2.11 Single, dual or multi chamber ICD follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM)

Scenario 2.11: Physician CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

and

93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

Scenario 2.11: Hospital Outpatient CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

and

93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

Scenario 2.11: Hospital Inpatient ICD-9-CM Codes

N/A

2 As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at http://www.bostonscientific.com/crm/reimbursement. Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at https://www.cms.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUppdate.pdf.


4 Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA's 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.

5 Heart Rhythm Society 2012 Coding Guide for Heart Rhythm Procedures and Services, Washington, DC.
3 Subcutaneous Implantable Defibrillator (S-ICD®)

Subcutaneous Implantable Defibrillator (S-ICD) Coding Overview 3-1

Commonly Billed Subcutaneous Implantable Defibrillator (S-ICD) Scenarios 3-2
Subcutaneous Implantable Defibrillator (S-ICD) Coding Overview

A STEP-BY-STEP DESCRIPTION OF A TYPICAL INITIAL S-ICD SYSTEM IMPLANT PROCEDURE

1. Determine the ideal location for the implanted PG by placing a demo device on the patient’s skin between the 5th and 6th intercostal space in the mid-axillary line.

2. Make the device pocket incision in accordance with the ideal device location identified in step 1.

3. Locate the tip of the xyphoid process and make a 2 – 3 centimeter horizontal incision beginning at the xyphoid midline extending horizontally to the left, toward the device pocket.

4. Using an electrode insertion tool, tunnel the lead electrode from the xyphoid incision to the pocket.

5. Complete the distal electrode insertion by making a two-centimeter insertion in the sternum and tunnel the distal tip electrode up from the xyphoid to the superior incision.

6. Connect the electrode to the device header and place the device in the pocket (0319T).

7. Automatic Setup of the device is performed and the device is prepared for defibrillation testing.

8. Testing of defibrillation thresholds (0326T), including arrhythmia induction, is performed.

9. The lead(s) and device are secured and the pulse generator pocket is closed.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Subcutaneous Implantable Defibrillator (S-ICD) Scenarios

Key:

- Moderate sedation (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2013 Current Procedural Terminology for specific guidelines.)

### 3.1 Initial S-ICD system implant, with defibrillator threshold testing at time of implant

#### Scenario 3.1: Physician Category III Codes

- 0319T Insertion or replacement of subcutaneous implantable defibrillator system with subcutaneous electrode
- 0326T Electrophysiologic evaluation of subcutaneous implantable defibrillator (includes defibrillation threshold evaluation, induction of arrhythmia, evaluation of sensing for arrhythmia termination, and programming or reprogramming of sensing or therapeutic parameters)

#### Scenario 3.1: Hospital Outpatient Category III Codes

- 0319T Insertion or replacement of subcutaneous implantable defibrillator system with subcutaneous electrode
- 0326T Electrophysiologic evaluation of subcutaneous implantable defibrillator (includes defibrillation threshold evaluation, induction of arrhythmia, evaluation of sensing for arrhythmia termination, and programming or reprogramming of sensing or therapeutic parameters)

#### Scenario 3.1: Hospital Inpatient ICD-9-CM Codes

37.94 Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD]

**Note:** Device testing during procedure – omit code

- Implantation of defibrillator with leads (epicardial patches), formation of pocket (abdominal fascia) (subcutaneous), any transvenous leads, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
- Techniques: lateral thoracotomy, medial sternotomy, subxiphoid procedure

Code also extracorporeal circulation, if performed (39.61)
Code also any concomitant procedure [e.g., coronary bypass (36.10 – 36.19) or CCM, total system (17.51)]

**Excludes:**
- Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)

See page ii for important information about the uses and limitations of this document. See page 3-4 for Sources and Footnotes pertaining to this section.

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3.2 Replacement of S-ICD pulse generator using existing lead with defibrillator threshold testing

<table>
<thead>
<tr>
<th>Scenario 3.2: Physician Category III Codes¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>☐ 0323T Removal of subcutaneous implantable defibrillator pulse generator with replacement of subcutaneous implantable defibrillator pulse generator only</td>
</tr>
<tr>
<td>☐ 0326T Electrophysiologic evaluation of subcutaneous implantable defibrillator (includes defibrillation threshold evaluation, induction of arrhythmia, evaluation of sensing for arrhythmia termination, and programming or reprogramming of sensing or therapeutic parameters)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Scenario 3.2: Hospital Outpatient Category III Codes²</th>
</tr>
</thead>
<tbody>
<tr>
<td>☐ 0323T Removal of subcutaneous implantable defibrillator pulse generator with replacement of subcutaneous implantable defibrillator pulse generator only</td>
</tr>
<tr>
<td>☐ 0326T Electrophysiologic evaluation of subcutaneous implantable defibrillator (includes defibrillation threshold evaluation, induction of arrhythmia, evaluation of sensing for arrhythmia termination, and programming or reprogramming of sensing or therapeutic parameters)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Scenario 3.2: Hospital Inpatient ICD-9-CM Codes³</th>
</tr>
</thead>
<tbody>
<tr>
<td>37.98 Replacement of automatic cardioverter-defibrillator pulse generator only</td>
</tr>
</tbody>
</table>

Note: Device testing during procedure – omit code

Excludes:
• Replacement of cardiac resynchronization defibrillator, pulse generator device only [CRT-D] (00.54)
3.3 **S-ICD Follow-up (in person)**

<table>
<thead>
<tr>
<th>Scenario 3.3: Physician Category III Codes¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>0327T Interrogation device evaluation (in person) with analysis, review and report, includes connection, recording and disconnection per patient encounter; implantable subcutaneous lead defibrillator system</td>
</tr>
<tr>
<td>or</td>
</tr>
<tr>
<td>0328T Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis; implantable subcutaneous lead defibrillator system</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Scenario 3.3: Hospital Outpatient Category III Codes²</th>
</tr>
</thead>
<tbody>
<tr>
<td>0327T Interrogation device evaluation (in person) with analysis, review and report, includes connection, recording and disconnection per patient encounter; implantable subcutaneous lead defibrillator system</td>
</tr>
<tr>
<td>or</td>
</tr>
<tr>
<td>0328T Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis; implantable subcutaneous lead defibrillator system</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Scenario 3.3: Hospital Inpatient ICD-9-CM Code³</th>
</tr>
</thead>
<tbody>
<tr>
<td>89.49 Automatic implantable cardioverter-defibrillator (AICD) check</td>
</tr>
</tbody>
</table>
  - Bedside check of an AICD or cardiac resynchronization defibrillator [CRT-D]
  - Checking pacing thresholds of device
  - Interrogation only without arrhythmia induction
  Excludes:  
  - Catheter based invasive electrophysiologic testing (37.26) 
  - Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20) |

**References**


² As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement). Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at [https://www.cms.gov/HospitalOutpatientPPS/Downloads/DeviceCats OPPSUpdate.pdf](https://www.cms.gov/HospitalOutpatientPPS/Downloads/DeviceCats OPPSUpdate.pdf).


⁴ CPT Category III codes 0319T-0328T were accepted at the May 2013 AMA CPT Editorial Panel meeting for the 2014 CPT production cycle. Therefore, these codes will not appear in the 2014 CPT codebook. However, due to the Category III code early release policy, these codes are effective on January 1, 2014. Readers should reference full details, descriptions and notes for CPT Category III codes 0319T-0328T on the AMA Web site at [www.ama-assn.org/go/cpt](http://www.ama-assn.org/go/cpt).

CPT Category III codes are temporary codes that allow data collection for emerging technology, services, and procedures. These codes are intended to be used to substantiate widespread usage. CPT Category III codes are not referred to the AMA-Specialty RVS Update Committee (RUC) for valuation because no relative value units (RVUs) are assigned to Category III codes. Physician and hospital outpatient payment is based on the policies of payers and not on a yearly fee schedule.
Cardiac Resynchronization Therapy Pacemakers (CRT-Ps)

Cardiac Resynchronization Therapy Pacemaker (CRT-P) Coding
Overview 4-1

Commonly Billed Cardiac Resynchronization Therapy Pacemaker (CRT-P) Scenarios 4-2
Cardiac Resynchronization Therapy Pacemakers (CRT-P) Coding Overview

A STEP-BY-STEP DESCRIPTION OF A TYPICAL INITIAL CRT-P SYSTEM IMPLANT PROCEDURE

1. The subclavian vein is accessed and a device pocket is formed.
2. Pacing leads are inserted into the right ventricle and right atrium, under fluoroscopy.
3. A guiding catheter is inserted into the subclavian vein.
4. The coronary sinus (CS) is cannulated with the guide catheter via the coronary sinus ostium (opening).
5. In most cases, a venogram is required to visualize the coronary venous system prior to inserting the left ventricular lead.
6. A guide wire is inserted through the guide catheter, into the coronary venous system to the desired branch vein.
7. Under fluoroscopy, the left ventricular coronary venous lead is inserted (+33225) over the guide wire and advanced into a branch of the coronary venous system.
8. Lead measurement tests, including pacing and sensing thresholds and lead impedances, are performed.
9. The guide wire is removed and replaced with a finishing wire to stabilize the lead upon removal of the guide catheter.
10. The guide catheter is removed while maintaining LV lead position.
11. The finishing wire is removed and the left ventricular coronary venous lead is secured.
12. A CRT-P pulse generator (33208) is connected to the three leads that are in place.
13. Additional testing of all lead combinations is completed.
14. The leads and device are secured, and the pulse generator pocket is closed.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Cardiac Resynchronization Therapy Pacemaker (CRT-P) Scenarios

Key:

- Moderate sedation (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)
- Add-on code

### Scenario 4.1: Initial CRT-P system implant, with venogram of the coronary sinus

<table>
<thead>
<tr>
<th>Scenario 4.1: Physician CPT® Codes¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>⊗ 33208 Insertion of new or replacement of permanent pacemaker with transvenous electrodes; atrial and ventricular</td>
</tr>
<tr>
<td>+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)</td>
</tr>
</tbody>
</table>

(List separately in addition to code for primary procedure)

<table>
<thead>
<tr>
<th>Scenario 4.1: Hospital Outpatient CPT® Codes²</th>
</tr>
</thead>
<tbody>
<tr>
<td>⊗ 33208 Insertion of new or replacement of permanent pacemaker with transvenous electrodes; atrial and ventricular</td>
</tr>
<tr>
<td>+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)</td>
</tr>
</tbody>
</table>

(List separately in addition to code for primary procedure)

<table>
<thead>
<tr>
<th>Scenario 4.1: Hospital Inpatient ICD-9-CM Codes³</th>
</tr>
</thead>
<tbody>
<tr>
<td>00.50 Implantation of cardiac resynchronization pacemaker without mention of defibrillation, total system [CRT-P]</td>
</tr>
<tr>
<td><strong>Note:</strong> Device testing during procedure – omit code</td>
</tr>
<tr>
<td>• Biventricular pacemaker</td>
</tr>
<tr>
<td>• Biventricular pacing without internal cardiac defibrillator</td>
</tr>
<tr>
<td>• BiV pacemaker</td>
</tr>
<tr>
<td>• Implantation of cardiac resynchronization (biventricular) pulse generator pacing device, formation of pocket, transvenous leads including placement of lead into left ventricular coronary venous system, and intraoperative procedures for evaluation of lead signals</td>
</tr>
<tr>
<td>• That with CRT-P generator and one or more leads</td>
</tr>
<tr>
<td><strong>Excludes:</strong></td>
</tr>
<tr>
<td>• Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)</td>
</tr>
<tr>
<td>• Insertion or replacement of any type pacemaker device (37.80–37.87)</td>
</tr>
<tr>
<td>• Replacement of cardiac resynchronization:</td>
</tr>
<tr>
<td>‒ defibrillator, pulse generator only [CRT-D] (00.54)</td>
</tr>
<tr>
<td>‒ pacemaker, pulse generator only [CRT-P] (00.53)</td>
</tr>
<tr>
<td>88.63 Phlebography of other intrathoracic veins using contrast material</td>
</tr>
</tbody>
</table>

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4.2 Upgrade of dual chamber pacemaker to CRT-P system, (using existing RA and RV leads), insertion of LV lead with venogram of the coronary sinus

Scenario 4.2: Physician CPT® Codes

○ 33229 Removal of permanent pulse generator with replacement of pacemaker pulse generator; multiple lead system

+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
   (List separately in addition to code for primary procedure)

Scenario 4.2: Hospital Outpatient CPT® Codes

○ 33229 Removal of permanent pulse generator with replacement of pacemaker pulse generator; multiple lead system

+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
   (List separately in addition to code for primary procedure)

Scenario 4.2: Hospital Inpatient ICD-9-CM Codes

00.53 Implantation or replacement of cardiac resynchronization pacemaker, pulse generator only [CRT-P]
   
   Note: Device testing during procedure – omit code
   
   • Implantation of CRT-P device with removal of any existing CRT-P or other pacemaker device

Excludes:

• Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)
• Implantation or replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)
• Insertion or replacement of any type pacemaker device (37.80-37.87)

00.52 Implantation or replacement of transvenous lead [electrode] into left ventricular coronary venous system

Excludes:

• Implantation of cardiac resynchronization:
• Defibrillator, total system [CRT-D] (00.51)
• Pacemaker, total system [CRT-P] (00.50)
• Initial insertion of transvenous lead [electrode] (37.70-37.72)
• Replacement of transvenous atrial and/or ventricular lead(s) [electrodes] (37.76)

88.63 Phlebography of other intrathoracic veins using contrast material

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4.3 **Replacement of CRT-P pulse generator only utilizing existing right atrial lead, right ventricular lead and left ventricular lead**

**Scenario 4.3: Physician CPT® Codes**
- 33229 Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; multiple lead system

**Scenario 4.3: Hospital Outpatient CPT® Codes**
- 33229 Removal of permanent pacemaker pulse generator with replacement of pacemaker pulse generator; multiple lead system

**Scenario 4.3: Hospital Inpatient ICD-9-CM Codes**
- 00.53 Implantation or replacement of cardiac resynchronization pacemaker, pulse generator only [CRT-P]
  
  **Note:** Device testing during procedure – omit code
  - Implantation of CRT-P device with removal of any existing CRT-P or other pacemaker device

**Excludes:**
- Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)
- Implantation or replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)
- Insertion or replacement of any type pacemaker device (37.80–37.87)
4.4 **Single chamber pacemaker upgrade to CRT-P, with insertion of right atrial lead, and insertion of left ventricular lead with coronary sinus venogram**

### Scenario 4.4: Physician CPT® Codes

- **33214** Upgrade of implanted pacemaker system, conversion of single chamber system to dual chamber system (includes removal of previously placed pulse generator, testing of existing lead, insertion of new lead, insertion of new pulse generator)
- **33225** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
  
  (List separately in addition to code for primary procedure)

### Scenario 4.4: Hospital Outpatient CPT® Codes

- **33214** Upgrade of implanted pacemaker system, conversion of single chamber system to dual chamber system (includes removal of previously placed pulse generator, testing of existing lead, insertion of new lead, insertion of new pulse generator)
- **33225** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
  
  (List separately in addition to code for primary procedure)

### Scenario 4.4: Hospital Inpatient ICD-9-CM Codes

- **00.50** Implantation of cardiac resynchronization pacemaker without mention of defibrillation, total system [CRT-P]

  **Note:** Device testing during procedure – omit code
  
  - Biventricular pacemaker
  - Biventricular pacing without internal cardiac defibrillator
  - BiV pacemaker
  - Implantation of cardiac resynchronization (biventricular) pulse generator pacing device, formation of pocket, transvenous leads including placement of lead into left ventricular coronary venous system, and intraoperative procedures for evaluation of lead signals
  - That with CRT-P generator and one or more leads

  **Excludes:**
  
  - Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)
  - Insertion or replacement of any type pacemaker device (37.80–37.87)
  - Replacement of cardiac resynchronization:
    - defibrillator, pulse generator only [CRT-D] (00.54)
    - pacemaker, pulse generator only [CRT-P] (00.53)

- **88.63** Phlebography of other intrathoracic veins using contrast material
4.5 CRT-P follow-up (in person) in clinic

**Scenario 4.5: Physician CPT® Codes**

1. **93288** Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

   or

2. **93281** Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead pacemaker system

**Scenario 4.5: Hospital Outpatient CPT® Codes**

1. **93288** Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

   or

2. **93281** Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead pacemaker system

**Scenario 4.5: Hospital Inpatient ICD-9-CM Codes**

N/A

4.6 CRT-P follow-up (remote)

**Scenario 4.6: Physician CPT® Codes**

1. **93294** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

2. **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

**Scenario 4.6: Hospital Outpatient CPT® Codes**

1. **93294** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

2. **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results
Scenario 4.6: Hospital Inpatient ICD-9-CM Codes
N/A

4.7 CRT-P follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM) data

Scenario 4.7: Physician CPT® Codes

93294 Interrogation device evaluation(s) (remote), up to 90 days: single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days: single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

and

93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

Scenario 4.7: Hospital Outpatient CPT® Codes

93294 Interrogation device evaluation(s) (remote), up to 90 days: single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days: single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

and

93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

Scenario 4.7: Hospital Inpatient ICD-9-CM Codes
N/A


2 As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at http://www.bostonscientific.com/crm/reimbursement. Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf.


4 Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA’s 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.
5 Cardiac Resynchronization Therapy Defibrillators (CRT-Ds)

Cardiac Resynchronization Therapy Defibrillator (CRT-D) Coding Overview 5-1

Commonly Billed Cardiac Resynchronization Therapy Defibrillator (CRT-D) Scenarios 5-2
Cardiac Resynchronization Therapy Defibrillators (CRT-Ds) Coding Overview

**CRT-D Implant Procedure**

The implant of a CRT-D system typically requires the use of a cardiac resynchronization therapy pulse generator and three electrodes, or leads. The three leads monitor and deliver electrical stimulation to the right atrium, right ventricle, and left ventricle. As in conventional implantable cardioverter-defibrillator (ICD) procedures, a defibrillation lead is inserted into the subclavian vein and positioned in the right ventricle. In some cases, the cephalic or internal jugular vein may be used as an alternative to the subclavian vein. In a similar manner, a pacing lead is positioned in the right atrium. In addition, a CRT-D system requires the implantation of a third lead into the coronary venous system of the left ventricle to coordinate, or resynchronize, ventricular contractions. This left ventricular lead is inserted into the subclavian vein, introduced into the coronary sinus, and advanced into a coronary vein located on the exterior wall of the left ventricle.

**A STEP-BY-STEP DESCRIPTION OF A TYPICAL INITIAL CRT-D SYSTEM IMPLANT PROCEDURE**

1. The subclavian vein is accessed and a pulse generator pocket is formed.
2. A pacing lead is inserted into the right atrium, and the defibrillation lead is inserted into the right ventricle, under fluoroscopy.
3. A guide catheter is inserted into the subclavian vein.
4. The coronary sinus (CS) is cannulated with the guide catheter via the coronary sinus ostium (opening).
5. In most cases a venogram is required in order to visualize the coronary venous system prior to inserting the left ventricular lead.
6. A guide wire is inserted through the guide catheter, into the coronary venous system to the desired branch vein.
7. Under fluoroscopy the left ventricular lead (+33225) is positioned over the guide wire and into a branch of the coronary venous system.
8. Lead measurement tests, including pacing and sensing thresholds and lead impedances, are performed.
9. The guide wire is removed and replaced with a finishing wire to stabilize the lead upon removal of the guide catheter.
10. The guide catheter is removed, maintaining LV lead position.
11. The finishing wire is removed, and the left ventricular lead is secured.
12. A CRT-D pulse generator (33249) is connected to the three leads that are in place.
13. Testing of defibrillation thresholds (93641), including arrhythmia induction, is conducted.
14. Additional testing of all lead combinations is completed.
15. The leads and device are secured, and the pulse generator pocket is closed.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Cardiac Resynchronization Therapy Defibrillator (CRT-D) Scenarios

Key:

- **Moderate sedation** (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)

- **Add-on code**

| Physician CPT® Codes¹ | Hospital Outpatient CPT® Codes² | Hospital Inpatient ICD-9-CM Codes³ |

### 5.1 Initial CRT-D system implant, with coronary sinus venogram, with defibrillator threshold testing at the time of implant

#### Scenario 5.1: Physician CPT® Codes¹

- **33249** Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- **33225** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
  (List separately in addition to code for primary procedure)
- **93641-26/51⁴** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

#### Scenario 5.1: Hospital Outpatient CPT® Codes²

- **33249** Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber
- **33225** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)
  (List separately in addition to code for primary procedure)
- **93641** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

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5–2
### Scenario 5.1: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 00.51 | Implantation of cardiac resynchronization defibrillator, total system [CRT-D]  
**Note:** Device testing during procedure – omit code  
- BiV defibrillator  
- Biventricular defibrillator  
- Biventricular pacing with internal cardiac defibrillator  
- BiV ICD  
- BiV pacemaker with defibrillator  
- BiV pacing with defibrillator  
- Implantation of cardiac resynchronization (biventricular) pulse generator with defibrillator [AICD], formation of pocket, transvenous leads, including placement of lead into left ventricular coronary venous system, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements  
- That with CRT-D generator and one or more leads  
**Excludes:**  
- Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)  
- Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD] (37.94)  
- Replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)  
| 88.63 | Phlebography of other intrathoracic veins using contrast material |

### 5.2 Initial CRT-D implant, with atrial and ventricular lead insertion, inability to place LV lead, with defibrillator threshold testing at the time of implant

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>⊗ 33249</td>
<td>Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber</td>
</tr>
</tbody>
</table>
| ⊗ 332553 | Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)  
(List separately in addition to code for primary procedure) |
| ⊗ 93641-26/51 | Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator |
Scenario 5.2: Hospital Outpatient CPT® Codes

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber

+ 33225-74* Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (e.g., for upgrade to dual chamber system)
  (List separately in addition to code for primary procedure)
  *Modifier -74 – Discontinued outpatient procedure after anesthesia administration

- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

Scenario 5.2: Hospital Inpatient ICD-9-CM Codes

00.51 Implantation of cardiac resynchronization defibrillator, total system [CRT-D]

  Note: Device testing during procedure – omit code
  • BiV defibrillator
  • Biventricular defibrillator
  • Biventricular pacing with internal cardiac defibrillator
  • BiV ICD
  • BiV pacemaker with defibrillator
  • BiV pacing with defibrillator
  • Implantation of cardiac resynchronization (biventricular) pulse generator with defibrillator [AICD], formation of pocket, transvenous leads, including placement of lead into left ventricular coronary venous system, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
  • That with CRT-D generator and one or more leads

  Excludes:
  • Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)
  • Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD] (37.94)
  • Replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)

88.63 Phlebography of other intrathoracic veins using contrast material
5.3 **Replacement of dual lead CRT-D pulse generator with defibrillator threshold testing at the time of implant**

**Scenario 5.3: Physician CPT® Codes**

- **33263** Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; dual lead system
- **93641-26/51** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 5.3: Hospital Outpatient CPT® Codes**

- **33263** Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; dual lead system
- **93641** Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

**Scenario 5.3: Hospital Inpatient ICD-9-CM Codes**

- **00.54** Implantation or replacement of cardiac resynchronization defibrillator, pulse generator device only [CRT-D]

  **Note:** Device testing during procedure – omit code
  - Implantation of CRT-D device with removal of any existing CRT-D, CRT-P, pacemaker, or defibrillator device

**Excludes:**
- Implantation of automatic cardioverter-defibrillator, pulse generator only (37.96)
- Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)
- Implantation or replacement of cardiac resynchronization pacemaker, pulse generator only [CRT-P] (00.53)
5.4 Single or dual chamber ICD upgrade to CRT-D (capping previous RA/RV leads, placing a new RA and/or RV lead(s), left ventricular lead insertion, with coronary sinus venogram with defibrillator threshold testing at the time of implant)

Scenario 5.4: Physician CPT® Codes

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber

+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg. for upgrade to dual chamber system)

  (List separately in addition to code for primary procedure)

- 33241-51 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator

Scenario 5.4: Hospital Outpatient CPT® Codes

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber

+ 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg. for upgrade to dual chamber system)

  (List separately in addition to code for primary procedure)

- 33241 Removal of pacing cardioverter-defibrillator pulse generator only

- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator
**Scenario 5.4: Hospital Inpatient ICD-9-CM Codes**

00.51 Implantation of cardiac resynchronization defibrillator, total system [CRT-D]

**Note:** Device testing during procedure – omit code

- BiV defibrillator
- Biventricular defibrillator
- Biventricular pacing with internal cardiac defibrillator
- BiV ICD
- BiV pacemaker with defibrillator
- BiV pacing with defibrillator
- Implantation of cardiac resynchronization (biventricular) pulse generator with defibrillator [AICD], formation of pocket, transvenous leads, including placement of lead into left ventricular coronary venous system, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
- That with CRT-D generator and one or more leads

**Excludes:**

- Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)
- Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD] (37.94)
- Replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)

88.63 Phlebography of other intrathoracic veins using contrast material

### 5.5 *Dual chamber* ICD upgrade to CRT-D (using existing RA and RV leads) with left ventricular lead insertion, coronary sinus venogram with defibrillator threshold testing at the time of implant

**Scenario 5.5: Physician CPT® Codes**

- 33264 Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; multiple lead system

- 33225 Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg for upgrade to dual chamber system)
  (List separately in addition to code for primary procedure)

- 93641-26/51 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator
### Scenario 5.5: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>33264</td>
<td>Removal of pacing cardioverter-defibrillator pulse generator with replacement of pacing cardioverter-defibrillator pulse generator; multiple lead system</td>
</tr>
<tr>
<td>33225</td>
<td>Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg. for upgrade to dual chamber system) (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>93641</td>
<td>Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator</td>
</tr>
</tbody>
</table>

### Scenario 5.5: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 00.51 | Implantation of cardiac resynchronization defibrillator, total system [CRT-D]  
*Note: Device testing during procedure – omit code*  
- BiV defibrillator  
- Biventricular defibrillator  
- Biventricular pacing with internal cardiac defibrillator  
- BiV ICD  
- BiV pacemaker with defibrillator  
- BiV pacing with defibrillator  
- Implantation of cardiac resynchronization (biventricular) pulse generator with defibrillator [AICD], formation of pocket, transvenous leads, including placement of lead into left ventricular coronary venous system, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements  
- That with CRT-D generator and one or more leads  
*Excludes:*  
- Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)  
- Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD] (37.94)  
- Replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54) |
| 88.63 | Phlebography of other intrathoracic veins using contrast material |
5.6 **Insertion of left ventricular** transvenous pacing lead only, with coronary sinus venogram, LV lead inserted into previously placed CRT-D device

**Scenario 5.6: Physician CPT® Codes**

- **33224** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, with attachment to previously placed pacemaker or pacing cardioverter-defibrillator pulse generator (including revision of pocket, removal, insertion, and/or replacement of existing generator)

**Scenario 5.6: Hospital Outpatient CPT® Codes**

- **33224** Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, with attachment to previously placed pacemaker or pacing cardioverter-defibrillator pulse generator (including revision of pocket, removal, insertion, and/or replacement of existing generator)

**Scenario 5.6: Hospital Inpatient ICD-9-CM Codes**

- **00.52** Implantation or replacement of transvenous lead [electrode] into left ventricular coronary venous system

  **Excludes:**
  - Implantation of cardiac resynchronization:
    - defibrillator, total system [CRT-D] (00.51)
    - pacemaker, total system [CRT-P] (00.50)
  - Initial insertion of transvenous lead [electrode] (37.70–37.72)
  - Replacement of transvenous atrial and/or ventricular lead(s) [electrodes] (37.76)

- **88.63** Phlebography of other intrathoracic veins using contrast material
5.7 **CRT-D (3-leads) follow-up (in person)**

### Scenario 5.7: Physician CPT® Codes¹

- 93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

  or

- 93284 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead implantable cardioverter-defibrillator system

### Scenario 5.7: Hospital Outpatient CPT® Codes²

- 93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

  or

- 93284 Programming device evaluation (in person) with iterative adjustments of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead implantable cardioverter-defibrillator system

### Scenario 5.7: Hospital Inpatient ICD-9-CM Codes³

- 89.49 Automatic implantable cardioverter-defibrillator (AICD) check
  - Bedside check of an AICD or cardiac resynchronization defibrillator [CRT-D]
  - Checking pacing thresholds of device
  - Interrogation only without arrhythmia induction

  Excludes:
  - Catheter based invasive electrophysiologic testing (37.26)
  - Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
### Scenario 5.8: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
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</tr>
</thead>
<tbody>
<tr>
<td>93295</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

### Scenario 5.8: Hospital Outpatient CPT® Codes

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### Scenario 5.8: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>N/A</td>
<td>N/A</td>
</tr>
</tbody>
</table>
5.9 CRT-D follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM) data

**Scenario 5.9: Physician CPT® Codes**

<table>
<thead>
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<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
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</tbody>
</table>

and

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
</tbody>
</table>

**Scenario 5.9: Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
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</thead>
<tbody>
<tr>
<td>93295</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
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</tr>
</tbody>
</table>

**Scenario 5.9: Hospital Inpatient ICD-9-CM Codes**

N/A

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2 As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement). Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at [https://www.cms.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf](https://www.cms.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf).


4 Modifiers -26 (professional component), -51 (multiple procedures) and -53 (discontinued procedure) are for physician billing only. See the AMA’s 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.
Intracardiac Electrophysiology and Related Scenarios

Intracardiac Electrophysiology Study Coding Overview 6-1

Commonly Billed Intracardiac Electrophysiology Study Scenarios 6-2

Intracardiac Catheter Ablation Coding Overview 6-7

Commonly Billed Intracardiac Catheter Ablation Scenarios 6-8
Intracardiac Electrophysiology Study Coding Overview

**Electrophysiology (EP) Studies**

Electrophysiology (EP) studies are done to assess a patient's cardiac arrhythmias. These studies are invasive diagnostic medical procedures requiring the insertion of several electrode catheters. EP studies are done to determine if an arrhythmia is the cause of the patient’s clinical symptoms and to assess the mechanism of the cardiac arrhythmia.

EP studies “include the insertion and repositioning of electrode catheters, recording of electrograms before and during pacing or programmed stimulation of multiple locations in the heart, analysis of recorded information, and report of the procedure. Electrophysiology studies are most often performed with three or more electrode catheters.”

The studies are performed using ECG, blood pressure, and pulse oximetry monitoring. Signal processing and amplification equipment to display and assess the intracardiac electrical recordings are used.

Intracardiac electrophysiology studies are coded using a variety of CPT® codes in the 93600–93656 CPT® code range.

**A STEP-BY-STEP DESCRIPTION OF A TYPICAL COMPREHENSIVE INTRACARDIAC ELECTROPYSIOLOGY STUDY:**

1. Introducer sheaths are inserted in the femoral vein.
2. Multiple electrode catheters are inserted into the sheaths and, under fluoroscopic guidance, are advanced into the right atrium, His bundle region, and right ventricle.
3. Once in position, the electrode catheters are attached to a monitor allowing display of the intracardiac electrograms obtained from the catheter.
4. Right atrial pacing and recording, His bundle recording, and right ventricular pacing and recording are performed. The catheters may be repositioned numerous times and pacing and recording are done at various areas within the heart.
5. If an arrhythmia is induced, it may be terminated by rapidly pacing the heart or by defibrillation or cardioversion.
6. Once all pacing and recording is completed, the catheters are withdrawn and the introducer sheaths are removed.
7. The physician documents the procedure and results of the study along with any recommendations for treatment.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Intracardiac Electrophysiology Study Scenarios

Key:

© Moderate sedation (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA's 2014 Current Procedural Terminology for specific guidelines.)

<table>
<thead>
<tr>
<th>Physician CPT® Codes¹</th>
<th>Hospital Outpatient CPT® Codes²</th>
<th>Hospital Inpatient ICD-9-CM Codes³</th>
</tr>
</thead>
</table>

6.1 Comprehensive EP Study with induction or attempted induction of arrhythmia

Scenario 6.1: Physician CPT® Codes¹

★ 93620-26⁴ Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording

Scenario 6.1: Hospital Outpatient CPT® Codes²

★ 93620 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording

Scenario 6.1: Hospital Inpatient ICD-9-CM Codes³

37.26 Catheter based invasive electrophysiologic testing

• Electrophysiologic studies [EPS]
• Code also any concomitant procedure

Excludes:

• Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
• His bundle recording (37.29)
• Non-invasive programmed electrical stimulation (NIPS) (37.20)
• That as part of intraoperative testing – omit code

See page ii for important information about the uses and limitations of this document. See page 6-7 for Sources and Footnotes pertaining to this section. CPT ©2013 American Medical Association. All rights reserved. CPT is a registered trademark of the American Medical Association.
6.2 Comprehensive EP Study with induction or attempted induction of arrhythmia and dual chamber ICD implant with defibrillation threshold testing at implant

**Scenario 6.2: Physician CPT® Codes**

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous leads(s), single or dual chamber
- 93641-26/514 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator
- 93620-514 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording

**Scenario 6.2: Hospital Outpatient CPT® Codes**

- 33249 Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous leads(s), single or dual chamber
- 93641 Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator
- 93620 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording

**Scenario 6.2: Hospital Inpatient ICD-9-CM Codes**

- 37.94 Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD]
  
  **Note:** Device testing during procedure – omit code
  
  - Implantation of defibrillator with leads (epicardial patches), formation of pocket (abdominal fascia) (subcutaneous), any transvenous leads, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
  
  - Techniques: lateral thoracotomy, medial sternotomy, subxiphoid procedure

  Code also extracorporeal circulation, if performed (39.61)

  Code also any concomitant procedure [e.g., coronary bypass (36.10–36.19) or CCM, total system (17.51)]

  **Excludes:**
  
  - Implantation of cardiac resynchronization defibrillator, total system [CRT-D] (00.51)

- 37.26 Catheter based invasive electrophysiologic testing
  
  - Electrophysiologic studies [EPS]

  Code also any concomitant procedure

  **Excludes:**
  
  - That as part of intraoperative testing – omit code
  
  - Device interrogation only without arrhythmia induction (bedside check)(89.45–89.49)
  
  - His bundle recording (37.29)
**Scenario 6.2: Hospital Inpatient ICD-9-CM Codes**

- Non-invasive programmed electrical stimulation (NIPS) (37.20)

**6.3 Comprehensive EP Study** with pacing and recording of multiple sites in the right atrium, right ventricle, His bundle and left atrium with induction of arrhythmia

**Scenario 6.3: Physician CPT® Codes**

- **93620-26** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording
- **93621-26** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with left atrial pacing and recording from coronary sinus or left atrium (List separately in addition to code for primary procedure)

**Scenario 6.3: Hospital Outpatient CPT® Codes**

- **93620** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording
- **93621** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with left atrial pacing and recording from coronary sinus or left atrium (List separately in addition to code for primary procedure)

**Scenario 6.3: Hospital Inpatient ICD-9-CM Codes**

- **37.26** Catheter based invasive electrophysiologic testing
  - Electrophysiologic studies [EPS]
  - Code also any concomitant procedure
  - **Excludes:**
    - Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
    - His bundle recording (37.29)
    - Non-invasive programmed electrical stimulation (NIPS) (37.20)
    - That as part of intraoperative testing – omit code
- **37.29** Other diagnostic procedures on heart and pericardium
  - **Excludes:**
    - Angiography (88.50–88.58)
    - Cardiac function tests (89.41–89.69)
    - Cardiovascular radiotopic scan and function study (92.05)
    - Coronary arteriography (88.55–88.57)
    - Diagnostic pericardiacentesis (37.0)
    - Diagnostic ultrasound of heart (88.72)
    - X-ray of heart (87.49)
6.4 Partial (limited) EP Study – pacing and recording in the RA and His bundle

Scenario 6.4: Physician CPT® Codes

- 93600-26 Bundle of His recording
- 93602-26 Intra-atrial recording
- 93610-26 Intra-atrial pacing

Scenario 6.4: Hospital Outpatient CPT® Codes

- 93600 Bundle of His recording
- 93602 Intra-atrial recording
- 93610 Intra-atrial pacing

Scenario 6.4: Hospital Inpatient ICD-9-CM Codes

- 37.26 Catheter based invasive electrophysiologic testing
  - Electrophysiologic studies [EPS]
  - Code also any concomitant procedure
  
  Excludes:
  - Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
  - His bundle recording (37.29)
  - Non-invasive programmed electrical stimulation (NIPS) (37.20)
  - That as part of intraoperative testing – omit code

- 37.29 Other diagnostic procedures on heart and pericardium
  
  Excludes:
  - Angiocardiography (88.50–88.58)
  - Cardiac function tests (89.41–89.69)
  - Cardiovascular radioisotopic scan and function study (92.05)
  - Coronary arteriography (88.55–88.57)
  - Diagnostic pericardiocentesis (37.0)
  - Diagnostic ultrasound of heart (88.72)
  - X-ray of heart (87.49)
6.5 **Follow-up EP Study** with attempted induction of arrhythmia to assess the efficacy of medication for suppression of arrhythmia

**Scenario 6.5: Physician CPT® Codes**

- 93624-26 Electrophysiologic follow-up study with pacing and recording to test effectiveness of therapy, including induction or attempted induction of arrhythmia

**Scenario 6.5: Hospital Outpatient CPT® Codes**

- 93624 Electrophysiologic follow-up study with pacing and recording to test effectiveness of therapy, including induction or attempted induction of arrhythmia

**Scenario 6.5: Hospital Inpatient ICD-9-CM Codes**

37.26 Catheter based invasive electrophysiologic testing

- Electrophysiologic studies [EPS]
- Code also any concomitant procedure

**Excludes:**

- Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
- His bundle recording (37.29)
- Non-invasive programmed electrical stimulation (NIPS) (37.20)
- That as part of intraoperative testing – omit code

**Note:** Some of the codes presented above may be used to code for a variety of procedures (diagnostic and therapeutic) employed in the field of electrophysiology, including atrial fibrillation, atrial flutter, AV Node, SVT and VT ablations. Please note that no Boston Scientific products are approved for sale in the US for atrial fibrillation ablations.

---


2. As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement). Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at [http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf](http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf).


4. Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA’s 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.
Intracardiac Catheter Ablation Coding Overview

**Intracardiac Catheter Ablation**

Intracardiac catheter ablation is a procedure in which electrode tip catheters are placed in the heart and energy is delivered through the catheter to destroy cardiac tissue that is either causing an arrhythmia or allowing an arrhythmia to perpetuate.

The ablation catheter is placed adjacent to the cardiac tissue responsible for the arrhythmia, and the tissue is destroyed using radiofrequency electrical energy, microwave, or extreme cold temperatures (cryoablation). The ablation creates a block through which the electrical impulses can no longer cross and is intended to restore the normal electrical pathways of the heart, allowing it to beat normally again.

Arrhythmias arising in the:

- Right atrium or right ventricle are ablated with catheters placed transvenously in the appropriate cardiac chamber
- Left atrium can be ablated using a catheter placed via a retrograde aortic approach (through the aorta, across the aortic valve, and through the mitral valve) or, more commonly, via a transseptal approach (across the intra-atrial septum).

**A STEP-BY-STEP DESCRIPTION OF A TYPICAL CATHETER ABLATION:***

1. Introducer sheaths are placed in the femoral vein.
2. Under fluoroscopic guidance, multiple electrode catheters are advanced through the sheaths into the heart.
3. The catheters are attached to a recording device allowing display of the intracardiac electrograms obtained from the catheter tip.
4. An arrhythmia is induced (or attempted), and the origin of the tachycardia is confirmed and localized
5. The ablation catheter tip is moved to the arrhythmogenic focus or pathway guided by the electrical recordings and fluoroscopy.
6. Radiofrequency electrical energy, microwave energy, or cryoablation is applied to the cardiac tissue, ablating the focus or pathway.
7. Post-ablation testing is performed to verify that the tachycardia cannot be induced.
8. The catheters and sheaths are withdrawn.

Note: This document is for reference purposes only and does not replace physicians’ medical documentation. Scenarios included within this document do not encompass all possible procedures.
Commonly Billed Intracardiac Catheter Ablation Scenarios

Key:

- **Moderate sedation** (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)
- **Add-on code**

<table>
<thead>
<tr>
<th>Physician CPT® Codes</th>
<th>Hospital Outpatient CPT® Codes</th>
<th>Hospital Inpatient ICD-9-CM Codes</th>
</tr>
</thead>
</table>

### Scenario 6.6: **AV node ablation with comprehensive EP study, induction of arrhythmia, mapping and dual chamber pacemaker implant**

<table>
<thead>
<tr>
<th><strong>93620-26</strong></th>
<th>Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>93609-26</strong></td>
<td>Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td><strong>93650</strong></td>
<td>Intracardiac catheter ablation of atrioventricular node function, atrioventricular conduction for creation of complete heart block, with or without temporary pacemaker placement</td>
</tr>
<tr>
<td><strong>33208-51</strong></td>
<td>Insertion of new or replacement of permanent pacemaker with transvenous electrodes; atrial and ventricular</td>
</tr>
</tbody>
</table>

### Scenario 6.6: **Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
<th><strong>93620</strong></th>
<th>Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of arrhythmia; with right atrial pacing and recording, right ventricular pacing and recording, His bundle recording</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>93609</strong></td>
<td>Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td><strong>93650</strong></td>
<td>Intracardiac catheter ablation of atrioventricular node function, atrioventricular conduction for creation of complete heart block, with or without temporary pacemaker placement</td>
</tr>
<tr>
<td><strong>33208</strong></td>
<td>Insertion of new or replacement of permanent pacemaker with transvenous electrodes; atrial and ventricular</td>
</tr>
</tbody>
</table>
### Scenario 6.6: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 37.26 | Catheter based invasive electrophysiologic testing  
- Electrophysiologic studies [EPS]  
- Code also any concomitant procedure  
**Excludes:**  
- Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)  
- His bundle recording (37.29)  
- Non-invasive programmed electrical stimulation (NIPS) (37.20)  
- That as part of intraoperative testing – omit code |
| 37.27 | Cardiac Mapping  
- Code also any concomitant procedure  
**Excludes:**  
- Electrocardiogram (89.52)  
- His bundle recording (37.29) |
| 37.34 | Excision or destruction of other lesion or tissue of heart, endovascular approach  
- Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave) (radiofrequency) (ultrasound), via peripherally inserted catheter  
- Modified maze procedure, percutaneous approach  
**Excludes:** Ablation, excision or destruction of lesion of tissue of heart:  
- Open approach (37.33)  
- Thoracoscopic approach (37.37) |
| 37.72 | Initial insertion of transvenous leads (electrodes) into atrium and ventricle  
**Excludes:**  
- Insertion of temporary transvenous pacemaker system (37.78)  
- Replacement of atrial and/or ventricular lead(s) (37.76) |
| 37.83 | Initial insertion of dual chamber device  
- Atrial ventricular sequential device  
**Excludes:**  
- Replacement of existing pacemaker device (37.85–37.87)
### Scenario 6.7: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>33249</td>
<td>Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber</td>
</tr>
<tr>
<td>33225</td>
<td>Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)</td>
</tr>
<tr>
<td>93641-26/51</td>
<td>Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator</td>
</tr>
<tr>
<td>93650</td>
<td>Intracardiac catheter ablation of atrioventricular node function, atrioventricular conduction for creation of complete heart block, with or without temporary pacemaker placement</td>
</tr>
</tbody>
</table>

### Scenario 6.7: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>33249</td>
<td>Insertion or replacement of permanent pacing cardioverter-defibrillator system with transvenous lead(s), single or dual chamber</td>
</tr>
<tr>
<td>33225</td>
<td>Insertion of pacing electrode, cardiac venous system, for left ventricular pacing, at time of insertion of pacing cardioverter-defibrillator or pacemaker pulse generator (eg, for upgrade to dual chamber system)</td>
</tr>
<tr>
<td>93641</td>
<td>Electrophysiologic evaluation of single or dual chamber pacing cardioverter-defibrillator leads including defibrillation threshold evaluation (induction of arrhythmia, evaluation of sensing and pacing for arrhythmia termination) at time of initial implantation or replacement; with testing of single or dual chamber pacing cardioverter-defibrillator pulse generator</td>
</tr>
<tr>
<td>93650</td>
<td>Intracardiac catheter ablation of atrioventricular node function, atrioventricular conduction for creation of complete heart block, with or without temporary pacemaker placement</td>
</tr>
</tbody>
</table>
Scenario 6.7: Hospital Inpatient ICD-9-CM Codes

00.51 Implantation of cardiac resynchronization defibrillator, total system [CRT-D]

Note: Device testing during procedure – omit code

- BiV defibrillator
- Biventricular defibrillator
- Biventricular pacing with internal cardiac defibrillator
- BiV ICD
- BiV pacemaker with defibrillator
- BiV pacing with defibrillator
- Implantation of cardiac resynchronization (biventricular) pulse generator with defibrillator [AICD], formation of pocket, transvenous leads, including placement of lead into left ventricular coronary venous system, intraoperative procedures for evaluation of lead signals, and obtaining defibrillator threshold measurements
- That with CRT-D generator and one or more leads

Excludes:

- Implantation of cardiac resynchronization pacemaker, total system [CRT-P] (00.50)
- Implantation or replacement of automatic cardioverter-defibrillator, total system [AICD] (37.94)
- Replacement of cardiac resynchronization defibrillator, pulse generator only [CRT-D] (00.54)

88.63 Phlebography of other intrathoracic veins using contrast material

37.34 Excision or destruction of other lesion or tissue of heart, endovascular approach

- Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave) (radiofrequency) (ultrasound), via peripherally inserted catheter
- Modified maze procedure, percutaneous approach

Excludes:

- Ablation, excision or destruction of lesion of tissue of heart:
  - Open approach (37.33)
  - Thoracoscopic approach (37.37)
6.8 **Comprehensive Electrophysiology Study with Ablation for AVNRT (SVT Ablation) and mapping**

<table>
<thead>
<tr>
<th>Scenario 6.8: Physician CPT® Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>⚫ 93653 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and HIS bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry</td>
</tr>
</tbody>
</table>

| ⚫ + 93609| Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure) |

<table>
<thead>
<tr>
<th>Scenario 6.8: Hospital Outpatient CPT® Codes</th>
</tr>
</thead>
<tbody>
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<td>⚫ 93653 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and HIS bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry</td>
</tr>
</tbody>
</table>

| ⚫ + 93609| Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure) |
**Scenario 6.8: Hospital Inpatient ICD-9-CM Codes**

37.26 Catheter based invasive electrophysiologic testing
   - Electrophysiologic studies [EPS]
   - Code also any concomitant procedure

**Excludes:**
- Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
- His bundle recording (37.29)
- Non-invasive programmed electrical stimulation (NIPS) (37.20)
- That as part of intraoperative testing – omit code

37.34 Excision or destruction of other lesion or tissue of heart, enodovascular approach
   - Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave) (radiofrequency) (ultrasound), via peripherally inserted catheter
   - Modified maze procedure, percutaneous approach

**Excludes:**
- Ablation, excision or destruction of lesion of tissue of heart:
  - Open approach (37.33)
  - Thoracoscopic approach (37.37)

37.27 Cardiac Mapping
   - Code also any concomitant procedure

**Excludes:**
- Electrocardiogram (89.52)
- His bundle recording (37.29)

---

**6.9 Comprehensive EP study and ablation of single accessory pathway with mapping**

**Scenario 6.9: Physician CPT® Codes**

- **93653** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and HIS bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry

- **+ 93609-26** Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure)

**Scenario 6.9: Hospital Outpatient CPT® Codes**

- **93653** Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and HIS bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry

- **+ 93609** Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia
### Scenario 6.9: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 37.26 | Catheter based invasive electrophysiologic testing  
  - Electrophysiologic studies (EPS)  
  - Code also any concomitant procedure  
  **Excludes:**  
  - Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)  
  - His bundle recording (37.29)  
  - Non-invasive programmed electrical stimulation (NIPS) (37.20)  
  - That as part of intraoperative testing – omit code |
| 37.34 | Excision or destruction of other lesion or tissue of heart, endovascular approach  
  - Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave)  
  - Radiofrequency) (ultrasound), via peripherally inserted catheter  
  - Modified maze procedure, percutaneous approach  
  **Excludes:**  
  - Ablation, excision or destruction of lesion of tissue of heart:  
    - Open approach (37.33)  
    - Thoracoscopic approach (37.37) |
| 37.27 | Cardiac Mapping  
  - Code also any concomitant procedure  
  **Excludes:**  
  - Electrocardiogram (89.52)  
  - His bundle recording (37.29) |

### 6.10 SVT ablation with comprehensive EP study, mapping and intracardiac echocardiography (ICE)

**Scenario 6.10: Physician CPT® Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>⊗ 93653</td>
<td>Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and His bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry</td>
</tr>
</tbody>
</table>
| ⊗+ 93609-26⁴ | Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia  
(List separately in addition to code for primary procedure) |
| + 93662-26⁴ | Intracardiac echocardiography during therapeutic/diagnostic intervention, including imaging supervision and interpretation  
(List separately in addition to code for primary procedure) |

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**Scenario 6.10: Hospital Outpatient CPT® Codes**

⊙  93653 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording (when necessary), and HIS bundle recording (when necessary) with intracardiac catheter ablation of arrhythmogenic focus; with treatment of supraventricular tachycardia by ablation of fast or slow atrioventricular pathway, accessory atrioventricular connection, cavo-tricuspid isthmus or other single atrial focus or source of atrial re-entry

⊙ +  93609 Intraventricular and/or intra-atrial mapping of tachycardia site(s) with catheter manipulation to record from multiple sites to identify origin of tachycardia (List separately in addition to code for primary procedure)

+  93662 Intracardiac echocardiography during therapeutic/diagnostic intervention, including imaging supervision and interpretation (List separately in addition to code for primary procedure)

**Scenario 6.10: Hospital Inpatient ICD-9-CM Codes**

37.26 Catheter based invasive electrophysiologic testing
  - Electrophysiologic studies [EPS]
  - Code also any concomitant procedure
  **Excludes:**
  - Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
  - His bundle recording (37.29)
  - Non-invasive programmed electrical stimulation (NIPS) (37.20)
  - That as part of intraoperative testing – omit code

37.27 Cardiac Mapping
  - Code also any concomitant procedure
  **Excludes:**
  - Electrocardiogram (89.52)
  - His bundle recording (37.29)

37.28 Intracardiac echocardiography
  - Echocardiography of heart chambers
  - ICE
  - Code also any synchronous Doppler flow mapping (88.72)
  **Excludes:**
  - Intravascular imaging of coronary vessels (intravascular ultrasound) (IVUS) (00.24)

37.34 Excision or destruction of other lesion or tissue of heart, enodovascular approach
  - Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave) (radiofrequency) (ultrasound), via peripherally inserted catheter
  - Modified maze procedure, percutaneous approach
  **Excludes:**
  - Ablation, excision or destruction of lesion of tissue of heart:
    - Open approach (37.33)
    - Thoracoscopic approach (37.37)
6.11 **VT ablation with 3D mapping and intracardiac echocardiography (ICE)**

<table>
<thead>
<tr>
<th>Scenario 6.11: Physician CPT® Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>☉ 93654 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording, HIS recording with intracardiac catheter ablation of arrhythmogenic focus; with treatment of ventricular tachycardia or focus of ventricular ectopy including intracardiac electrophysiologic 3D mapping, when performed, and left ventricular pacing and recording, when performed</td>
</tr>
<tr>
<td>☉ + 93662-26 Intracardiac echocardiography during therapeutic/diagnostic intervention, including imaging supervision and interpretation (List separately in addition to code for primary procedure)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Scenario 6.11: Hospital Outpatient CPT® Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>☉ 93654 Comprehensive electrophysiologic evaluation including insertion and repositioning of multiple electrode catheters with induction or attempted induction of an arrhythmia with right atrial pacing and recording, right ventricular pacing and recording, HIS recording with intracardiac catheter ablation of arrhythmogenic focus; with treatment of ventricular tachycardia or focus of ventricular ectopy including intracardiac electrophysiologic 3D mapping, when performed, and left ventricular pacing and recording, when performed</td>
</tr>
<tr>
<td>☉ + 93662 Intracardiac echocardiography during therapeutic/diagnostic intervention, including imaging supervision and interpretation (List separately in addition to code for primary procedure)</td>
</tr>
</tbody>
</table>
Scenario 6.11: Hospital Inpatient ICD-9-CM Codes

37.26 Catheter based invasive electrophysiologic testing
   • Electrophysiologic studies [EPS]
   • Code also any concomitant procedure

Excludes:
   • Device interrogation only without arrhythmia induction (bedside check) (89.45–89.49)
   • His bundle recording (37.29)
   • Non-invasive programmed electrical stimulation (NIPS) (37.20)
   • That as part of intraoperative testing – omit code

37.34 Excision or destruction of other lesion or tissue of heart, enodovascular approach
   • Ablation of heart tissue (cryoablation) (electrocurrent) (laser) (microwave) (radiofrequency) (ultrasound), via peripherally inserted catheter
   • Modified maze procedure, percutaneous approach

Excludes:
   • Ablation, excision or destruction of lesion of tissue of heart:
     – Open approach (37.33)
     – Thoracoscopic approach (37.37)

37.28 Intracardiac echocardiography
   • Echocardiography of heart chambers
   • ICE
   • Code also any synchronous Doppler flow mapping (88.72)

Excludes:
   • Intravascular imaging of coronary vessels (intravascular ultrasound) (IVUS) (00.24)

37.27 Cardiac Mapping
   • Code also any concomitant procedure

Excludes:
   • Electrocardiogram (89.52)
   • His bundle recording (37.29)

Note: For transseptal puncture, use code 93462 Left heart catheterization by transseptal puncture through intact septum or by transapical puncture. List separately in addition to code for primary procedure. Use 93462 in conjunction with 93452, 93453, 93456-93461, 93653, 93654. Do NOT report 93462 in conjunction with 93656.

Note: Some of the codes presented above may be used to code for a variety of procedures (diagnostic and therapeutic) employed in the field of electrophysiology, including atrial fibrillation, atrial flutter, AV Node, SVT and VT ablations. Please note that no Boston Scientific products are approved for sale in the US for atrial fibrillation ablations.


2 As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at http://www.bostonscientific.com/crm/reimbursement. Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPSSUpdate.pdf.


4 Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA’s 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.

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7 Cardiac Device Monitoring
Commonly Billed Cardiac Device Monitoring Scenarios 7-1
Commonly Billed Cardiac Device Monitoring Scenarios

Key:

- **Moderate sedation** (For these procedures, moderate [conscious] sedation is included and cannot be billed separately when provided by the same physician. See AMA’s 2014 Current Procedural Terminology for specific guidelines.)

<table>
<thead>
<tr>
<th>Physician CPT® Codes</th>
<th>Hospital Outpatient CPT® Codes</th>
<th>Hospital Inpatient ICD-9-CM Codes</th>
</tr>
</thead>
</table>

### 7.1 Single chamber pacemaker follow-up (in person)

**Scenario 7.1: Physician CPT® Codes**

- 93288 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

  or

- 93279 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

**Scenario 7.1: Hospital Outpatient CPT® Codes**

- 93288 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

  or

- 93279 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead pacemaker system

**Scenario 7.1: Hospital Inpatient ICD-9-CM Codes**

- 89.45 Artificial pacemaker rate check
  - Artificial pacemaker function check NOS
  - Bedside device check of pacemaker or cardiac resynchronization pacemaker [CRT-P]
  - Interrogation only without arrhythmia induction

  **Excludes:**
  - Catheter based invasive electrophysiologic testing (37.26)
  - Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
7.2  **Dual chamber pacemaker follow-up (in person)**

### Scenario 7.2:  **Physician CPT® Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93288</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system</td>
</tr>
</tbody>
</table>

or

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93280</td>
<td>Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent values with analysis, review and report by a physician or other qualified health care professional; dual lead pacemaker system</td>
</tr>
</tbody>
</table>

### Scenario 7.2:  **Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93288</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system</td>
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</table>

or

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<tbody>
<tr>
<td>93280</td>
<td>Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent values with analysis, review and report by a physician or other qualified health care professional; dual lead pacemaker system</td>
</tr>
</tbody>
</table>

### Scenario 7.2:  **Hospital Inpatient ICD-9-CM Codes**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>89.45</td>
<td>Artificial pacemaker rate check</td>
</tr>
<tr>
<td></td>
<td>• Artificial pacemaker function check NOS</td>
</tr>
<tr>
<td></td>
<td>• Bedside device check of pacemaker or cardiac resynchronization pacemaker [CRT-P]</td>
</tr>
<tr>
<td></td>
<td>• Interrogation only without arrhythmia induction</td>
</tr>
</tbody>
</table>

**Excludes:**

- Catheter based invasive electrophysiologic testing (37.26)
- Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
## 7.3 Dual chamber pacemaker follow-up (remote)

### Scenario 7.3: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93294</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

### Scenario 7.3: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93294</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

### Scenario 7.3: Hospital Inpatient ICD-9-CM Codes

N/A

## 7.4 Single chamber ICD follow-up (in person)

### Scenario 7.4: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93289</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements</td>
</tr>
<tr>
<td></td>
<td>or</td>
</tr>
<tr>
<td>93282</td>
<td>Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead implantable cardioverter-defibrillator system</td>
</tr>
</tbody>
</table>

### Scenario 7.4: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93289</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements</td>
</tr>
<tr>
<td></td>
<td>or</td>
</tr>
<tr>
<td>93282</td>
<td>Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; single lead implantable cardioverter-defibrillator system</td>
</tr>
</tbody>
</table>

See page ii for important information about the uses and limitations of this document. See page 7-14 for Sources and Footnotes pertaining to this section.

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### Scenario 7.4: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>389.49</td>
<td>Automatic implantable cardioverter-defibrillator (AICD) check&lt;br&gt;• Bedside check of an AICD or cardiac resynchronization defibrillator [CRT-D]&lt;br&gt;• Checking pacing thresholds of device&lt;br&gt;• Interrogation only without arrhythmia induction&lt;br&gt;&lt;br&gt;Excludes:&lt;br&gt;• Catheter based invasive electrophysiologic testing (37.26)&lt;br&gt;• Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)</td>
</tr>
</tbody>
</table>

### 7.5 Dual chamber ICD follow-up (in person)

#### Scenario 7.5: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93289</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements&lt;br&gt;&lt;br&gt;or&lt;br&gt;&lt;br&gt;93283</td>
</tr>
</tbody>
</table>

#### Scenario 7.5: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93289</td>
<td>Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements&lt;br&gt;&lt;br&gt;or&lt;br&gt;&lt;br&gt;93283</td>
</tr>
</tbody>
</table>

#### Scenario 7.5: Hospital Inpatient ICD-9-CM Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>89.49</td>
<td>Automatic implantable cardioverter-defibrillator (AICD) check&lt;br&gt;• Bedside check of an AICD or cardiac resynchronization defibrillator [CRT-D]&lt;br&gt;• Checking pacing thresholds of device&lt;br&gt;• Interrogation only without arrhythmia induction&lt;br&gt;&lt;br&gt;Excludes:&lt;br&gt;• Catheter based invasive electrophysiologic testing (37.26)&lt;br&gt;• Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)</td>
</tr>
</tbody>
</table>
### Scenario 7.6: Physician CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93295</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

### Scenario 7.6: Hospital Outpatient CPT® Codes

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93295</td>
<td>(If applicable) Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

### Scenario 7.6: Hospital Inpatient ICD-9-CM Codes

N/A
7.7 **ICD follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM)**

### Scenario 7.7: Physician CPT® Codes

- **93295** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

- **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

- **93297** Interrogation device evaluation(s) (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

### Scenario 7.7: Hospital Outpatient CPT® Codes

- **93295** (If applicable) Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

- **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

- **93297** Interrogation device evaluation(s) (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

### Scenario 7.7: Hospital Inpatient ICD-9-CM Codes

N/A
7.8 CRT-P (3 leads) follow-up (in person)

Scenario 7.8: Physician CPT® Codes

93288-26* Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

or

93281 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead pacemaker system

Scenario 7.8: Hospital Outpatient CPT® Codes

93288 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead pacemaker system

or

93281 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead pacemaker system

Scenario 7.8: Hospital Inpatient ICD-9-CM Codes

89.45 Artificial pacemaker rate check

- Artificial pacemaker function check NOS
- Bedside device check of pacemaker or cardiac resynchronization pacemaker [CRT-P]
- Interrogation only without arrhythmia induction

Excludes:

- Catheter based invasive electrophysiologic testing (37.26)
- Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)
7.9 **CRT-P follow-up (remote)**

### Scenario 7.9: **Physician CPT® Codes**

- **93294** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional
- **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

### Scenario 7.9: **Hospital Outpatient CPT® Codes**

- **93294** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional
- **93296** Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

### Scenario 7.9: **Hospital Inpatient ICD-9-CM Codes**

- N/A
7.10 **CRT-P follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM) data**

**Scenario 7.10: Physician CPT® Codes**

- 93294 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional
- 93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results
- 93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

**Scenario 7.10: Hospital Outpatient CPT® Codes**

- 93294 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional
- 93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results
- 93297 Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional

**Scenario 7.10: Hospital Inpatient ICD-9-CM Codes**

N/A

7.11 **CRT-D (3 leads) follow-up (in person)**

**Scenario 7.11: Physician CPT® Codes**

- 93289 Interrogation device evaluation (in person) with analysis, review and report by a physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements
- 93284 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead implantable cardioverter-defibrillator system

**Scenario 7.11: Hospital Outpatient CPT® Codes**

- 93289 Interrogation device evaluation (in person) with analysis, review and report by a
physician or other qualified health care professional, includes connection, recording and disconnection per patient encounter; single, dual, or multiple lead implantable cardioverter-defibrillator system, including analysis of heart rhythm derived data elements

or

93284 Programming device evaluation (in person) with iterative adjustment of the implantable device to test the function of the device and select optimal permanent programmed values with analysis, review and report by a physician or other qualified health care professional; multiple lead implantable cardioverter-defibrillator system.

Scenario 7.11: Hospital Inpatient ICD-9-CM Codes

89.49 Automatic implantable cardioverter-defibrillator (AICD) check
  • Bedside check of an AICD or cardiac resynchronization defibrillator [CRT-D]
  • Checking pacing thresholds of device
  • Interrogation only without arrhythmia induction

Excludes:
  • Catheter based invasive electrophysiologic testing (37.26)
  • Non-invasive programmed electrical stimulation [NIPS] (arrhythmia induction) (37.20)

7.12 CRT-D follow-up (remote)

Scenario 7.12: Physician CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

Scenario 7.12: Hospital Outpatient CPT® Codes

93295 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system interim analysis, review(s) and report(s) by a physician or other qualified health care professional

93296 Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

Scenario 7.12: Hospital Inpatient ICD-9-CM Codes

N/A
7.13 **CRT-D follow-up (remote) with analysis of Implantable Cardiovascular Monitor (ICM) data**

**Scenario 7.13: Physician CPT® Codes**

<table>
<thead>
<tr>
<th>CPT Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93295</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
</tbody>
</table>

**Scenario 7.13: Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
<th>CPT Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93295</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead implantable cardioverter-defibrillator system with interim analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93296</td>
<td>Interrogation device evaluation(s) (remote), up to 90 days; single, dual, or multiple lead pacemaker system or implantable cardioverter-defibrillator system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
</tbody>
</table>

**Scenario 7.13: Hospital Inpatient ICD-9-CM Codes**

N/A

7.14 **Remote analysis of Implantable Cardiovascular Monitor (ICM)**

**Scenario 7.14: Physician CPT® Codes**

<table>
<thead>
<tr>
<th>CPT Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis, review(s) and report(s) by a physician or other qualified health care professional</td>
</tr>
<tr>
<td>93299</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system or implantable loop recorder system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results</td>
</tr>
</tbody>
</table>

**Scenario 7.14: Hospital Outpatient CPT® Codes**

<table>
<thead>
<tr>
<th>CPT Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>93297</td>
<td>Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system, including analysis of 1 or more recorded physiologic cardiovascular data elements from all internal and external sensors, analysis,</td>
</tr>
</tbody>
</table>
review(s) and report(s) by a physician or other qualified health care professional

93299  Interrogation device evaluation(s), (remote) up to 30 days; implantable cardiovascular monitor system or implantable loop recorder system, remote data acquisition(s), receipt of transmissions and technician review, technical support and distribution of results

**Scenario 7.14: Hospital Inpatient ICD-9-CM Codes**

N/A

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2 As of January 1, 2005, the Centers for Medicare and Medicaid Services (CMS) require hospitals to report all device category codes (C-codes) on Medicare outpatient claims when medical devices are used in conjunction with procedure(s) billed. If C-codes are not identified on submitted Medicare outpatient claims, the claim(s) will be returned to the hospital for correction. Find C-codes for CRM devices at [http://www.bostonscientific.com/crm/reimbursement](http://www.bostonscientific.com/crm/reimbursement). Also find C-codes for CRM devices and related accessories (e.g., introducers, catheters, sheaths) at [http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf](http://www.cms.hhs.gov/HospitalOutpatientPPS/Downloads/DeviceCats_OPPSUpdate.pdf).


4 Modifiers 26 (professional component) and 51 (multiple procedures) are for physician billing only. See the AMA’s 2014 Current Procedural Terminology for complete descriptions. Always verify appropriate usage with payers.
C-Codes

Cardiac Rhythm Management Category Codes (C-Codes) for Medical Devices 8-2

Electrophysiology Category Codes (C-Codes) for Medical Devices 8-8
Background

Ambulatory Payment Classifications (APC) refers to the hospital outpatient payment system that took effect on August 1, 2000. This system, mandated by federal law to replace the former retrospective cost-based reimbursement system, utilizes pre-set, capped payments for each APC. APCs cluster outpatient procedures into groups based on comparable resource use and clinical similarities. APCs pertain to Medicare outpatient services only and have no bearing on Medicare inpatient or physician reimbursement.
**Cardiac Rhythm Management Category Codes (C-Codes) for Medical Devices**

**Pacemakers**

**Pacemaker, dual chamber, rate-response (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALTRUA® DR</td>
<td>S205, S602, S603,</td>
<td>C1785</td>
</tr>
<tr>
<td>VITALIO™ DR</td>
<td>K273, K274</td>
<td>C1785</td>
</tr>
<tr>
<td>INGENIO™ DR</td>
<td>K173, K174</td>
<td>C1785</td>
</tr>
<tr>
<td>ADVANTIO™ DR</td>
<td>K063, K064</td>
<td>C1785</td>
</tr>
</tbody>
</table>

**Pacemaker, single chamber, rate-response (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALTRUA SR</td>
<td>S204, S601</td>
<td>C1786</td>
</tr>
<tr>
<td>VITALIO™ SR</td>
<td>K272</td>
<td>C1786</td>
</tr>
<tr>
<td>INGENIO™ SR</td>
<td>K172</td>
<td>C1786</td>
</tr>
<tr>
<td>ADVANTIO™ SR</td>
<td>K062</td>
<td>C1786</td>
</tr>
</tbody>
</table>

**Pacemaker, other than single or dual chamber (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>INVIVE™ CRT-P</td>
<td>V173, V172</td>
<td>C2621</td>
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<tr>
<td>INTUA™ CRT-P</td>
<td>V273, V272</td>
<td>C2621</td>
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## Defibrillators

**Cardioverter-defibrillator, dual chamber (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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<tbody>
<tr>
<td>ENERGEN™ DR</td>
<td>E142, E143</td>
<td>C1721</td>
</tr>
<tr>
<td>INCEPTA™ DR</td>
<td>E162, E163</td>
<td>C1721</td>
</tr>
<tr>
<td>PUNCTUA™ DR</td>
<td>E053</td>
<td>C1721</td>
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**Cardioverter-defibrillator, single chamber (implantable)**

<table>
<thead>
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<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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<tbody>
<tr>
<td>ENERGEN VR</td>
<td>E140, E141</td>
<td>C1722</td>
</tr>
<tr>
<td>INCEPTA VR</td>
<td>E160, E161</td>
<td>C1722</td>
</tr>
<tr>
<td>PUNCTUA™ VR</td>
<td>E051</td>
<td>C1722</td>
</tr>
<tr>
<td>S-ICD®</td>
<td>1010</td>
<td>C1722</td>
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**Cardioverter-defibrillator, other than single or dual chamber (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>COGNIS® HE</td>
<td>N119</td>
<td>C1882</td>
</tr>
<tr>
<td>ENERGEN™ CRT-D</td>
<td>N140, N141</td>
<td>C1882</td>
</tr>
<tr>
<td>INCEPTA™ CRT-D</td>
<td>N160, N161, N164</td>
<td>C1882</td>
</tr>
<tr>
<td>PUNCTUA™ CRT-D</td>
<td>N051</td>
<td>C1882</td>
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</tbody>
</table>
**Leads**

*Lead, cardioverter-defibrillator, endocardial single coil (implantable)*

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
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<tbody>
<tr>
<td>ENDOTAK® RELIANCE S</td>
<td>0127, 0128, 0137, 0138</td>
<td>C1777</td>
</tr>
<tr>
<td>ENDOTAK RELIANCE SG</td>
<td>0170, 0171, 0172, 0180, 0181, 0182</td>
<td>C1777</td>
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<tr>
<td>ENDOTAKRELIANCE® 4-SITE GORE</td>
<td>0282, 0283, 0292, 0293, 0294</td>
<td>C1777</td>
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</tbody>
</table>

*Lead, cardioverter-defibrillator, endocardial dual coil (implantable)*

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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</thead>
<tbody>
<tr>
<td>ENDOTAK RELIANCE</td>
<td>0147, 0148, 149, 0157, 0158, 0159</td>
<td>C1895</td>
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<tr>
<td>ENDOTAK RELIANCE G</td>
<td>0174, 0175, 0176, 0177, 0184, 0185, 0186, 0187</td>
<td>C1895</td>
</tr>
<tr>
<td>ENDOTAK RELIANCE® 4-SITE GORE</td>
<td>0285, 0286, 0295, 0296</td>
<td>C1895</td>
</tr>
<tr>
<td>ENDOTAK RELIANCE® 4-SITE Non-GORE</td>
<td>0265, 0266, 0275, 0276</td>
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</table>

*Lead, cardioverter-defibrillator, other than endocardial single or dual coil (implantable)*

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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</thead>
<tbody>
<tr>
<td>SQ Lead</td>
<td>3400</td>
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</table>
**Leads** *(continued)*

**Lead, pacemaker, other than transvenous VDD single pass**

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<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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<tbody>
<tr>
<td>FLEXTEND®</td>
<td>4086, 4087, 4088</td>
<td>C1898</td>
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<tr>
<td>FINELINE® II EZ STEROX</td>
<td>4469, 4470, 4471, 4472, 4473, 4474</td>
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<tr>
<td>FINELINE II STEROX</td>
<td>4456, 4457, 4458, 4459, 4479, 4480</td>
<td>C1898</td>
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<tr>
<td>OSCOR®</td>
<td>4039</td>
<td>C1898</td>
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<tr>
<td>Greatbatch® Epicardial</td>
<td>4046, 4047</td>
<td>C1898</td>
</tr>
<tr>
<td>DEXTRUS®</td>
<td>4135, 4136, 4137</td>
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**Lead, coronary venous**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
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<tbody>
<tr>
<td>EASYTRAK® 2</td>
<td>4517, 4518, 4520</td>
<td>C1900</td>
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<tr>
<td>EASYTRAK 2 IS-1</td>
<td>4542, 4543, 4544</td>
<td>C1900</td>
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<tr>
<td>EASYTRAK 3</td>
<td>4524, 4525, 4527</td>
<td>C1900</td>
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<tr>
<td>EASYTRAK 3 IS-1</td>
<td>4548, 4549, 4550</td>
<td>C1900</td>
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<tr>
<td>ACUITY® Steerable</td>
<td>4554, 4555</td>
<td>C1900</td>
</tr>
<tr>
<td>ACUITY Spiral</td>
<td>4591, 4592, 4593</td>
<td>C1900</td>
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</table>

**Guiding Catheters and Accessories**

**Adaptor/extension, pacing lead or neurostimulator lead (implantable)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brady Adapter</td>
<td>6161, 6162, 6125, 6986, 6987</td>
<td>C1883</td>
</tr>
<tr>
<td>Left Ventricular Lead Adapter</td>
<td>4402, 4403</td>
<td>C1883</td>
</tr>
<tr>
<td>Tachy Adapter</td>
<td>6931, 6986, 6987</td>
<td>C1883</td>
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</tbody>
</table>

See page ii for important information about the uses and limitations of this document. See page 8-20 for Sources and Footnotes pertaining to this section.
### Guiding Catheters and Accessories (continued)

#### Catheter, guiding (may include infusion/perfusion capability)

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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<tbody>
<tr>
<td><strong>ACUITY™ Break-away™ (Outer)</strong></td>
<td>7067, 7068, 7069, 7070, 7071, 7072, 7073, 7074, 7075, 7076, 7077, 7078, 7079, 7080</td>
<td>C1887</td>
</tr>
<tr>
<td><strong>ACUITY™ Break-away™ (Inner)</strong></td>
<td>7063, 7064, 7065, 7066</td>
<td>C1887</td>
</tr>
<tr>
<td><strong>ACUITY™ Cut-Away®</strong></td>
<td>7008, 7009, 7011, 7012, 7014, 7015, 7017, 7018, 7020, 7021, 7023, 7024, 7026, 7027</td>
<td>C1887</td>
</tr>
<tr>
<td><strong>ACUITY™ Cut-Away® (Inner)</strong></td>
<td>7038, 7039, 7041, 7042</td>
<td>C1887</td>
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</tbody>
</table>

#### Introducer/sheath, other than guiding, intracardiac electrophysiological, non-laser

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
<th>C-Code</th>
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</thead>
<tbody>
<tr>
<td><strong>Hemostasis Introducer</strong></td>
<td>6264, 6265, 6266, 6267, 6268, 6269, 6270, 6271, 6273, 6274, 6275, 6276, 6277, 6278, 7461, 7462, 7463, 7464, 7465, 7466, 7467, 7469, 7470, 7472, 7473, 7475</td>
<td>C1894</td>
</tr>
<tr>
<td><strong>Non-Hemostasis Introducer</strong></td>
<td>7089, 7090, 7091, 7093, 7095, 7096, 7097, 7099, 7127, 7131, 7133,</td>
<td>C1894</td>
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</table>

#### Catheter, occlusion

<table>
<thead>
<tr>
<th>Device Name</th>
<th>Model #</th>
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<tbody>
<tr>
<td><strong>Balloon Catheter</strong></td>
<td>6714, 6747</td>
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</table>

#### Guiding Catheters and Accessories (continued)

#### Guide Wire

<table>
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<tr>
<td><strong>Guide Wire</strong></td>
<td>4640, 4641, 4642, 4643, 4647, 4648, 6411, 7081, 7082,</td>
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<tr>
<td><strong>SUPPORTRAK Finishing Wire</strong></td>
<td>6667, 6668, 6669, 6681, 6682, 6683, 6684, 6685</td>
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See page ii for important information about the uses and limitations of this document. See page 8-20 for Sources and Footnotes pertaining to this section.
Electrophysiology Category Codes (C-Codes) for Medical Devices

**Catheters – Advanced Mapping**

*Catheter, electrophysiology, diagnostic/ablation, 3-D or vector mapping*

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UNP</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
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</thead>
<tbody>
<tr>
<td>CONSTELLATION® MAPPING</td>
<td>M004US8031U0</td>
<td>US8031U</td>
<td>31mm/2mm/64 electrodes</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M004US8038U0</td>
<td>US8038U</td>
<td>38mm/3mm/64 electrodes</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M004US8048U0</td>
<td>US8048U</td>
<td>48mm/4mm/64 electrodes</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M004US8060U0</td>
<td>US8060U</td>
<td>60mm/5mm/64 electrodes</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M004US58075U0</td>
<td>US8075U</td>
<td>75mm/7mm/64 electrodes</td>
<td>C1732</td>
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</tbody>
</table>

*Catheter, electrophysiology, diagnostic, 3-D mapping (19 or fewer electrodes)*

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UNP</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
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<tbody>
<tr>
<td>POLARIS X™ STEERABLE MAPPING CATHETER</td>
<td>M0047000D0</td>
<td>70000D</td>
<td>Decapolar/6F/StandardCurve/2.5mm</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M0047001D0</td>
<td>70001D</td>
<td>Decapolar/6F/StandardCurve/5mm</td>
<td>C1732</td>
</tr>
<tr>
<td></td>
<td>M0047003D0</td>
<td>70003D</td>
<td>Decapolar/6F/StandardCurve/2.5/5/2.5mm</td>
<td>C1732</td>
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<tr>
<td></td>
<td>M0047004D0</td>
<td>70004D</td>
<td>Decapolar/6F/StandardCurve/2/8/2mm</td>
<td>C1732</td>
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<tr>
<td></td>
<td>M0047005D0</td>
<td>70005D</td>
<td>Decapolar/6F/StandardCurve/2/10/2mm</td>
<td>C1732</td>
</tr>
</tbody>
</table>
Catheters – Cool-tip Ablation

Catheters, electrophysiology, diagnostic/ablation, other than 3-D or ventor mapping cool-tip

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UPN</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>CHILLI II&lt;sup&gt;®&lt;/sup&gt; COOLED 7F/4MM TIP</td>
<td>M00490310</td>
<td>90310</td>
<td>7.5F/Standard Curve 2.5mm/Standard Distal</td>
<td>C2630</td>
</tr>
<tr>
<td></td>
<td>M0049031K20</td>
<td>9031K20</td>
<td>7.5F/Large Curve 2.5mm/Standard Distal</td>
<td>C2630</td>
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<tr>
<td></td>
<td>M0049031N40</td>
<td>9031N40</td>
<td>7.5F/Asymmetric 4 Curve 2.5mm/Standard Distal</td>
<td>C2630</td>
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</table>
# Catheters – Diagnostic

**Catheters, electrophysiology, diagnostic, other than 3-D mapping (20 or more electrodes)**

<table>
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<tr>
<th>Device Name</th>
<th>UPN</th>
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<th>Description</th>
<th>C-Code</th>
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<tbody>
<tr>
<td>BLAZER® DX-20 STEERABLE CATHETER AND CABLE</td>
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<tr>
<td>M00420SL2520</td>
<td>20SL252</td>
<td>DuoDeca/7F/Super Large Curve/2/5/2mm</td>
<td>C1731</td>
<td></td>
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<tr>
<td>M00420SL2820</td>
<td>20SL282</td>
<td>DuoDeca/7F/Super Large Curve/2/8/2mm</td>
<td>C1731</td>
<td></td>
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<tr>
<td>M00420SL2220</td>
<td>20SL222</td>
<td>DuoDeca/7F/Super Large Curve/2/2/2mm</td>
<td>C1731</td>
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<tr>
<td>M00420SL5550</td>
<td>20SL555</td>
<td>DuoDeca/7F/Super Large Curve/5/5/5mm</td>
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<tr>
<td>M00420SL21020</td>
<td>20SL2102</td>
<td>DuoDeca/7F/Super Large Curve/2/10/2mm</td>
<td>C1731</td>
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<tr>
<td>M00420SL28600</td>
<td>20SL2860</td>
<td>DuoDeca/7F/Super Large Curve/2/8/2/60mm</td>
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<tr>
<td>M00420SL220250</td>
<td>20SL22025</td>
<td>DuoDeca/7F/Super Large Curve/2/20/2/25mm</td>
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<tr>
<td>M00420M2520</td>
<td>20M252</td>
<td>DuoDeca/7F/Medium Curve/2/5/2mm</td>
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<tr>
<td>M00420M2220</td>
<td>20M222</td>
<td>DuoDeca/7F/Medium Curve/2/2/2mm</td>
<td>C1731</td>
<td></td>
</tr>
<tr>
<td>M00420M255050</td>
<td>20M25505</td>
<td>DuoDeca/7F/Medium Curve/2/5/50/5mm</td>
<td>C1731</td>
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</tr>
<tr>
<td>M00420M270280</td>
<td>20M27028</td>
<td>DuoDeca/7F/Medium Curve/2/70/2/8/2mm</td>
<td>C1731</td>
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<tr>
<td>M00420M54050</td>
<td>20M5405</td>
<td>DuoDeca/7F/Medium Curve/5/40/5mm</td>
<td>C1731</td>
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<tr>
<td>M00420M28400</td>
<td>20M2840</td>
<td>DuoDeca/7F/Medium Curve/2/8/2/40mm</td>
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<tr>
<td>M00420M210350</td>
<td>20M21035</td>
<td>DuoDeca/7F/Medium Curve/2/10/2/35mm</td>
<td>C1731</td>
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</table>

See page ii for important information about the uses and limitations of this document. See page 8-20 for Sources and Footnotes pertaining to this section.
# Catheters – Fixed Curve Diagnostic

**Catheters, electrophysiology, diagnostic, other than 3-D mapping (19 or fewer electrodes)**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UPN</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>EXPLORER ST™ FIXED CURVE DIAGNOSTIC CATHETER</td>
<td>M00454180</td>
<td>5418</td>
<td>Quad/5F/Josephson Curve/5mm</td>
<td>C1730</td>
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<tr>
<td></td>
<td>M00454200</td>
<td>5420</td>
<td>Quad/5F/Cournand Curve/5mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454490</td>
<td>5449</td>
<td>Quad/5F/Cournand Curve/2/5/2mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454510</td>
<td>5451</td>
<td>Quad/5F/Josephson Curve/2/5/2mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454570</td>
<td>5457</td>
<td>Deca/6F/Cournand Curve/2/5/2mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454580</td>
<td>5458</td>
<td>Deca/6F/Cournand Curve/2mm</td>
<td>C1730</td>
</tr>
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<td></td>
<td>M00454590</td>
<td>5459</td>
<td>Deca/6F/Cournand Curve/2/8/2mm</td>
<td>C1730</td>
</tr>
<tr>
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<td>M00454690</td>
<td>5469</td>
<td>Quad/6F/Josephson Curve/10mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454700</td>
<td>5470</td>
<td>Quad/6F/Josephson Curve/5mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454770</td>
<td>5477</td>
<td>Deca/6F/Damato Curve/2/8/2mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454800</td>
<td>5480</td>
<td>Quad/6F/Cournand Curve/10mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00454810</td>
<td>5481</td>
<td>Quad/6F/Cournand Curve/5mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00455610</td>
<td>5561</td>
<td>Quad/6F/Conduction Study Curve/5mm</td>
<td>C1730</td>
</tr>
<tr>
<td></td>
<td>M00455630</td>
<td>5563</td>
<td>Quad/6F/Multipurpose Curve/5mm</td>
<td>C1730</td>
</tr>
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<td></td>
<td>M00459200</td>
<td>5920</td>
<td>Quad/6F/K-Curve/2mm</td>
<td>C1730</td>
</tr>
<tr>
<td>EXPLORER 360 JR.™ FIXED CURVE DIAGNOSTIC</td>
<td>M0045404S0</td>
<td>5404S</td>
<td>Quad/5F/Josephson Curve/5mm</td>
<td>C1730</td>
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<tr>
<td></td>
<td>M0045409S0</td>
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<td>Quad/5F/Conduction Curve/5mm</td>
<td>C1730</td>
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<tr>
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<td>Quad/5F/Josephson Curve/2mm</td>
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<td>Quad/5F/Cournand Curve/5mm</td>
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See page ii for important information about the uses and limitations of this document. See page 8-20 for Sources and Footnotes pertaining to this section.
## Catheters – Fixed Curve Diagnostic (continued)

### Catheters, electrophysiology, diagnostic, other than 3-D mapping (19 or fewer electrodes) (continued)

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### Catheters – Steerable Diagnostic

**Catheters, electrophysiology, diagnostic, other than 3-D mapping (19 or fewer electrodes) (continued)**

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## Catheters – Temperature Ablation

*Catheters, electrophysiology, diagnostic/ablation, other than 3-D or vector mapping, other than cool-tip*

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### Catheters – Temperature Ablation (continued)

**Catheters, electrophysiology, diagnostic/ablation, other than 3-D or vector mapping, other than cool-tip (continued)**

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Catheters – Temperature Ablation (continued)

Catheters, electrophysiology, diagnostic/ablation, other than 3-D or vector mapping, other than cool-tip (continued)

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### Catheters – Temperature Ablation (continued)

**Catheters, electrophysiology, diagnostic/ablation, other than 3-D or vector mapping, other than cool-tip (continued)**

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**Catheters – Temperature Ablation** (continued)

*Catheters, electrophysiology, diagnostic/ablation, other than 3-D or vector mapping, other than cool-tip (continued)*

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### Sheaths – Transseptal

#### Fixed Curve Braided Transseptal Sheath

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<td>8.5F/55MH Curve/60cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 5660120S 0</td>
<td>5660120S</td>
<td>8.5F/120MH Curve(Short)/60cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 5660120L 0</td>
<td>5660120L</td>
<td>8.5F/120MH Curve(Long)/60cm</td>
<td>C1766</td>
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<tr>
<td></td>
<td>M004 5660150 0</td>
<td>5660150</td>
<td>8.5F/150MH Curve/60cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 567915 0</td>
<td>567915</td>
<td>8.5F/15MH Curve/79.4cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 567955 0</td>
<td>567955</td>
<td>8.5F/55MH Curve/79.4cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 567990 0</td>
<td>567990</td>
<td>8.5F/90MH Curve/79.4cm</td>
<td>C1766</td>
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<tr>
<td></td>
<td>M004 5679120 0</td>
<td>5679120</td>
<td>8.5F/120MH Curve(Long)/79.4cm</td>
<td>C1766</td>
</tr>
<tr>
<td></td>
<td>M004 5610155 0</td>
<td>5610155</td>
<td>8.5F/55MH Curve/101.5cm</td>
<td>C1766</td>
</tr>
</tbody>
</table>
# Catheters – Intracardiac Echocardiography

## Catheters, intracardiac echocardiography

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UPN</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>ULTRA ICE™ CATHETER</td>
<td>M00499000</td>
<td>9900</td>
<td>Ultrasound/9F/9MH Curve/110cm</td>
<td>C1759</td>
</tr>
</tbody>
</table>
### Pericardiocentesis

**Catheter, drainage**

<table>
<thead>
<tr>
<th>Device Name</th>
<th>UPN</th>
<th>Order #</th>
<th>Description</th>
<th>C-Code</th>
</tr>
</thead>
<tbody>
<tr>
<td>PERIVAC™ PERICARDIO-CENTESIS KIT</td>
<td>M00443051</td>
<td>4305</td>
<td>Pericardiocentesis Kit W/Pigtail</td>
<td>C1729</td>
</tr>
<tr>
<td>PERIVAC™ PERICARDIO-CENTESIS KIT</td>
<td>M00443151</td>
<td>4315</td>
<td>Pericardiocentesis Kit W/Straight</td>
<td>C1729</td>
</tr>
</tbody>
</table>

Correct coding should always be verified with your Medicare Administrative Contractors and fiscal intermediary and private payers.

**Note:** Some of the codes presented above may be used to code for a variety of procedures (diagnostic and therapeutic) employed in the field of electrophysiology, including atrial fibrillation, atrial flutter, AV Node, SVT and VT ablations. Please note that no Boston Scientific products are approved for sale in the US for atrial fibrillation ablations.

Direct questions regarding hospital outpatient C-Codes for Boston Scientific CRM and EP products or other reimbursement issues to the departments below.

For questions about market-released products: 1.800.CARDIAC (227.3422), ask for the Reimbursement Customer Support Line.

For questions about investigational products: Clinical Trial Reimbursement Services, 1.800.CARDIAC (227.3422)

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Appendix

Appendix A: CPT© Modifiers
**APPENDIX A**

**CPT® Modifiers**

The list below provides modifiers applicable to CPT® 2013 codes. See the AMA's 2014 *Current Procedural Terminology* Professional Edition for full definitions.¹

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>-22</td>
<td>Increased Procedural Services</td>
</tr>
<tr>
<td>-23</td>
<td>Unusual Anesthesia</td>
</tr>
<tr>
<td>-24</td>
<td>Unrelated Evaluation and Management Service by the Same Physician or Other Qualified Health Care Professional During a Postoperative Period</td>
</tr>
<tr>
<td>-25</td>
<td>Significant, Separately Identifiable Evaluation and Management Service by the Same Physician or Other Qualified Health Care Professional on the Same Day of a Procedure or Other Service</td>
</tr>
<tr>
<td>-26</td>
<td>Professional Component</td>
</tr>
<tr>
<td>-32</td>
<td>Mandated Services</td>
</tr>
<tr>
<td>-33</td>
<td>Preventive Services</td>
</tr>
<tr>
<td>-47</td>
<td>Anesthesia by Surgeon</td>
</tr>
<tr>
<td>-50</td>
<td>Bilateral Procedure</td>
</tr>
<tr>
<td>-51</td>
<td>Multiple Procedures</td>
</tr>
<tr>
<td>-52</td>
<td>Reduced Services</td>
</tr>
<tr>
<td>-53</td>
<td>Discontinued Procedure</td>
</tr>
<tr>
<td>-54</td>
<td>Surgical Care Only</td>
</tr>
<tr>
<td>-55</td>
<td>Postoperative Management Only</td>
</tr>
<tr>
<td>-56</td>
<td>Preoperative Management Only</td>
</tr>
<tr>
<td>-57</td>
<td>Decision for Surgery</td>
</tr>
<tr>
<td>-58</td>
<td>Staged or Related Procedure or Service by the Same Physician or Other Qualified Health Care Professional During the Postoperative Period</td>
</tr>
<tr>
<td>-59</td>
<td>Distinct Procedural Service</td>
</tr>
<tr>
<td>-62</td>
<td>Two Surgeons</td>
</tr>
<tr>
<td>-63</td>
<td>Procedure Performed on Infants less than 4 kg</td>
</tr>
<tr>
<td>-66</td>
<td>Surgical Team</td>
</tr>
<tr>
<td>-76</td>
<td>Repeat Procedure or Service by Same Physician or Other Qualified Health Care Professional</td>
</tr>
<tr>
<td>-77</td>
<td>Repeat Procedure by Another Physician or Other Qualified Health Care Professional</td>
</tr>
<tr>
<td>-78</td>
<td>Unplanned Return to the Operating/Procedure Room by the Same Physician or Other Qualified Health Care Professional Following Initial Procedure for a Related Procedure During the Postoperative Period</td>
</tr>
<tr>
<td>-79</td>
<td>Unrelated Procedure or Service by the Same Physician or Other Qualified Health Care Professional During the Postoperative Period</td>
</tr>
<tr>
<td>-80</td>
<td>Assistant Surgeon</td>
</tr>
<tr>
<td>-81</td>
<td>Minimum Assistant Surgeon</td>
</tr>
<tr>
<td>-82</td>
<td>Assistant Surgeon (when qualified resident surgeon not available)</td>
</tr>
<tr>
<td>-90</td>
<td>Reference (Outside) Laboratory</td>
</tr>
<tr>
<td>-91</td>
<td>Repeat Clinical Diagnostic Laboratory Test</td>
</tr>
<tr>
<td>-92</td>
<td>Alternative Laboratory Platform Testing</td>
</tr>
<tr>
<td>-99</td>
<td>Multiple Modifiers</td>
</tr>
</tbody>
</table>

¹ See page ii for important information about the uses and limitations of this document. CPT® 2013 American Medical Association. All rights reserved. CPT is a registered trademark of the American Medical Association.
### CPT® Modifiers for Hospital Outpatient Use

The list below provides modifiers approved for hospital outpatient use (Level 1 [CPT®]). See the AMA’s 2014 Current Procedural Terminology Professional Edition for full definitions.1

<table>
<thead>
<tr>
<th>Modifier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>-25</td>
<td>Significant, Separately Identifiable Evaluation, and Management Service by the Same Physician or Other Qualified Health Care Professional on the Same Day of the Procedure or Other Service</td>
</tr>
<tr>
<td>-27</td>
<td>Multiple Outpatient Hospital E/M Encounters on the Same Date</td>
</tr>
<tr>
<td>-50</td>
<td>Bilateral Procedure</td>
</tr>
<tr>
<td>-52</td>
<td>Reduced Service</td>
</tr>
<tr>
<td>-58</td>
<td>Staged or Related Procedure or Service by the Same Physician or Other Qualified Health Care Professional During the Postoperative Period</td>
</tr>
<tr>
<td>-59</td>
<td>Distinct Procedural Service</td>
</tr>
<tr>
<td>-73</td>
<td>Discontinued Outpatient Procedure Prior to Anesthesia Administration</td>
</tr>
<tr>
<td>-74</td>
<td>Discontinued Outpatient Procedure After Anesthesia Administration</td>
</tr>
<tr>
<td>-76</td>
<td>Repeat Procedure or Service by Same Physician or Other Qualified Health Care Professional</td>
</tr>
<tr>
<td>-77</td>
<td>Repeat Procedure by Another Physician or Other Qualified Health Care Professional</td>
</tr>
<tr>
<td>-78</td>
<td>Unplanned Return to the Operating/Procedure Room by the Same Physician or Other Qualified Health Care Professional Following Initial Procedure for a Related Procedure During the Postoperative Period</td>
</tr>
<tr>
<td>-79</td>
<td>Unrelated Procedure or Service by the Same Physician or Other Qualified Health Care Professional During the Postoperative Period</td>
</tr>
<tr>
<td>-91</td>
<td>Repeat Clinical Diagnostic Laboratory Test</td>
</tr>
</tbody>
</table>

---

Disclaimer

Please note: this coding information may include some codes for procedures for which Boston Scientific currently offers no cleared or approved products. In those instances, such codes have been included solely in the interest of providing users with comprehensive coding information and are not intended to promote the use of any Boston Scientific products for which they are not cleared or approved.

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