



Medicare Critical Access Hospital Version

KEY CONCEPTS OUTLINE

Module 3: Medical Necessity and Limitation on Liability Notices

- I. Overview of Medicare Coverage
 - A. In order to be covered by Medicare, items and services must:
 1. Fall into a Medicare benefit category;
 2. Not be statutorily excluded;
 3. Be reasonable and necessary; and
 4. Meet other Medicare program requirements for payment. < *Medicare Program Integrity Manual*, Chapter 3 § 3.6.2.1 >
 - B. Coverage guidance:
 1. The Social Security Act defines Medicare benefit categories and exclusions, supplemented by regulatory guidance (e.g. 42 *C.F.R.* §§ 409, 410) and sub-regulatory guidance (e.g., the *Medicare Benefit Policy Manual*) published by CMS.
 2. In some cases, CMS publishes National Coverage Determinations (NCDs), discussed later in this module, specifying the circumstances under which an item or service is reasonable and necessary. < *Medicare Program Integrity Manual*, Chapter 3 § 3.6.2.2 >
 3. If there is no NCD, MACs may publish Local Coverage Determinations (LCDs), discussed later in this module, specifying the circumstances under which an item or service is reasonable and necessary. < *Medicare Program Integrity Manual*, Chapter 3 § 3.6.2.2 >
 4. If there is no NCD or LCD applicable to an item or service, contractors determine if it is reasonable and necessary based on the following criteria:
 - a. It is safe and effective;

- b. It is not experimental or investigational;
- c. It is appropriate, including duration and frequency;
- d. It is furnished in accordance with accepted standards of medical practice for the diagnosis or treatment of the beneficiary's condition or to improve the function of a malformed body member;
- e. It is furnished in a setting appropriate to the beneficiary's medical needs and condition;
- f. It is ordered and furnished by qualified personnel; and
- g. It meets, but does not exceed, the beneficiary's medical need. < *Medicare Program Integrity Manual*, Chapter 3 § 3.6.2.2 >

II. National and Local Coverage Policies

A. Medicare Coverage Database

1. CMS hosts a comprehensive coverage website entitled the Medicare Coverage Database where they publish National and Local Coverage Determinations and related documents.

Link: Coverage Database (NCDs, NCAs, LCDs) under Medicare-Related Sites – General

2. Types of Documents on the Medicare Coverage Database

- a. National Coverage Determinations (NCDs)
 - i. NCDs describe national Medicare coverage policy and generally provide the conditions under which an item or service is considered to be covered. < *Medicare Program Integrity Manual*, Chapter 13 § 13.1.1 >
 - ii. NCDs are binding on all Medicare contractors and in most cases on ALJs in the appeals process. < 42 *C.F.R.* 405.1060; *Medicare Program Integrity Manual*, Chapter 13 § 13.1.1 >

- b. National Coverage Analyses (NCAs) and Decision Memoranda

*CMS publishes NCAs and Decision Memoranda describing CMS coverage decisions and providing the **clinical basis and rationale** of the decisions, including **clinical evidence and studies**.*

- i. NCAs and Coverage Decision Memoranda are not binding on Medicare Contractors or ALJs, but CMS directs contractors to consider them in their medical review activities. < *Medicare Program Integrity Manual*, Chapter 12 § 13.1.1 >
- c. Coding Analyses for Labs (CALs), Medicare Evidence Development & Coverage Advisory Committee (MEDCAC) Meeting minutes, Technology Assessments (TAs) and Medicare Coverage Documents (MCDs)
 - i. CALs, MEDCAC meeting minutes, TAs, and MCDs provide additional guidance on national Medicare coverage policies and decisions.
- d. Local Coverage Determination (LCDs)
 - i. MACs publish LCDs to describe local coverage policy and as educational tools to assist and furnish guidance to providers within their jurisdiction. < *Medicare Program Integrity Manual*, Chapter 13 § 13.1.3 >
 - a) Prior to 2019, LCDs often contained CPT codes and ICD-10 codes pertinent to the policies contained in the LCD. These codes now appear in Local Coverage Articles related to the LCDs. < *Medicare Program Integrity Manual*, Chapter 13 § 13.5.1 >
 - ii. LCDs are not binding on Medicare contractors or ALJs. Regulations require contractors and ALJs to give substantial deference to LCDs applicable to a case and if they do not follow an LCD, to explain why in their decision letter. < 42 *C.F.R.* 405.1062 >
- e. Local Coverage Articles
 - i. MACs publish coverage articles addressing local coverage, coding, billing, medical review, and claims considerations. The articles may include newly developed educational materials, coding instructions, or clarification of existing billing or claims policy.
- f. Handout 5 is a copy of the MLN Booklet – How to Use the Medicare Coverage Database, June 2020.

Link: Coverage Database (NCDs, NCAs, LCDs) under Medicare-Related Sites – General

Click on How to Use the Medicare Coverage Database to download the current version

Case Study 1

Facts: A Medicare patient is scheduled for a first diagnostic joint injection procedure described by HCPCS code 64635 (Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); lumbar or sacral, single facet joint) at a pain clinic that is an outpatient department of a PPS hospital.

The following documentation is in the pain clinic electronic record for the patient:

1. Pain clinic nursing assessment showing pain level 6/10.
2. Bilateral hip, sacroiliac (SI) joint, and lower back x-rays.
3. H&P signed by the pain clinic physician detailing the intended procedure, referencing x-rays, and referencing the nursing pain assessment.

Turn to the *LCD – Facet Joint Interventions for Pain Management L34892*) and *Article – Billing and Coding: Facet Joint Interventions for Pain Management (A56670)* in the materials behind the outline and review the requirements for coverage of the procedure. What additional documentation is needed to demonstrate coverage?

B. Coverage with Evidence Development (CED)

1. CED policies cover items or services on the condition they are furnished in the context of approved clinical studies or with the collection of additional clinical data. < *Guidance for the Public, Industry, and CMS Staff; Coverage with Evidence Development Document*, Issued on November 20, 2014 >

Link: Coverage with Evidence Development (CED) under Medicare-Related Sites – General

Use links on the left navigation to access an information page for each item or service covered under CED.

2. The routine costs of items and services, associated with services covered under CED, are also covered if the items or services are generally covered for Medicare beneficiaries. < *Guidance for the Public, Industry, and CMS Staff; Coverage with Evidence Development Document*, Issued on November 20, 2014 >
3. Billing information for clinical trial reporting under CED can be found in *MLN Matters Article MM8401* and the *Medicare Claims Processing Manual*, Chapter 32 §§ 68.2, 69.5, 69.6.

C. Laboratory NCD Manual

1. CMS publishes laboratory NCDs, along with additional coding and coverage information in a "*Lab NCD Manual*" entitled *Medicare National Coverage Determination (NCD) Coding Policy Manual and Change Report, Clinical Diagnostic Laboratory Services*.
 - a. Portions of the *Lab NCD Manual* are included in the materials behind the outline for reference.

Link: Clinical Diagnostic Laboratory NCD Manual under Medicare -Related Sites - General

2. The *Lab NCD Manual* contains a list of "Non-covered ICD-10-CM Codes for All Lab NCD Edits" that are never covered by Medicare for a diagnostic laboratory service. It is not clear whether the list applies to other NCDs or to laboratory tests not covered by an NCD. <*Lab NCD Manual*>
3. The Lab NCD Manual contains national policies and MACs may not issue or maintain local policies (i.e., LCDs) that are inconsistent with the Lab NCD Manual. <Program Memorandum AB-02-110>
4. There are three "lists" of diagnosis codes applicable to each NCD
 - a. Non-Covered ICD-10-CM Codes for All NCD Edits
 - i. This is a master list set forth at the beginning of the NCD manual.
 - ii. This list applies to all NCDs and represents diagnoses for which a laboratory test covered by an NCD will never be a covered Medicare benefit. <Program Memorandum AB-02-110>
 - a) It is not clear whether CMS takes the position that the list of "ICD-10-CM codes denied" also applies to laboratory tests that do not fall within the scope of one of the laboratory NCDs.
 - iii. Tests performed for one of these diagnoses may be billed to the patient without an ABN. <Medicare Claims Processing Manual, One-Time Notification Transmittal 11>
 - a) If a test performed for one of these diagnoses is billed to Medicare, the test should be billed with the -GY modifier (discussed in a later module). <Medicare Claims Processing Manual, One Time Notification Transmittal 11>

- b. ICD-10-CM Codes Covered by Medicare
 - i. This list is set forth in the body of each NCD.
 - ii. These codes are presumed to support medical necessity. <Program Memorandum AB-02-110>
- c. ICD-10-CM Codes That Do Not Support Medical Necessity
 - i. This list is set forth in the body of each NCD.
 - a) In many cases, this list includes all diagnosis codes not included in one of the two lists discussed above.
 - ii. These codes represent diagnoses that generally do not support medical necessity but for which there may be exceptions. <Program Memorandum AB-02-110>
 - iii. Tests performed for one of these diagnoses may be billed to the patient if the patient was given an effective ABN. <Program Memorandum AB-02-110>

III. Prior Authorization

A. Overview

1. For specified services, CMS requires a prior authorization as a condition of payment. The provider must submit a request for and receive a provisional affirmation of coverage in order for the specified service to be covered and paid. <See 42 *C.F.R.* 419.82; 84 *Fed. Reg.* 61447, 85 *Fed. Reg.* 86236-248>

Although CMS refers to this process as the “prior authorization” process in regulations and other guidance, they refer to the actual approval as a “provisional affirmation”.

2. The prior authorization process only applies to services paid through Medicare Fee-for-Service and provided in hospital outpatient departments. <84 *Fed. Reg.* 61453>
 - a. Critical Access Hospital (CAH) outpatient departments are not subject to prior authorization requirements. <Prior Authorization Process for Certain Hospital Outpatient Department (OPD) Services, Frequently Asked Questions, Q9>

- b. The prior authorization process does not apply to services provided outside a hospital outpatient department (e.g., ASC or physician office) or to services paid through a Medicare Advantage plan. <84 Fed. Reg. 61453>
3. CMS has published a "Prior Authorization (PA) Program for Certain Hospital Outpatient Department (OPD) Services Operational Guide", referred to in this section as the Operational Guide, in the downloads section of the Prior Authorization for Certain Hospital Outpatient Department (OPD) Services website.

Link: Prior Authorization for Certain Hospital Outpatient Department Services – HCPCS Codes under Medicare -Related Sites - General

- B. CMS has eight categories of services requiring prior authorization:
 1. Blepharoplasty;
 2. Rhinoplasty;
 3. Panniculectomy;
 4. Botulinum toxin injections;
 - a. Prior authorization is only required when one of the listed Botulinum Toxin codes is billed with one of the listed injection codes. Botulinum Toxin billed with other procedure codes will not require prior authorization. <Operational Guide, Section 6.2.2>
 5. Vein ablation. <84 Fed. Reg. 61448, 42 C.F.R. 419.83(a)(1)>
 6. Cervical Fusion with Disc Removal; and
 7. Implanted Spinal Neurostimulators. <85 Fed. Reg. 86246-248, 42 C.F.R. 419.83(a)(2)>
 8. Facet Joint Interventions <87 Fed. Reg. 72230, 42 C.F.R. 419.83(a)(3)>

IV. Medicare's Financial Liability Protections

- A. The Limitation on Liability ("LOL") statute is a Medicare law designed to protect beneficiaries from unexpected personal liability for a non-covered service if they are unaware the service is not covered by Medicare. < *Medicare Claims Processing Manual*, Chapter 30 § 10 >

The beneficiary may not be charged for a non-covered service if:

- *The service is denied for a reason specified in the "LOL" statute; AND*
- *The beneficiary did not have advance notice Medicare would not pay for it.*

- B. Circumstances When Limitation on Liability Applies (and Advance Notice is Mandatory to Charge the Patient)

1. The item or service is not reasonable and necessary. < *Medicare Claims Processing Manual*, Chapter 30 §§ 20, 20.1 >
 - a. The item or service is not considered by Medicare to be medically necessary under the circumstances.
 - b. The service is a preventative service that is usually covered but will not be covered in this instance because frequency limitations have been exceeded.
2. The service is custodial care. < *Medicare Claims Processing Manual*, Chapter 30 § 20.1 >
3. The item or service is experimental (e.g., research use only or experimental use only laboratory tests). < *Medicare Claims Processing Manual*, Chapter 30 §§ 20.1 and 40.2.2 >

- C. Circumstances When Limitation on Liability Does Not Apply (and Advance Notice is Voluntary, Beneficiary May be Charged Without Notice)

1. Although ABNs are not required for services that are statutorily excluded from coverage or that fail to meet a technical benefit requirement, CMS "strongly encourages" providers to issue ABNs in these circumstances. < *Medicare Claims Processing Manual*, Chapter 30 § 50.2.1 >
2. The LOL provisions do not apply to items or services that fail to meet a technical benefit requirement. < *Medicare Claims Processing Manual*, Chapter 30 § 20.2 >

- a. "Technical denials" occur if coverage requirements for an item or service are not met or there is a failure to meet a condition of payment required by regulation. <Medicare Claims Processing Manual, Chapter 30 § 20.2>

Example: Denial of a drug or biological because it is usually self-administered by the patient is considered a technical denial.

3. The LOL provisions do not apply to items or services that do not fit into a Medicare benefit (i.e., is statutorily excluded). <Medicare Claims Processing Manual, Chapter 30 § 20.2>
- a. "Categorical denials" occur when the denial is based on other statutory provisions not referenced in the LOL statute. <Medicare Claims Processing Manual, Chapter 30 §§ 20.1 and 20.2>

Items or services excluded from Medicare coverage include:

- *Routine physicals and most screening tests, except the Initial Preventative Physical Exam and Annual Wellness Visit*
- *Most vaccinations, except flu, COVID, pneumococcal and hepatitis B*
- *Routine eye care, examinations and most eyeglasses*
- *Hearing aids and hearing examinations*
- *Dental care and dentures*
- *Routine foot care and flat foot care*
- *Orthopedic shoes and orthotic foot supports*
- *Cosmetic surgery and surgery performed for cosmetic purposes*

V. Handout 6 is a summary and comparison of Medicare notices.

VI. Advance Beneficiary Notice of Non-coverage (ABN)

A. General Rule

1. A properly prepared and delivered ABN form satisfies the Limitation on Liability notice requirement for outpatient services that are not considered reasonable and necessary or are custodial. <Medicare Claims Processing Manual, Chapter 30 §§ 20, 30, and 50>

B. The ABN Form

1. The Advance Beneficiary Notice (CMS-R-131 (Exp. 06/30/2023)), available in English, Spanish, and large print, is the required form for providing notice of non-coverage for outpatient services. Handout 7 is the ABN Form.

2. The ABN may not be modified except as specifically allowed in the completion instructions. < *Medicare Claims Processing Manual*, Chapter 30 § 50.5, C >

Link: Beneficiary Notice Initiative under Medicare-Related Sites – General
Use the links on the left navigation to go to the FFS ABN page.

C. Delivery of the ABN

1. The ABN should be delivered in person to the beneficiary or their representative and the provider must answer all inquiries of the beneficiary, including the basis for the determination that the service is not covered. < *Medicare Claims Processing Manual*, Chapter 30 § 50.8, 50.8.1 >
 - a. If delivery in person is not possible, delivery may be by telephone, mail, secure fax, or email. < *Medicare Claims Processing Manual*, Chapter 30 § 50.8.1 >
 - i. If notice is by telephone, a copy should be mailed, faxed or emailed to the beneficiary for them to sign and return to the provider. In order to be effective, the beneficiary must not dispute the contact. < *Medicare Claims Processing Manual*, Chapter 30 § 50.8.1 >
2. Beneficiary Comprehension
 - a. An ABN will not be considered effective unless the beneficiary, or their authorized representative, comprehends the notice. < *Medicare Claims Processing Manual*, Chapter 30 § 50.8 >
 - b. The only printed versions of the form allowed are the OMB approved English and Spanish versions, and insertions should be made in the language of the printed form. < *Medicare Claims Processing Manual*, Chapter 30 § 50.5, A >
 - c. Oral assistance should be provided for languages other than English and Spanish and documented in the “Additional Information” section. < *Medicare Claims Processing Manual*, Chapter 30 § 50.5, A >
3. Beneficiary Representative
 - a. If the patient is unable to comprehend the notice, notice must be provided to a known legal representative if the patient has one. < *Medicare Claims Processing Manual*, Chapter 30 § 50.3 >

- i. If the beneficiary does not have a representative, one may be appointed following CMS guidelines and as permitted by State and Local laws. < *Medicare Claims Processing Manual*, Chapter 30 § 50.3>
 - b. If a representative signs on behalf of the beneficiary, the name of the representative should be printed on the form and the signature should be annotated with "rep" or "representative". < *Medicare Claims Processing Manual*, Chapter 30 § 50.3>
 - c. An authorized representative is an individual authorized under State or other applicable law to act on behalf of a beneficiary when the beneficiary is temporarily or permanently unable to act for themselves (e.g., a legally appointed representative or legal guardian). < *Medicare Claims Processing Manual*, Chapter 30 § 500>
 - i. In states with health care consent statutes providing for health care decision making by surrogates for individuals who lack advance directives or guardians, it is permissible to rely on individuals designated under those statutes to act as authorized representatives. < *Medicare Claims Processing Manual*, Chapter 30 § 500>
- 4. Timing of Delivery
 - a. The ABN must be provided far enough in advance of delivery of potentially non-covered items or services to allow the beneficiary time to consider all available options and make an informed decision without undue pressure. < *Medicare Claims Processing Manual*, Chapter 30 §§ 40.2.1, 50.8>
 - i. The ABN is not effective if it is provided during an emergency, the beneficiary is under great duress, or the beneficiary is coerced or misled by the notifier, the notice, or the manner of delivery. < *Medicare Claims Processing Manual*, Chapter 30 § 40.2>
 - b. A valid ABN remains effective as long as there has been no change in:
 - i. The care described on the original ABN;
 - ii. The beneficiary's health status which would require a change in the treatment for the condition; and/or
 - iii. The Medicare coverage guidelines for the non-covered item or service (i.e., updates or changes to the coverage policy of the item or service). < *Medicare Claims Processing Manual*, Chapter 30 § 50.8, A>

- c. For items or services that are repetitive or continuous, a new ABN may be issued after one year, however, it is not required unless a change has occurred making the ABN no longer effective. < *Medicare Claims Processing Manual*, Chapter 30 § 50.8, A >

Caution: Medicare Claims Processing Manual, Chapter 30 § 40.2 continues to state that notice is not effective if delivered more than a year before the item or service is provided. This section was published in 2019 and is presumably superseded by the above guidance published in 2021.

5. Completion of the Form

Unless noted otherwise, information in this section is from the "Form Instructions, Advance Beneficiary Notice of Non-coverage (ABN), OMB Approval Number: 0938-0566" available on the FFS ABN webpage and included in the materials behind the outline.

- a. "Notifier(s)"
 - i. If the notifier in the header is an entity other than the billing entity, the notifier should annotate the Additional Information section of the ABN with information for contacting the billing entity for questions. < *Medicare Claims Processing Manual*, Chapter 30 § 50.3 >
 - ii. If multiple entities are involved in rendering or billing for the care (e.g., one entity provides the technical component and another entity provides the professional component), separate ABNs are not necessary. < *Medicare Claims Processing Manual*, Chapter 30 § 50.3 >
- b. "Blank D"
 - i. The "Blank D" field is filled in with one of the following general categories as applicable: Item, Service, Laboratory Test, Test, Procedure, Care, Equipment. All "Blank D" fields must be filled in for the ABN to valid.
 - ii. In the column under "Blank D", describe the specific item or service that is non-covered, including the frequency or duration of repetitive or continuous services. Items can be grouped, e.g., "wound care supplies" or "observation services" rather than listed individually.

c. "Reason Medicare May Not Pay:"

- i. Explain the reason the item may not be covered by Medicare.
- ii. Simply stating "medically unnecessary" or the equivalent is not acceptable. <Medicare Claims Processing Manual, Chapter 30 § 40.2.1, C>

Tip: Be specific about the reason for denial, for example:

- "Medicare does not pay for custodial care, except for some hospice services"
- "Medicare does not pay for this test for your condition".

d. "Estimated Cost"

- i. Provide a good faith estimate of the cost of the non-covered services to the patient. The cost to the patient is the provider's usual and customary charge and is not limited by the Medicare allowable or payment amount.

Caution: The final amount billed to the patient may be affected by state laws requiring providers to give uninsured patients a discount, including discounts based on financial need or equal to the discount given to their largest payer.

- a) An estimate will be considered to be made in good faith if the estimate is within the greater of \$100 or 25% of the cost of the service to the patient (i.e., amount billed to the patient) and may be given as a range or may exceed the final amount billed.

Examples of good faith cost estimates for a service with a \$1000 charge:

- Any estimate greater than \$750
- Between \$750 - \$1100
- No more than \$1200

- ii. Multiple services may be grouped together into a single cost estimate.
- iii. An average daily cost estimate may be provided for complex projections (i.e., observation services).

iv. Unknown costs

- a) The hospital may not have a policy of routinely or frequently failing to provide a cost estimate, however, the patient may sign an ABN without a cost estimate in limited circumstances.
 - 1) If additional services may be required (i.e., reflexive testing), the cost of the initial services should be given, along with a notation that additional services may be provided.
 - 2) If the costs cannot be determined, make a notation in the cost estimate area that no cost estimate is available.

Case Study 2

Facts: A Medicare patient presents to the CAH for a CT scan of the head without and with contrast. Under the applicable Medicare coverage policy, the test was not considered medically necessary for the patient's condition. The patient signed an ABN before the procedure was performed. The ABN was properly prepared and the CAH provided an estimated cost of \$2,000 - \$2,200 for the intended procedure.

The CAH performed the procedure; however, the patient required a more expensive contrast than what is normally used, resulting in a total charge of \$2,400 for the procedure. The CAH billed Medicare and Medicare denied coverage. The patient was billed \$2,400 for the CT scan.

The patient called the hospital and said she is only going to pay \$850. She claims that the hospital is overcharging her because she found out from the Medicare beneficiary services hotline that Medicare typically only pays \$850 for the service.

- Assuming no state laws affect the amount to be collected by the provider, how much may the CAH collect from the patient?

e. "Options"

- i. The beneficiary or their representative must check one of the options or have the provider check the option if they are unable to do so.

- a) The provider should make a note on the ABN if they checked the option at the request of the beneficiary.
- ii. If the beneficiary refuses to choose an option, the ABN should be annotated with the refusal and the annotation should be witnessed. < Medicare Claims Processing Manual, Chapter 30 § 40.2.2, B and 50.6, A.2.>
- iii. Special Instructions for Dually Eligible Beneficiary
 - a) Dually eligible beneficiaries have both Medicare and Medicaid, including patients enrolled in a Qualified Medicare Beneficiary (QMB) Program.
 - b) For dually eligible beneficiaries, Option 1 must be modified by lining through certain language as designated in the "Form Instructions", included in the materials behind the outline. This is an exception to the general prohibition on modifying the ABN form.
 - c) Dually eligible beneficiaries should be instructed to choose Option 1 in order for the claim to be submitted for Medicare adjudication and, if denied, submitted to Medicaid for a determination.
 - 1) If both Medicare and Medicaid deny coverage, refer to the "Form Instructions" or MLN Booklet *Dually Eligible Beneficiaries under Medicare and Medicaid*, included in the materials behind the outline, for more information on potential beneficiary liability.
 - iv. If there are multiple items on the ABN and the beneficiary wants to select different options for each of the items, more than one ABN should be used to accommodate the beneficiary's choices.
- f. "Additional Information"
 - i. May be used for witness signatures or to make annotations, such as advising the beneficiary to notify their provider of tests or services that were ordered but not received. If items are added after the date of the ABN, they must be dated.
- g. "Signature"
 - i. The beneficiary or their representative should sign and date the notice.

- ii. If the beneficiary refuses to sign but still desires to receive the item or service, the ABN should be annotated with the refusal and the annotation should be witnessed. < *Medicare Claims Processing Manual*, Chapter 30 § 40.2.2, B and 50.6, A.2.>

Case Study 3

Facts: A Medicare patient presented to the hospital for a minor non-cosmetic surgical procedure related to varicose veins. Under an applicable LCD, the procedure is not considered medically necessary for patients with a diagnosis of varicose veins. A properly prepared ABN was reviewed with the patient.

The patient refused to sign the ABN, but still wants to proceed with the procedure based on the recommendation of his physician. Two witnesses acknowledged in writing on the ABN form that the ABN had been reviewed with the patient, but that he refused to sign it. May the hospital bill the patient for the procedure, even though they did not sign the ABN?

6. Copy of the ABN

- a. The hospital should retain the original ABN and give a copy to the beneficiary. < *Medicare Claims Processing Manual*, Chapter 30 §§ 40.2.1 B and 50.5 C>
 - i. The ABN should be retained for 5 years, or longer as required by state law. < *Medicare Claims Processing Manual*, Chapter 30 § 50.7>

Caution: The ABN should be retained even if the beneficiary refuses the service or refuses to sign or choose an option.

- b. Carbon copies, fax copies, electronically scanned copies, and photocopies are all acceptable. < *Medicare Claims Processing Manual*, Chapter 30 §§ 40.2.1 B and 50.5 C>

D. Other Considerations for an Effective ABN

1. Interplay between the ABN and EMTALA requirements

- a. EMTALA Requirements Take Priority over ABN Requirements

- i. Under the Emergency Medical Treatment and Active Labor Act (EMTALA) hospitals have an obligation to complete a medical screening examination (MSE) and stabilize a patient presenting to its emergency department, or in certain circumstance, presenting to other areas of the hospital. < *Medicare Claims Processing Manual*, Chapter 30 § 40.4>
 - a) CMS and the OIG take the position that where EMTALA applies, it is improper to present an ABN to a patient before completing the MSE and stabilizing the patient. < *Medicare Claims Processing Manual*, Chapter 30 § 40.4>
 - b. Contractor’s Medical Necessity Determinations for EMTALA required care
 - i. The MAC is required to make medical necessity determinations of EMTALA screening/stabilization services based on the “information available to the treating physician or practitioner (including the patient’s presenting symptoms or complaint) at the time the item or service was ordered or furnished”. < *Social Security Act § 1862 (d)*>
 - ii. The Intermediary should not apply frequency edits to EMTALA screening/stabilization services. < *Social Security Act § 1862 (d)*>
2. Medicare Advantage Plan Beneficiaries
- a. The ABN form may not be used for services provided under Medicare Advantage Plans. < *Medicare Claims Processes Manual*, Chapter 30 § 50.1>
3. Prohibition on Routine, Blanket and Generic ABNs
- a. In general, “generic” ABNs (i.e. merely stating denial is possible”), “routine,” ABNs (i.e. no specific reason Medicare will not pay), and “blanket” ABNs (i.e. given for all claims) will not be considered to be effective. < *Medicare Claims Processing Manual*, Chapter 30 § 40.2.2 C>

Routine ABNs may be given for frequency limited service (e.g., screening mammography) if the ABN states the frequency limitation (e.g., “Medicare does not pay for this service more often than _____.”)

CASE STUDIES WITH ANALYSIS

Case Study 1

Facts: A Medicare patient is scheduled for a first diagnostic joint injection procedure described by HCPCS code 64635 (Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); lumbar or sacral, single facet joint) at a pain clinic that is an outpatient department of a PPS hospital.

The following documentation is in the pain clinic electronic record for the patient:

1. Pain clinic nursing assessment showing pain level 6/10.
2. Bilateral hip, sacroiliac (SI) joint, and lower back x-rays.
3. H&P signed by the pain clinic physician detailing the intended procedure, referencing the x-rays, and referencing the nursing pain assessment.

Turn to the *LCD – Facet Joint Interventions for Pain Management L34892*) and *Article – Billing and Coding: Facet Joint Interventions for Pain Management (A56670)* in the materials behind the outline and review the requirements for coverage of the procedure. What additional documentation is needed to demonstrate coverage?

Analysis: The pain clinic physician must document that there is no untreated radiculopathy or neurogenic claudication and no non-facet pathology that could explain the source of the patient’s pain, as required by the LCD.

In addition to the pain assessment, the LCD requires a disability scale at baseline for functional assessment. The billing and coding article lists some acceptable disability scales in the “Documentation Requirements” section.

The patient’s history of pain and conservative care that has been tried must also be documented. The LCD requires the patient has had pain for a minimum of 3 months with documented failure of conservative management, defined in the policy as use of NSAIDs, acetaminophen, physical therapy, acupuncture, or spinal manipulation.

The provider must also document that the patient has a covered diagnosis, supported by the medical record. The Article contains a list of covered diagnosis codes.

Note: This procedure will require a prior authorization when performed in a PPS hospital outpatient department beginning July 1, 2023.

Note: This case study is based on a Novitas LCD, and billing and coding Article provided for illustrative purposes only. The LCDs and billing and coding Articles for facet joint injections vary by jurisdiction, including the documentation required to

demonstrate coverage. Verify coverage, including documentation requirements, with the LCD for the applicable jurisdiction and timeframe when determining coverage.

Version 03/03/2023
Check for Updates

Case Study 2

Facts: A Medicare patient presents to the CAH for a CT scan of the head without and with contrast. Under the applicable Medicare coverage policy, the test was not considered medically necessary for the patient's condition. The patient signed an ABN before the procedure was performed. The ABN was properly prepared and the CAH provided an estimated cost of \$2,000 - \$2,200 for the intended procedure.

The CAH performed the procedure; however, the patient required a more expensive contrast than what is normally used, resulting in a total charge of \$2,400 for the procedure. The CAH billed Medicare and Medicare denied coverage. The patient was billed \$2,400 for the CT scan.

The patient called the hospital and said she is only going to pay \$850. She claims that the hospital is overcharging her because she found out from the Medicare beneficiary services hotline that Medicare typically only pays \$850 for the service.

- Assuming no state laws affect the amount to be collected by the provider, how much may the CAH collect from the patient?

Analysis: The patient is financially liable for \$2,400. The cost estimate was made in good faith because it is within the greater of \$100 or 25% of the final cost to the patient – 25% of the final cost of \$2,400 would be \$600, meaning any estimate greater than \$1,800 would be within 25% of the final cost.

Case Study 3

Facts: A Medicare patient presented to the hospital for a minor non-cosmetic surgical procedure related to varicose veins. Under an applicable LCD, the procedure is not considered medically necessary for patients with a diagnosis of varicose veins. A properly prepared ABN was reviewed with the patient.

The patient refused to sign the ABN, but still wants to proceed with the procedure based on the recommendation of his physician. Two witnesses acknowledged in writing on the ABN form that the ABN had been reviewed with the patient, but that he refused to sign it. May the hospital bill the patient for the procedure, even though they did not sign the ABN?

Analysis: Under the "LOL" provisions, an ABN does not need to be signed to be effective as long as the patient read and understood the notice. Two witnesses should document the patient's refusal to sign, which was done in this case. < *Medicare Claims Processing Manual*, Chapter 30 § 40.2.2, B and 50.6, A.2.>

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| CONTRACTOR NAME | CONTRACT TYPE | CONTRACT NUMBER | JURISDICTION | STATES | 3 - 23 |
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LCD Information

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LCD Title

FACET JOINT Interventions for Pain Management

Proposed LCD in Comment Period

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CMS National Coverage Policy

This LCD supplements but does not replace, modify or supersede existing Medicare applicable National Coverage Determinations (NCDs) or payment policy rules and regulations for FACET JOINT interventions for pain management. Federal statute and subsequent Medicare regulations regarding provision and payment for medical services are lengthy. They are not repeated in this LCD. Neither Medicare payment policy rules nor this LCD replace, modify or

Editor's note: This LCD was abbreviated for these materials. The full LCD with applicable jurisdictions and full Summary of Evidence is available in the Medicare Coverage Database.

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supersede applicable state statutes regarding medical practice or other health practice professions acts, definitions and/or scopes of practice. All providers who report services for Medicare payment must fully understand and follow all existing laws, regulations and rules for Medicare payment for FACET JOINT interventions for pain management and must properly submit only valid claims for them. Please review and understand them and apply the medical necessity provisions in the policy within the context of the manual rules. Relevant CMS manual instructions and policies may be found in the following Internet-Only Manuals (IOMs) published on the CMS Web site:

IOM Citations:

- CMS IOM Publication 100-02, *Medicare Benefit Policy Manual*,
 - Chapter 15, Section 50 Drugs and Biologicals
- CMS IOM Publication 100-03, *Medicare National Coverage Determinations (NCD) Manual*, Chapter 1,
 - Part 1, Section 30.3 for Acupuncture
 - Part 2, Section 150.7 for Prolotherapy, Joint Sclerotherapy, and Ligamentous Injections with Sclerosing Agents
 - Part 4, Section 220.1 for Computed Tomography (CT)
- CMS IOM Publication 100-08, *Medicare Program Integrity Manual*,
 - Chapter 13, Section 13.5.4 Reasonable and Necessary Provision in an LCD

Social Security Act (Title XVIII) Standard References:

- Title XVIII of the Social Security Act, Section 1861(s)(2)(K), medical or surgical services provided by a physician, certified nurse-midwife services, qualified psychologist services, and services of a certified registered nurse anesthetist;
- Title XVIII of the Social Security Act, Section 1862(a)(1)(A) states that no Medicare payment shall be made for items or services which are not reasonable and necessary for the diagnosis or treatment of illness or injury.
- Title XVIII of the Social Security Act, Section 1862(a)[14], which are other than physicians' services described by section 1861(s)(2)(K)
- Title XVIII of the Social Security Act, Section 1862(a)(7). This section excludes routine physical examinations.

Code of Federal Regulations (CFR) References:

- CFR, Title 42, Volume 2, Chapter IV, Part 410.74 Physician assistants' services.
- CFR, Title 42, Volume 2, Chapter IV, Part 410.75 Nurse practitioners' services.
- CFR, Title 42, Volume 2, Chapter IV, Part 410.76 Clinical nurse specialists' services.
- CFR, Title 42, Volume 3, Chapter IV, Part 419.22 Hospital services excluded from payment under the hospital outpatient prospective payment system.
- FR, Volume 65, Number 68, Page 18543. April 7, 2000, non-physician providers services, as defined

Coverage Guidance

Coverage Indications, Limitations, and/or Medical Necessity

Compliance with the provisions in this LCD may be monitored and addressed through post payment data analysis and subsequent medical review audits.

History/Background and/or General Information

The spine is the most common source of chronic pain. Chronic axial spinal pain is one of the major causes of

disability and accounts for a substantial U.S. health burden. Chronic spine pain poses a peculiar diagnostic and therapeutic challenge due to multiple pain sources, overlapping clinical features, and nonspecific radiological findings.

The FACET JOINTs can cause axial spinal pain and referred pain in the extremities. The pathology of the pain source is due to FACET JOINTs being richly innervated by the nerve fibers from the medial branch of the dorsal ramus of spinal nerves. Each facet has a dual nerve supply. One exception is at the C2–C3 zygapophysial joint, which has a singular nerve supply from the third occipital nerve (the superficial medial branch of C3 dorsal ramus).¹

FACET JOINT interventions may be used in pain management for chronic cervical/thoracic and back pain arising from the paravertebral FACET JOINTs. The facet block procedure is an injection of a local anesthetic, with or without a steroid medication, either into the FACET JOINT (intra-articular) or outside the joint space around the nerve supply to the joint (the medial branch nerve) known as medial branch block (MBB). Imaging guidance (fluoroscopy or CT per code descriptor) is used to assure accurate placement of the needle for the injection. Paravertebral FACET JOINT denervation is a therapeutic intervention used to provide both long-term pain relief and reduce the likelihood of recurrence of chronic cervical/thoracic or back pain confirmed as originating in the FACET JOINT's medial branch nerve.¹

There are various methods that may be used in performing FACET JOINT denervation. Percutaneous radiofrequency ablation (RFA) is a minimally invasive procedure done with imaging guidance (fluoroscopy or CT per code descriptor) and involves using energy in the radiofrequency range to cause necrosis of specific nerves (medial branches of the dorsal rami), preventing the neural transmission of pain. Conventional radiofrequency ablation (non-pulsed or continuous) applies thermal energy of typically 80 to 85 degrees Celsius. The terms RFA and radiofrequency neurotomy are used interchangeably. Both terms refer to a procedure that destroys the functionality of the nerve using radiofrequency energy. Non-thermal methods of denervation include chemical (chemodenervation), low-grade thermal energy (less than 80 degrees Celsius), pulsed RFA, laser neurolysis, and cryoablation.¹

Throughout this document, societal recommendations with the grading of evidence are referenced. There are multiple systems to grade or rank the quality of medical evidence and develop evidence-based recommendations. Not all grading systems are equivalent, so while there are typically similarities in the grades or recommendations from various grading systems, they must be considered independent of the other. The references in this document refer to the following grading systems.

1. GRADE Guidelines used in some systematic reviews, the basis for NASS recommendations align with GRADE.
2. A Modified approach to the grading of evidence² and development of interventional pain management specific instrument³ used in American Society of Interventional Pain Physicians (ASIPP) Guidelines and some systematic reviews.
3. The U.S. Preventive Services Task Force grading of evidence guidelines used by 2020 Consensus Guidelines by Cohen et al⁴.
4. Levels of Evidence for Primary Research Question and Grades of Recommendation for Summaries or Review of Studies adopted by North American Spine Society (NASS).⁵

A Multi-MAC Subject Matter Expert (SME) Panel on FACET JOINT and Medial Nerve Branch Procedures meeting was held on 5/28/2020.

Covered Indications

A. FACET JOINT Interventions:

FACET JOINT Interventions generally consist of four types of procedures: Intraarticular (IA) FACET JOINT Injections, Medial Branch Blocks (MBB), Radiofrequency Ablations (RFA) and Facet cyst rupture/aspiration.

 FACET JOINT Interventions are considered medically reasonable and necessary for the diagnosis and treatment of chronic pain in patients who meet **ALL** of the following criteria:

1. Moderate to severe chronic neck or low back pain, predominantly axial, that causes functional deficit measured on pain or disability scale¹¹; **AND***
2. Pain that has been present for a minimum of 3 months with documented failure to respond to noninvasive conservative care management (as tolerated)^{4,12}; **AND**
3. Absence of untreated radiculopathy or neurogenic claudication (except for radiculopathy caused by FACET JOINT synovial cyst)^{4,7}; **AND**
4. There is no non-facet pathology per clinical assessment or radiology studies that could explain the source of the patient's pain, including but not limited to fracture, tumor, infection, or significant deformity.¹¹

 *Pain assessment must be performed and documented at baseline, after each diagnostic procedure using the same pain scale for each assessment. A disability scale must also be obtained at baseline to be used for functional assessment (if patient qualifies for treatment).

B. Diagnostic FACET JOINT Injection Procedures (IA or MBB):

The primary indication of a diagnostic FACET JOINT procedure is to diagnose whether the patient has facet syndrome.^{1,4,7,12,16} Intraarticular (IA) facet block(s) are considered medically reasonable and necessary as a diagnostic test only if MBB cannot be performed due to specific documented anatomic restrictions or there is an indication to proceed with therapeutic intraarticular injections. These restrictions must be clearly documented in the medical record and made available upon request.

Diagnostic procedures should be performed with the intent that if successful, a RFA procedure would be considered the primary treatment goal at the diagnosed level(s).¹¹

A second diagnostic facet procedure is considered medically reasonable and necessary to confirm validity of the initial diagnostic facet procedure when administered at the same level. The second diagnostic procedure may only be performed a minimum of 2 weeks after the initial diagnostic procedure.⁷

-  1. An initial diagnostic FACET JOINT procedure will be considered medically reasonable and necessary when the patient meets the criteria outlined under the indications for FACET JOINT interventions.
2. A second confirmatory diagnostic FACET JOINT procedure is considered medically reasonable and necessary in patients who meet **BOTH** of the following criteria:
- The patient meets the criteria for the first diagnostic procedure; **AND**
 - After the first diagnostic FACET JOINT procedure, there must be a consistent positive response of at least 80% relief of primary (index) pain (with the duration of relief being consistent with the agent used).¹¹

Frequency limitation: For each covered spinal region no more than four (4) diagnostic joint sessions will be considered medically reasonable and necessary per rolling 12 months, in recognition that the pain generator cannot always be identified with the initial and confirmatory diagnostic procedure.

C. Therapeutic FACET JOINT Injection Procedures (IA):

Therapeutic FACET JOINT procedures are considered medically reasonable and necessary for patients who meet **ALL** of the following criteria:

1. The patient has had two (2) medically reasonable and necessary diagnostic FACET JOINT procedures with each one providing a consistent minimum of 80% relief of primary (index) pain (with the duration of relief being consistent with the agent used); **AND**
2. Subsequent therapeutic FACET JOINT procedures at the same anatomic site result in at least consistent 50% pain relief for at least three (3) months from the prior therapeutic procedure or at least 50% consistent improvement in the ability to perform previously painful movements and ADLs as compared to baseline measurement using the same scale¹¹; **AND**
3. Documentation of why the patient is not a candidate for radiofrequency ablation (such as established spinal pseudarthrosis, implanted electrical device) is in the medical record.^{5,42,43,44}

Frequency limitation: For each covered spinal region no more than four (4) therapeutic FACET JOINT injection (IA) sessions will be reimbursed per rolling 12 months.

D. FACET JOINT Denervation:

An initial thermal RFA of cervical, thoracic, or lumbar paravertebral FACET JOINT (medial branch) nerves is considered medically reasonable and necessary for patients who have had at least two (2) medically reasonable and necessary diagnostic MBBs, with each one providing a consistent minimum of 80% sustained relief of primary (index) pain (with the duration of relief being consistent with the agent used).

Repeat thermal¹¹ FACET JOINT RFA at the same anatomic site is considered medically reasonable and necessary provided the patient had a minimum of consistent 50% improvement in pain for at least six (6) months **or** at least 50% consistent improvement in the ability to perform previously painful movements and ADLs as compared to baseline measurement using the same scale.

Frequency limitation: For each covered spinal region no more than two (2) radiofrequency sessions will be reimbursed per rolling 12 months.

E. Facet Cyst Aspiration/Rupture

Intra-articular FACET JOINT injection performed with synovial cyst aspiration is considered medically reasonable and necessary when **BOTH** of the following criteria are met:

1. Advanced diagnostic imaging study (e.g., MRI/CT/myelogram) confirm compression or displacement of the corresponding nerve root by a FACET JOINT synovial cyst; **AND**
2. Clinical and physical symptoms related to synovial facet cyst are documented in the medical record.

Frequency limitation: Cyst aspiration/rupture may be repeated once and only if there is 50% or more consistent improvement in pain for at least three (3) months.¹¹

Limitations

1. FACET JOINT interventions done without CT or fluoroscopic guidance are considered not medically reasonable and necessary. This includes FACET JOINT interventions done without any guidance,

- performed under ultrasound guidance,^{4,11} or with Magnetic Resonance Imaging (MRI).⁴
2. General anesthesia is considered not medically reasonable and necessary for FACET JOINT interventions. Neither conscious sedation nor Monitored Anesthesia Care (MAC) is routinely necessary for intraarticular FACET JOINT injections or medial branch blocks and are not routinely considered medically reasonable and necessary. Individual consideration may be given on redetermination (appeal) for payment in rare, unique circumstances if the medical necessity of sedation is unequivocal and clearly documented in the medical record. Frequent reporting of these services together may trigger focused medical review.⁴
 3. It is not expected that patients will present with pain in both cervical/thoracic and lumbar spinal regions. Therefore, FACET JOINT interventions (both diagnostic and therapeutic) are limited to one spinal region per session.
 4. It is not routinely necessary for multiple blocks (e.g., epidural injections, sympathetic blocks, trigger point injections, etc.) to be provided to a patient on the same day as FACET JOINT procedures. Multiple blocks on the same day could lead to improper or lack of diagnosis. If performed, the medical necessity of each injection (at the same or a different level[s]) must be clearly documented in the medical record. For example, the performance of both paravertebral FACET JOINT procedures and a transforaminal epidural steroid injection (TFESI) at the same or close spinal level at the same encounter would not be expected unless a synovial cyst is compressing the nerve root. In this situation, TFESI may provide relief for the radicular pain, while the facet cyst rupture allows nerve root decompression. Frequent reporting of multiple blocks on the same day may trigger a focused medical review.
 5. FACET JOINT intraarticular injections and medial branch blocks may involve the use of anesthetic, corticosteroids, anti-inflammatories and/or contrast agents and do not include injections of biologicals or other substances not FDA designated for this use.
 6. One to two levels, either unilateral or bilateral, are allowed per session per spine region. The need for a three or four-level procedure bilaterally may be considered under unique circumstances and with sufficient documentation of medical necessity on appeal. A session is a time period, which includes all procedures (i.e., MBB, IA, facet cyst ruptures, and RFA ablations) that are performed during the same day.
 7. If there is an extended period of time, two years or more, since the last RFA and/or there is a question as to the source of the recurrent pain then diagnostic procedures must be repeated.
 8. Therapeutic intraarticular facet injections are not considered medically reasonable and necessary unless there is documentation explaining why RFA cannot be performed.^{5,42,43,44}
 9. FACET JOINT procedures in patients for the indication of generalized pain conditions (such as fibromyalgia) or chronic centralized pain syndromes are considered not medically reasonable and necessary. Individual consideration may be considered under unique circumstances and with sufficient documentation of medical necessity on appeal.⁵⁶
 10. In patients with implanted electrical devices, providers must follow manufacturer instructions and extra planning as indicated to ensure safety of procedure.

The following are considered not medically reasonable and necessary:

1. Intraarticular and extraarticular FACET JOINT prolotherapy^{5,42,43,44}
2. Non-thermal modalities for FACET JOINT denervation including chemical, low-grade thermal energy (less than 80 degrees Celsius), laser neurolysis, and cryoablation
3. Intra-facet implants⁵⁸
4. FACET JOINT procedure performed after anterior lumbar interbody fusion (ALIF)
5. Definitive clinical and/or imaging findings pointing to a specific diagnosis other than FACET JOINT syndrome
6. Diagnostic injections or MBB at the same level as the previously successful RFA procedure

Notice: Services performed for any given diagnosis must meet all the indications and limitations stated in this LCD, the general requirements for medical necessity as stated in CMS payment policy manuals, any and all existing CMS

national coverage determinations, and all Medicare payment rules.



Provider Qualifications

Patient safety and quality of care mandate that healthcare professionals who perform facet injections/procedures are appropriately trained and/or credentialed by a formal residency/fellowship program and/or are certified by either an accredited and nationally recognized organization or by a post-graduate training course accredited by an established national accrediting body or accredited professional training program. If the practitioner works in a hospital facility at any time and/or is credentialed by a hospital for any procedure, the practitioner must be credentialed to perform the same procedure in the outpatient setting. At a minimum, training must cover and develop an understanding of anatomy and drug pharmacodynamics and kinetics as well as proficiency in diagnosis and management of disease, the technical performance of the procedure, and utilization of the required associated imaging modalities.

In addition to the above requirements, non-physician providers, such as certified nurse anesthetist, with certain exceptions, may certify, order and establish the plan of care as authorized by State law. (See Sections 1861[s][2] and 1862[a][14] of Title XVIII of the Social Security Act; 42 CFR, Sections 410.74, 410.75, 410.76 and 419.22; FR Vol. 65, No. 68 page18543, April 7, 2000). Each practitioner must provide only those services within the scope of practice for each state.

Definitions

Acute Pain: The temporal definition of pain persisting for up to 4 weeks after the onset of the pain.

Axial: Relating to or situated in the central part of the body, in the head and trunk as distinguished from the limbs, e.g., axial skeleton.

Biopsychosocial Model: Interdisciplinary model that looks at the interconnection between biology, pathology and socioenvironmental factors.

Central Neuropathic Pain: Pain, which is causally related to a lesion or disease of the central somatosensory nerves.

Centralized Pain: A neurological chronic pain syndrome of the central nervous system (brain, brainstem, and spinal cord) which commonly presents with widespread generalized allodynia which is causally related to the increased responsiveness of nociceptive nerves in the central nervous system to the normal threshold or subthreshold stimulation from the afferent nerves. The condition has also been called "central sensitization," "central amplification," and "central pain syndrome." Fibromyalgia is considered one of the most common centralized pain syndromes.

Cervical Facet Pain: Pain located in the cervical spine, which may be characterized by chronic headaches, restricted motion, and axial neck pain, which may radiate sub-occipitally to the shoulders or mid-back.

Chronic Pain: The temporal definition of pain persisting for greater than or equal to 12 weeks after the onset of the pain.

Dual Diagnostic Blocks: The diagnostic technique of injecting the same spinal nerve on two separate occasions to be used as an efficacy comparison to increase diagnostic accuracy.

Epidural Steroid Injection: The administration via injection of steroid medicine into the potential epidural space in the spinal column to deliver steroids to the spinal nerves.

FACET JOINT Intraarticular Injections, Diagnostic: The placement of local anesthetic and possibly a corticosteroid into the FACET JOINT to diagnose FACET JOINT pain.

FACET JOINT Intraarticular Injections, Therapeutic: The placement of local anesthetic and possibly a corticosteroid into the FACET JOINT to produce the beneficial effect of pain reduction.

FACET JOINT: A diarthrodial joint in the spinal column (also called the zygapophysial joint or z-joint), producing the articulation of the posterior elements of one vertebra with its neighboring vertebra. There are bilateral superior and inferior articular surfaces at each spinal level. The terminology or nomenclature of the FACET JOINT is classified by the specific vertebrae level that forms it (e.g., C4-5 or L2-3). There are two (2) FACET JOINTs, right and left, at each spinal level.

Facet Injection: (also called facet block) A general term used to describe the injection of local anesthetic and possibly a corticosteroid in the FACET JOINT capsule or along the medial branch nerves supplying the FACET JOINTs.

FACET JOINT Denervation or Radiofrequency Ablation (RFA): A general term used to describe the minimally invasive procedure that uses thermal energy generated by the radiofrequency current to deprive the FACET JOINT of its nerve supply. The procedure is also known as a Medial Branch Radiofrequency Neurotomy (Ablation) because it is used to thermally remove the medial branch nerve by using electrical current to create thermal energy to coagulate the adjacent tissues around the targeted medial branch nerve.

FACET JOINT Syndrome: A set of concurrent signs or symptoms to describe FACET JOINT pain as the pain generator. The typical clinical signs or symptoms of a facet syndrome may include local paraspinal tenderness; pain that is brought about or increased on hyperextension, rotation, and lateral bending; low back stiffness; absence of neurologic deficit; absence of root tension signs (non-radiating below the knee, absence of paresthesia). Cervical facet pain is often characterized by chronic headaches, restricted motion, and axial neck pain, which may radiate sub-occipitally to the shoulders or mid-back.

Facet Level: Refers to the zygapophyseal joint or the two medial branch (MB) nerves that innervate that zygapophyseal joint. Each level has a pair of FACET JOINTs: one on the right side and one of the left side of the spine.

Intra-Articular Injection (IA): The injection of local anesthetic and possibly a corticosteroid into the FACET JOINT capsule.

Medial Branch: The dorsal ramus is the dorsal branch of a spinal nerve that forms from the dorsal root of the nerve after it emerges from the spinal cord.

Medial Branch Block (MBB): The placement of local anesthetic and possibly a corticosteroid near the medial branch nerve which supplies the sensory innervation to a specific FACET JOINT.

Neuropathic Pain: The pain which is caused by a lesion or disease of the somatosensory nerves.

Neurogenic Claudication: Intermittent leg pain from impingement of the nerves emanating from the spinal cord

(also called pseudoocclusion).

New Onset of Spinal Pain: The new onset of the spinal pain must be materially and significantly different in location, type, duration and character from the previously treated spine pain.

Noninvasive Conservative Management: The use of nonsteroidal anti-inflammatory drugs (NSAIDs), acetaminophen, physical therapy, acupuncture (applies to only chronic low back pain), or spinal manipulation. This management should include the application of a biopsychosocial treatment technique.

Non-Radicular Back Pain: The radiating non-neuropathic pain which is not causally related to a spinal nerve root irritation and does not produce reproducible neuropathic symptoms in an objective dermatomal pattern.

Peripheral Neuropathic Pain: Pain, which is causally related to a lesion or disease of the peripheral somatosensory nerves.

Radicular Back Pain: The radiating neuropathic pain causally related to the spinal nerve root irritation which extends into the distal distribution, typically the lower extremity, producing neuropathic pain in a dermatomal pattern.

Radiculopathy: Radiating neuropathic pain causally related to the spinal nerve root irritation, which extends distal producing neuropathic pain in a dermatomal pattern.

Region: The segments of the back involved will be defined in this policy as two regions:

1. Cervical/Thoracic region = C1-C7/T1-T12
2. Lumbar/Sacral region = L1-L5/S1-S5

Session: A time period, which includes all procedures (i.e., MBB, IA, facet cyst ruptures, and RFA ablations) performed during one day.

Subacute Pain: The temporal definition of pain occurring during the 4-12-week time period.

Transforaminal Epidural Steroid Injection (TFESI): An epidural injection performed via a paramedian approach to enter the epidural space by placing the needle in the posterior-superior quadrant of the intervertebral foramen (neuroforamen) to inject near the dorsal root ganglion and exiting spinal nerve root (previously known as a selective nerve root block).

Summary of Evidence

Diagnostic FACET JOINT Injections

Due to the lack of reliable history, physical exam, or imaging to predict response, providers must rely on facet interventions diagnostic injections given for diagnostic purposes to determine if the FACET JOINT is the source of suspected spinal pain. There is controversy over optimal patient selection for diagnostic injections, which measures successful response and type and number of diagnostic injections performed.

Numerous investigations have been undertaken to correlate symptoms and physical exam findings with facet pathology and have concluded conventional clinical findings are unreliable in identifying FACET JOINT success.

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Article Information

General Information

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Article Title

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Article Type

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CMS National Coverage Policy

Internet-Only Manuals (IOMs)

- CMS IOM Publication 100-04, *Medicare Claims Processing Manual*,

- Chapter 12, Section 40.7 Claims for Bilateral Surgeries
- Chapter 13, Section 10.1 Billing Part B Radiology Services and Other Diagnostic Procedures, Section 20 Payment Conditions for Radiology Services, and Section 30 Computerized Axial Tomography (CT) Procedures

Social Security Act (Title XVIII) Standard References:

- Title XVIII of the Social Security Act, Section 1833(e) states that no payment shall be made to any provider of services or other person under this part unless there has been furnished such information as may be necessary in order to determine the amounts due such provider or other person under this part for the period with respect to which the amounts are being paid or for any prior period.

Article Guidance

Article Text

This Billing and Coding Article provides billing and coding guidance for Local Coverage Determination (LCD) L34892, FACET JOINT Interventions for Pain Management. Please refer to the LCD for reasonable and necessary requirements.

Coding Guidance

Notice: It is not appropriate to bill Medicare for services that are not covered (as described by the entire LCD) as if they are covered. When billing for non-covered services, use the appropriate modifier.

Diagnostic and Therapeutic Procedures:

Each paravertebral facet level refers to either the FACET JOINT, also called the zygapophyseal joint **OR** the two medial branch nerves that innervate each zygapophyseal joint.

Each facet level has a pair of FACET JOINTS (one on the right side and one on the left side of the spine). Unilateral or bilateral facet interventions may be performed during the FACET JOINT procedure (a diagnostic nerve block, a therapeutic FACET JOINT [intraarticular] injection, or a medial branch block injection, in one session. A bilateral facet nerve intervention is still considered a single level intervention.

For paravertebral spinal nerves and branch injections, image guidance (fluoroscopy or CT) is required for the performance of CPT codes 64490, 64491, 64493, and 64494 with any injection contrast, which is an included component of the code.

As defined by the Current Procedural Terminology (CPT) Professional edition code book, there are two distinct anatomic spinal regions for paravertebral facet injections: cervical /thoracic (codes 64490, 64491) and lumbar/sacral (codes 64493, 64494).

For each initial, single level injection, diagnostic or therapeutic, performed with image guidance (fluoroscopy or CT), use code 64490 (cervical or thoracic) or code 64493 (lumbar or sacral).

For any additional diagnostic or therapeutic procedures on the same day, use add-on codes 64491 (cervical/thoracic) or 64494 (lumbar/sacral) to report second level injections performed with image guidance (fluoroscopy or CT) in addition to the primary procedure codes 64490 or 64493.

Note: Each unilateral or bilateral intervention at any level should be reported as one unit of service (UOS).

If an initial (64490 or 64493) or second level add-on (64491 or 64494) paravertebral facet injection procedure is performed bilaterally, report the procedure with modifier -50 as a single line item using one UOS. Do not use modifier RT or LT when performing these procedures bilaterally (modifier -50).

For services performed in the Ambulatory Surgical Center (ASC), do not use modifier 50. Report the applicable procedure code on two separate lines, with one unit each and append the -RT and -LT modifiers to each line.

When an intraarticular FACET JOINT injection is used for facet cyst aspiration/rupture, it should be reported with CPT code 64999. Providers are required to indicate in block 19 of the 1500 claim form or the EMC Equivalent the date of the initial injection procedure and if the injection procedure is being repeated.

For CPT codes 64492 and 64495, the need for a three-level procedure may be considered under unique circumstances and with sufficient documentation of medical necessity on appeal.

KX modifier requirements:

The KX modifier should be appended to the line for all diagnostic injections. In most cases the KX modifier will only be used for the two initial diagnostic injections. If the initial diagnostic injections do not produce a positive response as defined by the LCD and are not indicative of identification of the pain generator, and it is necessary to perform additional diagnostic injections, at a different level, append the KX modifier to the line. Aberrant use of the KX modifier may trigger focused medical review.

Neurolytic Destruction Procedures (Radiofrequency Ablation):

For destruction of paravertebral FACET JOINT medial branch nerves, image guidance and localization (fluoroscopy or CT) are required and inclusive in codes 64633, 64634, 64635, and 64636.

Per the current CPT Professional edition code book, codes 64633, 64634, 64635, and 64636 are reported per joint, not per nerve. Although two nerves innervate each FACET JOINT, only one unit per code may be reported for each joint denervated, regardless of the number of nerves treated. There are two distinct anatomic spinal regions for paravertebral facet destruction: cervical/thoracic (codes 64633, 64634) and lumbar/sacral (codes 64635, 64636). For each initial, single level thermal radiofrequency destruction performed with image guidance (fluoroscopy or CT), use code 64633 (cervical or thoracic) or code 64635 (lumbar or sacral).

For any additional thermal radiofrequency destruction performed on the same day, use add-on codes 64634 (cervical/thoracic) or 64636 (lumbar/sacral) in addition to the primary procedure codes 64633 or 64635.

Note: Each unilateral or bilateral intervention at any level should be reported as one UOS.

If initial (64633 or 64635) or each additional add-on (64634 or 64636) paravertebral neurolytic destruction procedure is performed bilaterally, report the procedure with modifier -50 as a single line item using one UOS. Do not use modifier RT or LT when performing these services bilaterally (modifier -50).

For services performed in the Ambulatory Surgical Center (ASC), do not use modifier 50. Report the applicable procedure code on two separate lines, with one unit each and append the -RT and -LT modifiers to each line.

Non-thermal FACET JOINT denervation (including chemical, low grade thermal energy [<80 degrees Celsius] or any other form of pulsed radiofrequency) should not be reported with CPT codes 64633, 64634, 64635 or 64636. These services should be reported with CPT code 64999.

Note: Report CPT code 64999 when facet cyst aspiration/rupture is performed.

Note: CPT code 64999 is non covered when used to report non thermal FACET JOINT denervation including chemical, low grade thermal energy (less than 80 degrees Celsius) or any form of pulsed radiofrequency.

Note: When reporting CPT code 64999 ensure that the description of the service is included on the claim.

If FACET JOINTs are injected with biologicals or other substances not designated for this use the entire claim may deny per CMS IOM *Medicare Benefit Policy Manual*, Chapter 16, Section 180-Services Related to and Required as a Result of Services Which Are Not Covered Under Medicare.

Utilization Parameters

Note: A session is defined as all procedures (i.e., MBB, IA, facet cyst ruptures, and destruction by a neurolytic agent (e.g., RFA) performed on the same date of service.

CPT codes 64490 through 64494 will be limited to no more than four (4) sessions, per region, per rolling 12 months.

CPT code 64490 through 64494 with the KX modifier will be limited to no more than four (4) sessions, per region, per rolling 12 months.



CPT codes 64633 through 64636 will be limited to no more than two (2) sessions, per region, per rolling 12 months.

Consistent with the LCD, CPT code 64999 may only be reported twice for an intraarticular FACET JOINT injection for a facet cyst aspiration/rupture.

Documentation Requirements

1. All documentation must be maintained in the patient's medical record and made available to the contractor upon request.
2. Every page of the record must be legible and include appropriate patient identification information (e.g., complete name, dates of service[s]). The documentation must include the legible signature of the physician or non-physician practitioner responsible for and providing the care to the patient.
3. The submitted medical record must support the use of the selected ICD-10-CM code(s). The submitted CPT/HCPCS code must describe the service performed.
4. The medical record must include the assessment of the patient by the performing provider as it relates to the complaint of the patient for that visit, relevant medical history, and the results of any pertinent tests/procedures.
5. Documentation of why the patient is not a candidate for radiofrequency ablation (RFA) must be submitted for therapeutic injection procedures.
6. The scales used to assess the measurement of pain and/or disability must be documented in the medical record. Acceptable scales include but are not limited to: verbal rating scales, Numerical Rating Scale (NRS), Visual Analog Scale (VAS) for pain assessment, Pain Disability Assessment Scale (PDAS), Oswestry Disability Index (ODI), Oswestry Low Back Pain Disability Questionnaire (OLBPDQ), Quebec Back Pain Disability Score (QBPDS), Roland Morris Pain Scale, Back Pain Functional Scale (BPFS), and the Patient-Reported Outcomes Measurement Information System (PROMIS) profile domains to assess function.

Coding Information

CPT/HCPCS Codes**Group 1 Paragraph:**

Providers are reminded to refer to the long descriptors of the CPT codes in their CPT book.

Note: Report CPT code **64999** when facet cyst aspiration/rupture is performed.

The following CPT codes need to be listed separately in addition to a code for the primary procedure: **64491, 64494, 64634, and 64636.**

Group 1 Codes: (9 Codes)

| CODE | DESCRIPTION |
|-------|------------------------------|
| 64490 | Inj paravert f jnt c/t 1 lev |
| 64491 | Inj paravert f jnt c/t 2 lev |
| 64493 | Inj paravert f jnt l/s 1 lev |
| 64494 | Inj paravert f jnt l/s 2 lev |
| 64633 | Destroy cerv/thor facet jnt |
| 64634 | Destroy c/th facet jnt addl |
| 64635 | Destroy lumb/sac facet jnt |
| 64636 | Destroy l/s facet jnt addl |
| 64999 | Nervous system surgery |

Group 2 Paragraph:

Note: CPT code **64999** is non-covered when used to report non-thermal facet joint denervation including chemical, low grade thermal energy (less than 80 degrees Celsius), or any form of pulsed radiofrequency.

The following CPT/HCPCS codes do not support medical necessity and will not be covered by Medicare.

Group 2 Codes: (13 Codes)

| CODE | DESCRIPTION |
|-------|------------------------------|
| 64492 | Inj paravert f jnt c/t 3 lev |
| 64495 | Inj paravert f jnt l/s 3 lev |
| 64999 | Nervous system surgery |
| 0213T | Njx paravert w/us cer/thor |
| 0214T | Njx paravert w/us cer/thor |
| 0215T | Njx paravert w/us cer/thor |
| 0216T | Njx paravert w/us lumb/sac |
| 0217T | Njx paravert w/us lumb/sac |

| CODE | DESCRIPTION |
|-------|----------------------------|
| 0218T | Njx paravert w/us lumb/sac |
| 0219T | Plmt post facet implt cerv |
| 0220T | Plmt post facet implt thor |
| 0221T | Plmt post facet implt lumb |
| 0222T | Plmt post facet implt addl |

CPT/HCPCS Modifiers**Group 1 Paragraph:**

N/A

Group 1 Codes: (4 Codes)

| CODE | DESCRIPTION |
|------|---|
| 50 | BILATERAL PROCEDURE: UNLESS OTHERWISE IDENTIFIED IN THE LISTINGS, BILATERAL PROCEDURES THAT ARE PERFORMED AT THE SAME OPERATIVE SESSION SHOULD BE IDENTIFIED BY ADDING THE MODIFIER -50 TO THE APPROPRIATE FIVE DIGIT CODE OR BY USE OF THE SEPARATE FIVE DIGIT MODIFIER CODE 09950 |
| KX | REQUIREMENTS SPECIFIED IN THE MEDICAL POLICY HAVE BEEN MET |
| LT | LEFT SIDE (USED TO IDENTIFY PROCEDURES PERFORMED ON THE LEFT SIDE OF THE BODY) |
| RT | RIGHT SIDE (USED TO IDENTIFY PROCEDURES PERFORMED ON THE RIGHT SIDE OF THE BODY) |

ICD-10-CM Codes that Support Medical Necessity**Group 1 Paragraph:**

It is the provider's responsibility to select codes carried out to the highest level of specificity and selected from the ICD-10-CM code book appropriate to the year in which the service is rendered for the claim(s) submitted.

The following ICD-10-CM codes support medical necessity and provide coverage for CPT codes: **64490, 64491, 64493, 64494, 64633, 64634, 64635, 64636, and 64999** (facet cyst aspiration/rupture).

Note: ICD-10 Codes **M71.30** or **M71.38** is allowed for facet cyst rupture procedures only.

Group 1 Codes: (20 Codes)

| CODE | DESCRIPTION |
|---------|--|
| M47.812 | Spondylosis without myelopathy or radiculopathy, cervical region |

| CODE | DESCRIPTION |
|---------|---|
| M47.813 | Spondylosis without myelopathy or radiculopathy, cervicothoracic region |
| M47.814 | Spondylosis without myelopathy or radiculopathy, thoracic region |
| M47.815 | Spondylosis without myelopathy or radiculopathy, thoracolumbar region |
| M47.816 | Spondylosis without myelopathy or radiculopathy, lumbar region |
| M47.817 | Spondylosis without myelopathy or radiculopathy, lumbosacral region |
| M47.892 | Other spondylosis, cervical region |
| M47.893 | Other spondylosis, cervicothoracic region |
| M47.894 | Other spondylosis, thoracic region |
| M47.895 | Other spondylosis, thoracolumbar region |
| M47.896 | Other spondylosis, lumbar region |
| M47.897 | Other spondylosis, lumbosacral region |
| M48.12 | Ankylosing hyperostosis [Forestier], cervical region |
| M48.13 | Ankylosing hyperostosis [Forestier], cervicothoracic region |
| M48.14 | Ankylosing hyperostosis [Forestier], thoracic region |
| M48.15 | Ankylosing hyperostosis [Forestier], thoracolumbar region |
| M48.16 | Ankylosing hyperostosis [Forestier], lumbar region |
| M48.17 | Ankylosing hyperostosis [Forestier], lumbosacral region |
| M71.30 | Other bursal cyst, unspecified site |
| M71.38 | Other bursal cyst, other site |

ICD-10-CM Codes that DO NOT Support Medical Necessity

Group 1 Paragraph:

All those not listed under the "ICD-10 Codes that Support Medical Necessity" section of this article.

Group 1 Codes: (1 Code)

| CODE | DESCRIPTION |
|-------|----------------|
| XX000 | Not Applicable |

ICD-10-PCS Codes

N/A

Additional ICD-10 Information

Bill Type Codes

Contractors may specify Bill Types to help providers identify those Bill Types typically used to report this service. Absence of a Bill Type does not guarantee that the article does not apply to that Bill Type. Complete absence of all Bill Types indicates that coverage is not influenced by Bill Type and the article should be assumed to apply equally to all claims.

| CODE | DESCRIPTION |
|------|----------------|
| 999x | Not Applicable |

Revenue Codes

Contractors may specify Revenue Codes to help providers identify those Revenue Codes typically used to report this service. In most instances Revenue Codes are purely advisory. Unless specified in the article, services reported under other Revenue Codes are equally subject to this coverage determination. Complete absence of all Revenue Codes indicates that coverage is not influenced by Revenue Code and the article should be assumed to apply equally to all Revenue Codes.

| CODE | DESCRIPTION |
|-------|----------------|
| 99999 | Not Applicable |

Other Coding Information

N/A

Revision History Information

| REVISION HISTORY DATE | REVISION HISTORY NUMBER | REVISION HISTORY EXPLANATION |
|-----------------------|-------------------------|---|
| 04/25/2021 | R4 | Article revised and published on 05/27/2021 effective for dates of service on and after 04/25/2021. The "Diagnostic and Therapeutic Procedures:" and Neurolytic Destruction Procedures (Radiofrequency Ablation):" sections of the article were revised to clarify coding guidance for the add-on CPT codes 64491/64494 (second level) and 64634/64636 (each additional) when billing bilaterally. Also, the "Internet-Only Manuals" section of the article was updated to include Pub. 100-04, <i>Medicare Claims Processing Manual</i> , Chapter 12, Section 40.7 Claims for Bilateral Surgeries. |
| 04/25/2021 | R3 | Article revised and published on 04/22/2021 effective for dates of service on and after 04/25/2021. This revision was to add clarifying language to the paragraph under the "KX modifier requirements:" section of the Article. Also, minor formatting change made |



**Medicare National Coverage
Determinations (NCD)
Coding Policy Manual and
Change Report (ICD-10-CM)
*January 2023**



Clinical Diagnostic Laboratory Services

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Centers for Medicare & Medicaid Services**

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NCD Manual Changes

| CR Date | Reason | Release | Change | Edit |
|-----------|---|----------|--------|---|
| *01/01/23 | *Per CR 12888 add the specified ICD-10-CM codes from the list of ICD-10-CM codes that are denied for the Urine Culture, Bacterial (190.12) NCD. *Transmittal #11583 | *2023100 | | *190.12 Urine Culture, Bacterial |
| *01/01/23 | *Per CR 12888 add the specified ICD-10-CM codes from the list of ICD-10-CM codes that are covered for the Urine Culture, Bacterial (190.12) NCD. *Transmittal #11583 | *2023100 | | *190.12 Urine Culture, Bacterial |
| *01/01/23 | *Per CR 12888 add the specified ICD-10-CM codes from the list of ICD-10-CM codes that are denied for the Human Immunodeficiency Virus (HIV) Testing (Prognosis Including Monitoring) (190.13) NCD. *Transmittal #11583 | *2023100 | | *190.13 Human Immunodeficiency Virus (HIV) Testing (Prognosis Including Monitoring) |

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Non-covered ICD-10-CM Codes for All Lab NCDs

This section lists codes that are never covered by Medicare for a diagnostic lab testing service. If a code from this section is given as the reason for the test, the test may be billed to the Medicare beneficiary without billing Medicare first because the service is not covered by statute, in most instances because it is performed for screening purposes and is not within an exception. The beneficiary, however, does have a right to have the claim submitted to Medicare, upon request.

The ICD-10-CM codes in the table below can be viewed on CMS' website as part of
Downloads: Lab Code List, at
<http://www.cms.gov/Medicare/Coverage/CoverageGenInfo/LabNCDsICD10.html>

| Code | Description |
|---------|---|
| R99 | Ill-defined and unknown cause of mortality |
| Z00.00 | Encounter for general adult medical examination without abnormal findings |
| Z00.01 | Encounter for general adult medical examination with abnormal findings |
| Z00.110 | Health examination for newborn under 8 days old |
| Z00.111 | Health examination for newborn 8 to 28 days old |
| Z00.121 | Encounter for routine child health examination with abnormal findings |
| Z00.129 | Encounter for routine child health examination without abnormal findings |
| Z00.5 | Encounter for examination of potential donor of organ and tissue |
| Z00.6 | Encounter for examination for normal comparison and control in clinical research program |
| Z00.70 | Encounter for examination for period of delayed growth in childhood without abnormal findings |
| Z00.71 | Encounter for examination for period of delayed growth in childhood with abnormal findings |
| Z00.8 | Encounter for other general examination |
| Z02.0 | Encounter for examination for admission to educational institution |
| Z02.1 | Encounter for pre-employment examination |
| Z02.2 | Encounter for examination for admission to residential institution |
| Z02.3 | Encounter for examination for recruitment to armed forces |
| Z02.4 | Encounter for examination for driving license |
| Z02.5 | Encounter for examination for participation in sport |
| Z02.6 | Encounter for examination for insurance purposes |
| Z02.71 | Encounter for disability determination |
| Z02.79 | Encounter for issue of other medical certificate |
| Z02.81 | Encounter for paternity testing |
| Z02.82 | Encounter for adoption services |

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| Code | Description |
|--------|--|
| Z02.83 | Encounter for blood-alcohol and blood-drug test |
| Z02.89 | Encounter for other administrative examinations |
| Z02.9 | Encounter for administrative examinations, unspecified |
| Z04.6 | Encounter for general psychiatric examination, requested by authority |
| Z04.81 | Encounter for examination and observation of victim following forced sexual exploitation |
| Z04.82 | Encounter for examination and observation of victim following forced labor exploitation |
| Z04.89 | Encounter for examination and observation for other specified reasons |
| Z04.9 | Encounter for examination and observation for unspecified reason |
| Z11.0 | Encounter for screening for intestinal infectious diseases |
| Z11.1 | Encounter for screening for respiratory tuberculosis |
| Z11.2 | Encounter for screening for other bacterial diseases |
| Z11.3 | Encounter for screening for infections with a predominantly sexual mode of transmission |
| Z11.4 | Encounter for screening for human immunodeficiency virus [HIV] |
| Z11.51 | Encounter for screening for human papillomavirus (HPV) |
| Z11.52 | Encounter for screening for COVID-19 |
| Z11.59 | Encounter for screening for other viral diseases |
| Z11.6 | Encounter for screening for other protozoal diseases and helminthiases |
| Z11.7 | Encounter for testing for latent tuberculosis infection |
| Z11.8 | Encounter for screening for other infectious and parasitic diseases |
| Z11.9 | Encounter for screening for infectious and parasitic diseases, unspecified |
| Z12.0 | Encounter for screening for malignant neoplasm of stomach |
| Z12.10 | Encounter for screening for malignant neoplasm of intestinal tract, unspecified |
| Z12.13 | Encounter for screening for malignant neoplasm of small intestine |
| Z12.2 | Encounter for screening for malignant neoplasm of respiratory organs |
| Z12.6 | Encounter for screening for malignant neoplasm of bladder |
| Z12.71 | Encounter for screening for malignant neoplasm of testis |
| Z12.72 | Encounter for screening for malignant neoplasm of vagina |
| Z12.73 | Encounter for screening for malignant neoplasm of ovary |
| Z12.79 | Encounter for screening for malignant neoplasm of other genitourinary organs |
| Z12.81 | Encounter for screening for malignant neoplasm of oral cavity |
| Z12.82 | Encounter for screening for malignant neoplasm of nervous system |
| Z12.83 | Encounter for screening for malignant neoplasm of skin |
| Z12.89 | Encounter for screening for malignant neoplasm of other sites |
| Z12.9 | Encounter for screening for malignant neoplasm, site unspecified |

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| Code | Description |
|---------|---|
| Z13.0 | Encounter for screening for diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism |
| Z13.21 | Encounter for screening for nutritional disorder |
| Z13.220 | Encounter for screening for lipoid disorders |
| Z13.228 | Encounter for screening for other metabolic disorders |
| Z13.29 | Encounter for screening for other suspected endocrine disorder |
| Z13.30 | Encounter for screening examination for mental health and behavioral disorders, unspecified |
| Z13.31 | Encounter for screening for depression |
| Z13.32 | Encounter for screening for maternal depression |
| Z13.39 | Encounter for screening examination for other mental health and behavioral disorders |
| Z13.40 | Encounter for screening for unspecified developmental delays |
| Z13.41 | Encounter for autism screening |
| Z13.42 | Encounter for screening for global developmental delays (milestones) |
| Z13.49 | Encounter for screening for other developmental delays |
| Z13.5 | Encounter for screening for eye and ear disorders |
| Z13.71 | Encounter for nonprocreative screening for genetic disease carrier status |
| Z13.79 | Encounter for other screening for genetic and chromosomal anomalies |
| Z13.810 | Encounter for screening for upper gastrointestinal disorder |
| Z13.811 | Encounter for screening for lower gastrointestinal disorder |
| Z13.818 | Encounter for screening for other digestive system disorders |
| Z13.820 | Encounter for screening for osteoporosis |
| Z13.828 | Encounter for screening for other musculoskeletal disorder |
| Z13.83 | Encounter for screening for respiratory disorder NEC |
| Z13.84 | Encounter for screening for dental disorders |
| Z13.850 | Encounter for screening for traumatic brain injury |
| Z13.858 | Encounter for screening for other nervous system disorders |
| Z13.88 | Encounter for screening for disorder due to exposure to contaminants |
| Z13.89 | Encounter for screening for other disorder |
| Z13.9 | Encounter for screening, unspecified |
| Z36.0 | Encounter for antenatal screening for chromosomal anomalies |
| Z36.1 | Encounter for antenatal screening for raised alphafetoprotein level |
| Z36.2 | Encounter for other antenatal screening follow-up |
| Z36.3 | Encounter for antenatal screening for malformations |

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| Code | Description |
|--------|---|
| Z36.4 | Encounter for antenatal screening for fetal growth retardation |
| Z36.5 | Encounter for antenatal screening for isoimmunization |
| Z36.81 | Encounter for antenatal screening for hydrops fetalis |
| Z36.82 | Encounter for antenatal screening for nuchal translucency |
| Z36.83 | Encounter for fetal screening for congenital cardiac abnormalities |
| Z36.84 | Encounter for antenatal screening for fetal lung maturity |
| Z36.85 | Encounter for antenatal screening for Streptococcus B |
| Z36.86 | Encounter for antenatal screening for cervical length |
| Z36.87 | Encounter for antenatal screening for uncertain dates |
| Z36.88 | Encounter for antenatal screening for fetal macrosomia |
| Z36.89 | Encounter for other specified antenatal screening |
| Z36.8A | Encounter for antenatal screening for other genetic defects |
| Z36.9 | Encounter for antenatal screening, unspecified |
| Z40.00 | Encounter for prophylactic removal of unspecified organ |
| Z40.01 | Encounter for prophylactic removal of breast |
| Z40.02 | Encounter for prophylactic removal of ovary(s) |
| Z40.09 | Encounter for prophylactic removal of other organ |
| Z40.8 | Encounter for other prophylactic surgery |
| Z40.9 | Encounter for prophylactic surgery, unspecified |
| Z41.1 | Encounter for cosmetic surgery |
| Z41.2 | Encounter for routine and ritual male circumcision |
| Z41.3 | Encounter for ear piercing |
| Z41.8 | Encounter for other procedures for purposes other than remedying health state |
| Z41.9 | Encounter for procedure for purposes other than remedying health state, unspecified |
| Z46.1 | Encounter for fitting and adjustment of hearing aid |
| Z56.0 | Unemployment, unspecified |
| Z56.2 | Threat of job loss |
| Z56.3 | Stressful work schedule |
| Z56.4 | Discord with boss and workmates |
| Z56.5 | Uncongenial work environment |
| Z56.6 | Other physical and mental strain related to work |
| Z56.81 | Sexual harassment on the job |
| Z56.82 | Military deployment status |
| Z56.89 | Other problems related to employment |

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| Code | Description |
|----------------|--|
| Z56.9 | Unspecified problems related to employment |
| Z57.0 | Occupational exposure to noise |
| Z57.1 | Occupational exposure to radiation |
| Z57.2 | Occupational exposure to dust |
| Z57.31 | Occupational exposure to environmental tobacco smoke |
| Z57.39 | Occupational exposure to other air contaminants |
| Z57.4 | Occupational exposure to toxic agents in agriculture |
| Z57.5 | Occupational exposure to toxic agents in other industries |
| Z57.6 | Occupational exposure to extreme temperature |
| Z57.7 | Occupational exposure to vibration |
| Z57.8 | Occupational exposure to other risk factors |
| Z57.9 | Occupational exposure to unspecified risk factor |
| Z58.6 | Inadequate drinking-water supply |
| Z59.00 | Homelessness unspecified |
| Z59.01 | Sheltered homelessness |
| Z59.02 | Unsheltered homelessness |
| Z59.1 | Inadequate housing |
| Z59.2 | Discord with neighbors, lodgers and landlord |
| Z59.3 | Problems related to living in residential institution |
| Z59.41 | Food insecurity |
| Z59.48 | Other specified lack of adequate food |
| Z59.5 | Extreme poverty |
| Z59.6 | Low income |
| Z59.7 | Insufficient social insurance and welfare support |
| Z59.811 | Housing instability, housed, with risk of homelessness |
| Z59.812 | Housing instability, housed, homelessness in past 12 months |
| Z59.819 | Housing instability, housed unspecified |
| *Z59.82 | *Transportation insecurity |
| *Z59.86 | *Financial insecurity |
| *Z59.87 | *Material hardship |
| Z59.89 | Other problems related to housing and economic circumstances |
| Z59.9 | Problem related to housing and economic circumstances, unspecified |
| Z60.2 | Problems related to living alone |
| Z62.21 | Child in welfare custody |

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| Code | Description |
|--------|--|
| Z71.0 | Person encountering health services to consult on behalf of another person |
| Z74.1 | Need for assistance with personal care |
| Z74.2 | Need for assistance at home and no other household member able to render care |
| Z74.3 | Need for continuous supervision |
| Z74.8 | Other problems related to care provider dependency |
| Z74.9 | Problem related to care provider dependency, unspecified |
| Z75.5 | Holiday relief care |
| Z76.0 | Encounter for issue of repeat prescription |
| Z76.1 | Encounter for health supervision and care of foundling |
| Z76.2 | Encounter for health supervision and care of other healthy infant and child |
| Z76.3 | Healthy person accompanying sick person |
| Z76.4 | Other boarder to healthcare facility |
| Z76.81 | Expectant parent(s) prebirth pediatrician visit |
| Z80.1 | Family history of malignant neoplasm of trachea, bronchus and lung |
| Z80.2 | Family history of malignant neoplasm of other respiratory and intrathoracic organs |
| Z80.49 | Family history of malignant neoplasm of other genital organs |
| Z80.51 | Family history of malignant neoplasm of kidney |
| Z80.52 | Family history of malignant neoplasm of bladder |
| Z80.59 | Family history of malignant neoplasm of other urinary tract organ |
| Z80.6 | Family history of leukemia |
| Z80.7 | Family history of other malignant neoplasms of lymphoid, hematopoietic and related tissues |
| Z80.8 | Family history of malignant neoplasm of other organs or systems |
| Z80.9 | Family history of malignant neoplasm, unspecified |
| Z81.0 | Family history of intellectual disabilities |
| Z81.1 | Family history of alcohol abuse and dependence |
| Z81.2 | Family history of tobacco abuse and dependence |
| Z81.3 | Family history of other psychoactive substance abuse and dependence |
| Z81.4 | Family history of other substance abuse and dependence |
| Z81.8 | Family history of other mental and behavioral disorders |
| Z82.0 | Family history of epilepsy and other diseases of the nervous system |
| Z82.1 | Family history of blindness and visual loss |
| Z82.2 | Family history of deafness and hearing loss |
| Z82.3 | Family history of stroke |

***January 2023 Changes
ICD-10-CM Version – Red**



**Medicare National Coverage Determinations (NCD)
Coding Policy Manual and Change Report (ICD-10-CM)**

| Code | Description |
|---------|---|
| Z82.41 | Family history of sudden cardiac death |
| Z82.49 | Family history of ischemic heart disease and other diseases of the circulatory system |
| Z82.5 | Family history of asthma and other chronic lower respiratory diseases |
| Z82.61 | Family history of arthritis |
| Z82.62 | Family history of osteoporosis |
| Z82.69 | Family history of other diseases of the musculoskeletal system and connective tissue |
| Z82.71 | Family history of polycystic kidney |
| Z82.79 | Family history of other congenital malformations, deformations and chromosomal abnormalities |
| Z82.8 | Family history of other disabilities and chronic diseases leading to disablement, not elsewhere classified |
| Z83.0 | Family history of human immunodeficiency virus [HIV] disease |
| Z83.1 | Family history of other infectious and parasitic diseases |
| Z83.2 | Family history of diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism |
| Z83.3 | Family history of diabetes mellitus |
| Z83.41 | Family history of multiple endocrine neoplasia [MEN] syndrome |
| Z83.49 | Family history of other endocrine, nutritional and metabolic diseases |
| Z83.511 | Family history of glaucoma |
| Z83.518 | Family history of other specified eye disorder |
| Z83.52 | Family history of ear disorders |
| Z83.6 | Family history of other diseases of the respiratory system |
| Z83.71 | Family history of colonic polyps |
| Z83.79 | Family history of other diseases of the digestive system |
| Z84.0 | Family history of diseases of the skin and subcutaneous tissue |
| Z84.1 | Family history of disorders of kidney and ureter |
| Z84.2 | Family history of other diseases of the genitourinary system |
| Z84.3 | Family history of consanguinity |
| Z84.81 | Family history of carrier of genetic disease |
| Z84.89 | Family history of other specified conditions |

***January 2023 Changes
ICD-10-CM Version – Red**



Reasons for Denial for All Lab NCDs

NOTE: This section includes CMS's interpretation of its longstanding policies pertaining to nationally covered laboratory services, and is included for informational purposes.

- Tests for screening purposes that are performed in the absence of signs, symptoms, complaints, or personal history of disease or injury are not covered except as explicitly authorized by statute.
- Tests for administrative purposes, including exams required by insurance companies, business establishments, government agencies, or other third parties, are not covered.
- Tests that are not reasonable and necessary for the diagnosis or treatment of an illness or injury are not covered by statute.
- Failure to provide documentation of the medical necessity of tests might result in denial of claims. The documentation may include notes documenting relevant signs, symptoms, or abnormal findings that substantiate the medical necessity for ordering the tests. In addition, failure to provide independent verification that the test was ordered by the treating physician (or qualified nonphysician practitioner) through documentation in the physician's office might result in denial.
- A claim for a test for which there is a national coverage policy will be denied as not reasonable and necessary if the claim is submitted without an ICD-10-CM code or narrative diagnosis listed as covered in the policy unless other medical documentation justifying the necessity is submitted with the claim.
- If a national coverage policy identifies a frequency expectation, a claim for a test that exceeds that expectation may be denied as not reasonable and necessary, unless it is submitted with documentation justifying increased frequency.
- Tests that are not ordered by a treating physician or other qualified treating nonphysician practitioner acting within the scope of their license and in compliance with Medicare requirements will be denied as not reasonable and necessary.
- Failure of the clinical laboratory performing the test to have the appropriate Clinical Laboratory Improvement Amendments of 1988 (CLIA) certificate will result in denial of claims.



Coding Guidelines for All Lab NCDs

1. On and after the implementation date for ICD-10-CM coding of Medicare billing claims, a claim for a clinical diagnostic laboratory service must include a valid ICD-10-CM diagnosis code. When a diagnosis has not been established by the physician, codes that describe symptoms and signs, as opposed to diagnoses, should be provided (see also bullet #5 below).

Please note that ICD-10-CM codes for diagnoses are not required (and will not be effective) for Medicare billing transactions prior to October 1, 2015. Please use ICD-9-CM codes for diagnoses prior to that date.

Please check the CMS website www.cms.gov/ICD10 for more information on the implementation of ICD-10-CM codes.

2. Medicare distinguishes ‘screening’ from ‘diagnostic uses’ of tests. ‘Screening’ is testing for disease or disease precursors so that early detection and treatment can be provided for those who test positive for the disease. Screening tests are performed when no specific sign, symptom, or diagnosis is present and the beneficiary has not been exposed to a disease.

In contrast, ‘diagnostic’ testing is testing to rule out or to confirm a suspected diagnosis because of a sign and/or symptom in the beneficiary. In these cases, the sign or symptom should be used to explain the reason for the test.

Some laboratory tests are covered by the Medicare program for screening purposes (for example, NCD # 210.1, Prostate Cancer Screening Tests). However, this manual focuses only on coding policies for diagnostic uses of laboratory services (for example, the test for prostate specific antigen (PSA)).

3. When the reason for performing a test is because the beneficiary has had contact with, or exposure to, a communicable disease, the appropriate code from category Z20, ‘Contact with or exposure to communicable diseases’, should be assigned. However, on review, the test might still be considered screening and not covered by Medicare.

4. All digits required by ICD-10-CM coding conventions must be used. A code is invalid if it has not been coded with all digits/characters required for that code.

5. The beneficiary’s condition(s) and/or diseases should be coded in ICD-10-CM to the highest degree of certainty for that encounter/visit, such as signs, symptoms, abnormal test results, or other reasons for the visit. When a non-specific ICD-10-CM code is submitted, the underlying sign, symptom, or condition must be related to the indications for the test.



Additional Coding Guideline(s)

Note: For any additional guideline(s) about ICD-10-CM coding for a specific diagnostic test service, please see the section “Limitations” in each NCD following the code list table.

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Check for Updates

the efficiency of our prior authorization processes, increase provider willingness to submit requests electronically, reduce provider burden, decrease delays in patient care, and promote high-quality, affordable health care.

In sum, we continue to believe prior authorization is an effective mechanism to ensure Medicare beneficiaries receive medically necessary care while protecting the Medicare Trust Funds from unnecessary increases in volume by virtue of improper payments without adding onerous new documentation requirements. A broad program integrity

strategy must use a variety of tools to best account for potential fraud, waste, and abuse, including unnecessary increases in volume. We believe prior authorization for these services will be an effective method for controlling unnecessary increases in the volume of these services and expect that it will reduce the instances in which Medicare pays for services that are determined not to be medically necessary.

After consideration of the public comments we received, we are finalizing our proposal to add the Facet joint interventions service category to

the list of hospital outpatient department services requiring prior authorization with modification. In particular, we are finalizing an implementation date for prior authorization for the Facet joint interventions service category of July 1, 2023, rather than the March 1, 2023 implementation date we proposed and making this change in the proposed regulation text at § 419.83(a)(3). Other than this change in the implementation date, we are finalizing the proposed regulation text changes as proposed.

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TABLE 103: FINAL LIST OF OUTPATIENT DEPARTMENT SERVICES THAT REQUIRE PRIOR AUTHORIZATION

| Beginning for service dates on or after July 1, 2020 | |
|--|---|
| Code | (i) Blepharoplasty, Blepharoptosis Repair, and Brow Ptosis Repair ³³⁰ |
| | |
| 15820 | Blepharoplasty, lower eyelid |
| 15821 | Blepharoplasty, lower eyelid; with extensive herniated fat pad |
| 15822 | Blepharoplasty, upper eyelid |
| 15823 | Blepharoplasty, upper eyelid; with excessive skin weighting down lid |
| 67900 | Repair of brow ptosis (supraciliary, mid-forehead or coronal approach) |
| 67901 | Repair of blepharoptosis; frontalis muscle technique with suture or other material (eg, banked fascia) |
| 67902 | Repair of blepharoptosis; frontalis muscle technique with autologous fascial sling (includes obtaining fascia) |
| 67903 | Repair of blepharoptosis; (tarso) levator resection or advancement, internal approach |
| 67904 | Repair of blepharoptosis; (tarso) levator resection or advancement, external approach |
| 67906 | Repair of blepharoptosis; superior rectus technique with fascial sling (includes obtaining fascia) |
| 67908 | Repair of blepharoptosis; conjunctivo-tarso-Muller's muscle-levator resection (eg, Fasanella-Servat type) |
| | |
| Code | (ii) Botulinum Toxin Injection |
| | |
| 64612 | Chemodenervation of muscle(s); muscle(s) innervated by facial nerve, unilateral (eg, for blepharospasm, hemifacial spasm) |
| 64615 | Chemodenervation of muscle(s); muscle(s) innervated by facial, trigeminal, cervical spinal and accessory nerves, bilateral (eg, for chronic migraine) |
| J0585 | Injection, onabotulinumtoxin, 1 unit |
| J0586 | Injection, abobotulinumtoxin, 5 units |
| J0587 | Injection, rimabotulinumtoxinb, 100 units |
| J0588 | Injection, incobotulinumtoxin a, 1 unit |

| | |
|-------|---|
| Code | (iii) Panniculectomy, Excision of Excess Skin and Subcutaneous Tissue (Including Lipectomy), and related services |
| 15830 | Excision, excessive skin and subcutaneous tissue (includes lipectomy); abdomen, infraumbilical panniculectomy |
| 15847 | Excision, excessive skin and subcutaneous tissue (includes lipectomy), abdomen (eg, abdominoplasty) (includes umbilical transposition and fascial plication) |
| 15877 | Suction assisted lipectomy; trunk |
| Code | (iv) Rhinoplasty, and related services ³³¹ |
| 20912 | Cartilage graft; nasal septum |
| 21210 | Graft, bone; nasal, maxillary or malar areas (includes obtaining graft) |
| 30400 | Rhinoplasty, primary; lateral and alar cartilages and/or elevation of nasal tip |
| 30410 | Rhinoplasty, primary; complete, external parts including bony pyramid, lateral and alar cartilages, and/or elevation of nasal tip |
| 30420 | Rhinoplasty, primary; including major septal repair |
| 30430 | Rhinoplasty, secondary; minor revision (small amount of nasal tip work) |
| 30435 | Rhinoplasty, secondary; intermediate revision (bony work with osteotomies) |
| 30450 | Rhinoplasty, secondary; major revision (nasal tip work and osteotomies) |
| 30460 | Rhinoplasty for nasal deformity secondary to congenital cleft lip and/or palate, including columellar lengthening; tip only |
| 30462 | Rhinoplasty for nasal deformity secondary to congenital cleft lip and/or palate, including columellar lengthening; tip, septum, osteotomies |
| 30465 | Repair of nasal vestibular stenosis (eg, spreader grafting, lateral nasal wall reconstruction) |
| 30520 | Septoplasty or submucous resection, with or without cartilage scoring, contouring or replacement with graft |
| Code | (v) Vein Ablation, and related services |
| 36473 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, mechanochemical; first vein treated |
| 36474 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, mechanochemical; subsequent vein(s) treated in a single extremity, each through separate access sites |
| 36475 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, radiofrequency; first vein treated |
| 36476 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, radiofrequency; subsequent vein(s) treated in a single extremity, each through separate access sites |
| 36478 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, laser; first vein treated |
| 36479 | Endovenous ablation therapy of incompetent vein, extremity, inclusive of all imaging guidance and monitoring, percutaneous, laser; subsequent vein(s) treated in a single extremity, each through separate access sites |

| | |
|---|--|
| 36482 | Endovenous ablation therapy of incompetent vein, extremity, by transcatheter delivery of a chemical adhesive (eg, cyanoacrylate) remote from the access site, inclusive of all imaging guidance and monitoring, percutaneous; first vein treated |
| 36483 | Endovenous ablation therapy of incompetent vein, extremity, by transcatheter delivery of a chemical adhesive (eg, cyanoacrylate) remote from the access site, inclusive of all imaging guidance and monitoring, percutaneous; subsequent vein(s) treated in a single extremity, each through separate access sites |
| Beginning for service dates on or after July 1, 2021 | |
| Code | (i) Cervical Fusion with Disc Removal |
| 22551 | Arthrodesis, anterior interbody, including disc space preparation, discectomy, osteophylectomy and decompression of spinal cord and/or nerve roots; cervical below C2 |
| 22552 | Arthrodesis, anterior interbody, including disc space preparation, discectomy, osteophylectomy and decompression of spinal cord and/or nerve roots; cervical below C2, each additional interspace |
| Code | (ii) Implanted Spinal Neurostimulators ³³² |
| 63650 | Percutaneous implantation of neurostimulator electrode array, epidural |
| Beginning for service dates on or after July 1, 2023 | |
| Code | Facet Joint Interventions |
| 64490 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), cervical or thoracic; single level |
| 64491 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), cervical or thoracic; second level |
| 64492 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), cervical or thoracic; third and any additional level(s) |
| 64493 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), lumbar or sacral; single level |
| 64494 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), lumbar or sacral; second level |
| 64495 | Injection(s), diagnostic or therapeutic agent, paravertebral facet (zygapophyseal) joint (or nerves innervating that joint) with image guidance (fluoroscopy or CT), lumbar or sacral; third and any additional level(s) |
| 64633 | Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); cervical or thoracic, single facet joint |

| | |
|-------|---|
| 64634 | Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); cervical or thoracic, each additional facet joint |
| 64635 | Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); lumbar or sacral, single facet joint |
| 64636 | Destruction by neurolytic agent, paravertebral facet joint nerve(s), with imaging guidance (fluoroscopy or CT); lumbar or sacral, each additional facet joint |

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XXI. Overall Hospital Quality Star Rating**A. Background**

The Overall Hospital Quality Star Rating provides a summary of certain existing hospital quality information based on publicly available quality measure results reported through CMS programs in a way that is simple and easy for patients to understand, by assigning hospitals between one and five stars (85 FR 86193). The Overall Hospital Quality Star Rating was first introduced and reported on our Hospital Compare website in July 2016³³³ (now reported on its successor website at <https://www.medicare.gov/care-compare> and referred to as Care Compare) and has been refreshed multiple times, with the most current refresh planned for 2022.^{334 335 336 337 338 339 340} In the CY

³³⁰CPT 67911 (Correction of lid retraction) was removed on January 7, 2022.

³³¹CPT 21235 (Obtaining ear cartilage for grafting) was removed on June 10, 2020.

³³²CPT codes 63685 (Insertion or replacement of spinal neurostimulator pulse generator or receiver) and 63688 (Revision or removal of implanted spinal neurostimulator pulse generator or receiver) were temporarily removed from the list of OPD services that require prior authorization, as finalized in the CY 2021 OPPS/ASC final rule comment period.

³³³Centers for Medicare & Medicaid Services. (2016, July 27). First Release of the Overall Hospital Quality Star Rating on Hospital Compare. Retrieved from CMS.gov newsroom at: <https://www.cms.gov/newsroom/fact-sheets/first-release-overall-hospital-quality-star-rating-hospital-compare>.

³³⁴Centers for Medicare & Medicaid Services. (2016, May). Overall Hospital Quality Star Rating on Hospital Compare: July 2016 Updates and Specifications Report.

³³⁵Centers for Medicare & Medicaid Services. (2016, October). Overall Hospital Quality Star Rating on Hospital Compare: December 2016 Updates and Specifications Report.

³³⁶Centers for Medicare & Medicaid Services. (2017, October). Overall Hospital Quality Star Rating on Hospital Compare: July 2017 Updates and Specifications Report.

³³⁷Centers for Medicare & Medicaid Services. (2019, November 4). Overall Hospital Quality Star Rating on Hospital Compare: January 2020 Updates and Specifications Report. Retrieved from [qualitynet.org: https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2](https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2).

³³⁸Centers for Medicare & Medicaid Services. (2018, November 30). Overall Hospital Quality Star Rating on Hospital Compare: February 2019 Updates and Specifications Report. Retrieved from [qualitynet.org: https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2](https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2).

2021 OPPS/ASC final rule with comment period (85 FR 86182), we finalized a methodology to calculate the Overall Hospital Quality Star Rating. We refer readers to section XVI (Overall Hospital Quality Star Rating Methodology for Public Release in CY 2021 and Subsequent Years) of the CY 2021 OPPS/ASC final rule with comment period and 42 CFR 412.190 for details.

In the CY 2023 OPPS/ASC proposed rule (87 FR 44807-44809), we: (1) provided information on the previously finalized policy for inclusion of quality measure data from Veterans Health Administration (VHA) hospitals; (2) proposed to amend the language of § 412.190(c) to state that we would use publicly available measure results on Hospital Compare or its successor websites from a quarter within the prior twelve months; and (3) conveyed that although CMS intends to publish Overall Hospital Quality Star Ratings in 2023, we may apply the suppression policy if applicable.

B. Veterans Health Administration Hospitals

In the CY 2021 OPPS/ASC final rule with comment period (85 FR 86197 and 86198), we finalized a policy to include Veterans Health Administration hospitals' (VHA hospitals) quality measure data for the purpose of calculating the Overall Hospital Quality Star Ratings beginning with the 2023 refresh. In that final rule, we also stated that we intended to provide more information about the statistical impact of adding VHA hospitals to the Overall Star Rating and discuss procedural aspects in a future rule (85 FR 48999). Since the publication of the CY 2021 OPPS/ASC final rule, we conducted an internal analysis from February 28, 2022, through March 30, 2022, with

³³⁹Centers for Medicare & Medicaid Services. (2017, November). Star Methodology Enhancement for December 2017 Public Release. Retrieved from [www.qualitynet.org: https://qualitynet.org/outpatient/public-reporting/overall-ratings/resources](https://qualitynet.org/outpatient/public-reporting/overall-ratings/resources).

³⁴⁰Centers for Medicare & Medicaid Services. (2022, May 17). Overall Hospital Quality Star Rating on Hospital Compare: July 2022 Updates and Specifications Report. Retrieved from [qualitynet.org: https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2](https://qualitynet.org/inpatient/public-reporting/overall-ratings/resources#tab2).

measure data from all VHA hospitals in the calculation of the Overall Hospital Quality Star Ratings methodology. The internal analysis included a period of confidential reporting and feedback during which VHA hospitals reviewed their Overall Hospital Quality Star Ratings internal analysis results, and in addition, further familiarized themselves with the Overall Hospital Quality Star Ratings methodology and had the opportunity to ask questions. All VHA hospitals were made aware of the internal analysis and were provided the opportunity to participate. For the internal analysis, the Overall Hospital Quality Star Ratings were calculated using VHA hospital measure data along with subsection (d) hospitals and CAHs. The internal analysis included the same measures used for the April 2021 refresh of Overall Hospital Quality Star Ratings on our public reporting website, Care Compare. At the time of the 2022 VHA internal analysis, VHA hospitals in each peer group reported a similar number of measures when compared to non-VHA hospitals for most measure groups. VHA hospitals in the five-measure group peer group reported a lower median number of Safety and Readmission measures. VHA hospitals in all three peer groups reported fewer measures in the Timely and Effective Care measure group. The measurement periods for VHA and non-VHA hospitals were the same, except for the HAI-1, HAI-2, PSI 04, PSI 90, and OP-22 measures. The specific performance periods for these measures were provided to VHA hospitals during the internal analysis. The reasons for the differing measure reporting periods are:

- The HAI-1 and HAI-2 measures were first publicly reported for VHA hospitals in July 2021, but only included one quarter of measure data. Therefore, we chose to use the next public reporting, April 2022, which included four quarters of these measures' data.
- For the PSI 04 and PSI 90 measures, we used measure data that were publicly reported in July 2021. VHA hospitals first publicly reported these measures in October 2020; however, a different software was used for the measure calculations than the software used to calculate subsection (d) hospitals and CAHs measure data. We

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2. Whether the beneficiary and/or the healthcare provider or supplier knew or could reasonably have been expected to know that the item or service was not covered.

| Knowledge of the Non-covered Item/Service | Liability | Payment Responsibility |
|---|--|--|
| If the beneficiary knew, or should have known (e.g. a valid liability notice such as an ABN, Form CMS-R-131 was issued and the beneficiary consented to receiving the item or service). | Rests with the beneficiary | The beneficiary is responsible for making payment for the usual and customary charges to the healthcare provider or supplier for the denied item and/or service. |
| If the beneficiary did not know (and should not have known), and the healthcare provider or supplier knew, or should have known. | Rests with the healthcare provider or supplier | The beneficiary may not be charged for any costs related to the denied item and/or service, including copayments and deductibles. |
| If neither the beneficiary nor the healthcare provider or supplier knew, and could not reasonably be expected to have known. | Neither the beneficiary or the healthcare provider or supplier | The Medicare program makes payment for the assigned claim. |

20.1 - LOL Coverage Denials (Rev. 1, 10-01-03) (Rev.:4197; Issued: 01-11-19; Effective: 04-15-19; Implementation: 04-15-19)

A. Statutory Basis

The following table provides examples of denials based on §1862(a)(1), §1862(a)(9), §1879(e), or §1879(g) of the Act:

| Statutory Provision (section of the Act) | Description |
|---|--|
| §1862(a)(1)(A) | Items and services found to be not reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member. |
| §1862(a)(1)(B) & §1861(s)(10) | Pneumococcal vaccine and its administration, influenza vaccine and its administration, and hepatitis B vaccine and its administration furnished to an individual at high or intermediate risk of contracting hepatitis B, that are not reasonable and necessary for the prevention of illness. |
| §1862(a)(1)(C) | In the case of hospice care, items and services that are not reasonable and necessary for the palliation or management of terminal illness. |

| Statutory Provision (section of the Act) | Description |
|---|---|
| §1862(a)(1)(E) | Items and services that, in the case of research conducted pursuant to §1142 of the Act, are not reasonable and necessary to carry out the purposes of that section (which concerns research on outcomes of health care services and procedures). |
| §1862(a)(1)(F) | Screening mammography that is performed more frequently than is covered under §1834(c)(2) of the Act or that is not conducted by a facility described in §1834(c)(1)(B) of the Act and screening pap smears and screening pelvic exams performed more frequently than is provided for under §1861(nn) of the Act. |
| §1862(a)(1)(F) | Screening for glaucoma, which is performed more frequently than is provided under §1861(uu) of the Act. |
| §1862(a)(1)(G) | Prostate cancer screening tests (as defined in §1861(oo) of the Act), which are performed more frequently than is covered under such section. |
| §1862(a)(1)(H) | Colorectal cancer screening tests, which are performed more frequently than is covered under §1834(d) of the Act. |
| §1862(a)(1)(I) | The frequency and duration of home health services which are in excess of normative guidelines that the Secretary shall establish by regulation. |
| §1862(a)(1)(J) | Drugs or biologicals specified in §1847A(c)(6)(C) of the Act, for which payment is made under part B, furnished in a competitive area under §1847B of the Act, but not furnished by an entity under a contract under §1847(B) of the Act. |
| §1862(a)(1)(K) | An initial preventive physical examination, which is performed more than 1 year after the date the individual's first coverage period begins under Medicare Part B. |
| §1862(a)(1)(L) | Cardiovascular screening blood tests (as defined in §1861(xx)(1) of the Act), which are performed more frequently than is covered under §1861(xx)(2). |
| §1862(a)(1)(M) | A diabetes screening test (as defined in §1861(yy)(1) of the Act), which is performed more frequently than is covered under §1861(yy)(3) of the Act. |
| §1862(a)(1)(N) | An ultrasound screening for abdominal aortic aneurysm which is performed more frequently than is provided for under §1861(s)(2)(AA) of the Act. |
| §1862(a)(1)(O) | Kidney disease education services (as defined in §1861(ggg)(1) of the Act) which are furnished in excess of the number of sessions covered under §1861(ggg)(4) of the Act. |
| §1861(dd)(3)(A) | Hospice care determined to be non-covered because the beneficiary was not "terminally ill," as referenced by §1879(g)(2) of the Act since the Balanced Budget Act of 1997. |
| §1862(a)(1)(O) | Personalized prevention plan services (as defined in § 1861(hhh)(1) of the Act), which are performed more frequently than is covered under such section. |

| Statutory Provision (section of the Act) | Description |
|---|---|
| §1814(a)(2)(C) & §1835(a)(2)(A) on or after July 1, 1987 §1879(g)(1) before December 31, 1995 | Home health services determined to be non-covered because the beneficiary was not “homebound” or did not require “intermittent” skilled nursing care. |
| §1879(e) | Inpatient hospital services or extended care services if payment is denied solely because of an unintentional, inadvertent, or erroneous action that resulted in the beneficiary’s transfer from a certified bed (one that does not meet the requirements of §1861(e) or (j) of the Act) in a skilled nursing facility (SNF) or hospital. |
| §1862(a)(9) | Custodial care, unless otherwise permitted under paragraph §1862(a)(1)(C) of the Act. |

 **20.2 - Denials When the LOL Provision Does Not Apply**
(Rev.: 4197; Issued: 01-11-19; Effective: 04-15-19; Implementation: 04-15-19)

| Type of Denial | Description | Example(s) |
|--------------------|--|---|
| Categorical | Categorical Denials are circumstances in which the LOL provision does not apply because the Medicare payment denial is based on a statutory provision not referenced in §1879 of the Act. Refer to §1862(a) of the Act for a complete listing. | <ul style="list-style-type: none"> • Personal comfort items (§1862(a)(6) of the Act). • Routine physicals and most screening tests (§1862(a)(7) of the Act). • Most immunizations (vaccinations) (§1862(a)(7) of the Act). • Routine eye care, most eyeglasses and examinations (§1862(a)(7) of the Act). • Hearing aids and hearing aid examinations (§1862(a)(7) of the Act). • Cosmetic surgery (§1862(a)(10) of the Act). • Orthopedic shoes and foot supports (orthotics) (§1862(a)(8) of the Act). |

| Type of Denial | Description | Example(s) |
|-------------------------|---|--|
| | | <p>NOTE: §22.1 of this chapter provides a more expansive list of examples.</p> |
| <p>Technical</p> | <p>When coverage requirements are not met for a particular item or service, it is not a Medicare benefit; therefore, Medicare denies payment or when payment for a medically unreasonable or unnecessary item or service that is also barred because of failure to meet a condition of payment required by regulations.</p> | <ul style="list-style-type: none"> • Payment for the additional cost of a private room in a hospital or SNF is denied when the private accommodations are not required for medical reasons (§1861(v)(2) of the Act). • Payment for a dressing is denied because it does not meet the definition for “surgical dressings” (§1861(s)(5) of the Act). • Payment for SNF stays not preceded by the required 3-day hospital stay or Payment for SNF stay because the beneficiary did not meet the requirement for transfer to a SNF and for receiving covered services within 30 days after discharge from the hospital and because the special requirements for extension of the 30 days were not met (§1861(i) of the Act). • Drugs and biologicals which are usually self-administered by the patient. • Ambulance services denied because transportation by other means is not contraindicated or because regulatory criteria specified in 42 CFR 410.40, such as those relating to destination or nearest appropriate facility, are not met. (See the Medicare Benefit Policy Manual, Chapter 10) • Other items or services that must be denied under 42 CFR 410.12 through 410.105 of the Medicare regulations. |

20.2.1 - Categorical Denials

(Rev.: 4197; Issued: 01-11-19; Effective: 04-15-19; Implementation: 04-15-19)

Below is a more expansive list of examples of categorical denials:

| Statutory Provision (section of the Act) | Description |
|---|--|
| §1862(a)(12) | Dental care and dentures (in most cases). |
| §1862(a)(13) | Routine foot care and flat foot care. |
| §1862(a)(19) | Services under a physician's private contract. |
| §1862(a)(3) | Services paid for by a governmental entity that is not Medicare. |
| §1862(a)(4) | Health care received outside of the U. S. not covered by Medicare. |
| §1862(a)(11) | Services by immediate relatives. |
| §1862(a)(5) | Services required as a result of war. |
| §1862(a)(2) | Services for which there is no legal obligation to pay. |
| §1862(a)(21) | Home health services furnished under a plan of care, if the agency does not submit the claim. |
| §1862(a)(16) | Items and services excluded under the Assisted Suicide Funding Restriction Act of 1997. |
| §1862(a)(17) | Items or services furnished in a competitive acquisition area by any entity that does not have a contract with the Department of Health and Human Services (except in a case of urgent need). |
| §1862(a)(14) | Physicians' services performed by a physician assistant, midwife, psychologist, or nurse anesthetist, when furnished to an inpatient, unless they are furnished under arrangement with the hospital. |
| §1862(a)(18) | Items and services furnished to an individual who is a resident of a skilled nursing facility or of a part of a facility that includes a skilled nursing facility, unless they are furnished under arrangements by the skilled nursing facility. |
| §1862(a)(15) | Services of an assistant at surgery without prior approval from the peer review organization. |
| §1862(a)(20) | Outpatient occupational and physical therapy services furnished incident to a physician's services. |
| §1862(a)(22) | Claims submitted other than in an electronic form specified by the Secretary, subject to the exceptions set forth in §1862(h) of the Act. |

| Statutory Provision (section of the Act) | Description |
|---|---|
| §1862(a)(23) | Claims for the technical component of advanced diagnostic imaging services described in §1834(e)(1)(B) of the Act for which payment is made under the fee schedule established under §1848(b) of the Act and that are furnished by a supplier (as defined in §1861(d) of the Act), if such supplier is not accredited by an accreditation organization designated by the Secretary under §1834(e)(2)(B) of the Act. |
| §1862(a)(24) | Claims for renal dialysis services (as defined in §1881(b)(14)(B) of the Act) for which payment is made under such section unless such payment is made under such section to a provider of services or a renal dialysis facility for such services. |

30 - Determining Liability for Disallowed Claims Under §1879
(Rev.: 4197; Issued: 01-11-19; Effective: 04-15-19; Implementation: 04-15-19)

When a Medicare contractor determines that a review under the LOL provisions is appropriate under §20 of this chapter, the Medicare contractor must next determine who is liable, based on who knew, or should have known that Medicare was going to deny payment on the item or service. In order to make this determination, the contractor must take the following steps:

Version 04/15/2019
 Check for updates

Form Instructions
Advance Beneficiary Notice of Non-coverage (ABN)
OMB Approval Number: 0938-0566

Overview

The ABN is a notice given to beneficiaries in Original Medicare to convey that Medicare is not likely to provide coverage in a specific case. “Notifiers” include:

- Physicians, providers (including institutional providers like outpatient hospitals), practitioners and suppliers paid under Part B (including independent laboratories);
- Hospice providers and religious non-medical health care institutions (RNHCIs) paid exclusively under Part A; and
- Home health agencies (HHAs) providing care under Part A or Part B.

All of the aforementioned healthcare providers and suppliers must complete the ABN as described below in order to transfer potential financial liability to the beneficiary, and deliver the notice prior to providing the items or services that are the subject of the notice.

Medicare inpatient hospitals and skilled nursing facilities (SNFs) use other approved notices for Part A items and services when notice is required in order to shift potential financial liability to the beneficiary; however, these facilities must use the ABN for Part B items and services.

The ABN must be reviewed with the beneficiary or his/her representative and any questions raised during that review must be answered before it is signed. The ABN must be delivered far enough in advance that the beneficiary or representative has time to consider the options and make an informed choice. Employees or subcontractors of the notifier may deliver the ABN. ABNs are never required in emergency or urgent care situations. Once all blanks are completed and the form is signed, a copy is given to the beneficiary or representative. In all cases, the notifier must retain a copy of the ABN delivered to the beneficiary on file.

The ABN may also be used to provide notification of financial liability for items or services that Medicare never covers. When the ABN is used in this way, it is not necessary for the beneficiary to choose an option box or sign the notice.

ABN Changes

The ABN is a formal information collection subject to approval by the Executive Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA). As part of this process, the notice is subject to public comment and re-approval every 3 years. With the latest PRA submission, a change has been made to the ABN. In accordance with Title 18 of the Social Security Act, guidelines for Dual Eligible beneficiaries have been added to the ABN form instructions.

Completing the Notice

ABNs may be downloaded from the CMS website at: <http://www.cms.gov/Medicare/Medicare-General-Information/BNI/ABN.html>. Instructions for completion of the form are set forth below:

ABNs must be reproduced on a single page. The page may be either letter or legal-size, with additional space allowed for each blank needing completion when a legal-size page is used.

There are 10 blanks for completion in this notice, labeled from (A) through (J). We recommend that notifiers remove the lettering labels from the blanks before issuing the ABN to beneficiaries. Blanks (A)-(F) and blank (H) may be completed prior to delivering the notice, as appropriate. Entries in the blanks may be typed or hand-written, but should be large enough (i.e., approximately 12-point font) to allow ease in reading. (Note that 10 point font can be used in blanks when detailed information must be given and is otherwise difficult to fit in the allowed space.) The notifier must also insert the blank (D) header information into all of the blanks labeled (D) within the Option Box section, Blank (G). One of the check boxes in the Option Box section, Blank (G), must be selected by the beneficiary or his/her representative. Blank (I) should be a cursive signature, with printed annotation if needed in order to be understood.

Header:

Blanks A-C, the header of the notice, must be completed by the notifier prior to delivering the ABN.

- 1. Blank (A) Notifier(s):** Notifiers must place their name, address, and telephone number (including TTY number when needed) at the top of the notice. This information may be incorporated into a notifier's logo at the top of the notice by typing, hand-writing, pre-printing, using a label or other means.

If the billing and notifying entities are not the same, the name of more than one entity may be given in the Header as long as it is specified in the Additional Information (H) section who should be contacted for billing questions.

- 2. Blank (B) Patient Name:** Notifiers must enter the first and last name of the beneficiary receiving the notice, and a middle initial should also be used if there is one on the beneficiary's Medicare card. The ABN will not be invalidated by a misspelling or missing initial, as long as the beneficiary or representative recognizes the name listed on the notice as that of the beneficiary.
- 3. Blank (C) Identification Number:** Use of this field is optional. Notifiers may enter an identification number for the beneficiary that helps to link the notice with a related claim. The absence of an identification number does not invalidate the ABN. An internal filing number created by the notifier, such as a medical record number, may

be used. Medicare numbers (HICNs), Medicare beneficiary identifiers (MBIs), or Social Security numbers should not appear on the notice.

Body:

4. Blank (D): The following descriptors may be used in the Blank (D) fields:

Item
Service
Laboratory test
Test
Procedure
Care
Equipment

- The notifier must list the specific names of the items or services believed to be non-covered in the column directly under the header of Blank (D).
- In the case of partial denials, notifiers must list in the column under Blank (D) the excess component(s) of the item or service for which denial is expected.
- For repetitive or continuous non-covered care, notifiers must specify the frequency and/or duration of the item or service.
- General descriptions of specifically grouped supplies are permitted in this column. For example, “wound care supplies” would be a sufficient description of a group of items used to provide this care. An itemized list of each supply is generally not required.
- When a reduction in service occurs, notifiers must provide enough additional information so that the beneficiary understands the nature of the reduction. For example, entering “wound care supplies decreased from weekly to monthly” would be appropriate to describe a decrease in frequency for this category of supplies; just writing “wound care supplies decreased” is insufficient.
- Please note that there are a total of 7 Blank (D) fields that the notifier must complete on the ABN. Notifiers are encouraged to populate all of the Blank (D) fields in advance when a general descriptor such as “Item(s)/Service(s)” is used. All Blank (D) fields must be completed on the ABN in order for the notice to be considered valid.

- 5. Blank (E) Reason Medicare May Not Pay:** In the column under this header, notifiers must explain, in beneficiary friendly language, why they believe the items or services listed in the column under Blank (D) may not be covered by Medicare. Three commonly used reasons for non-coverage are:

“Medicare does not pay for this test for your condition.”

“Medicare does not pay for this test as often as this (denied as too frequent).”

“Medicare does not pay for experimental or research use tests.”

To be a valid ABN, there must be at least one reason applicable to each item or service listed in the column under Blank (D). The same reason for non-coverage may be applied to multiple items in Blank (D) when appropriate.

- 6. Blank (F) Estimated Cost:** Notifiers must complete the column under Blank (F) to ensure the beneficiary has all available information to make an informed decision about whether or not to obtain potentially non-covered services.

Notifiers must make a good faith effort to insert a reasonable estimate for all of the items or services listed under Blank (D). In general, we would expect that the estimate should be within \$100 or 25% of the actual costs, whichever is greater; however, an estimate that exceeds the actual cost substantially would generally still be acceptable, since the beneficiary would not be harmed if the actual costs were less than predicted.

Multiple items or services that are routinely grouped can be bundled into a single cost estimate. For example, a single cost estimate can be given for a group of laboratory tests, such as a basic metabolic panel (BMP). An average daily cost estimate is also permissible for long term or complex projections. As noted above, providers may also pre-print a menu of items or services in the column under Blank (D) and include a cost estimate alongside each item or service. If a situation involves the possibility of additional tests or procedures (such as in laboratory reflex testing), and the costs associated with such tests cannot be reasonably estimated by the notifier at the time of ABN delivery, the notifier may enter the initial cost estimate and indicate the possibility of further testing. Finally, if for some reason the notifier is unable to provide a good faith estimate of projected costs at the time of ABN delivery, the notifier may indicate in the cost estimate area that no cost estimate is available. We would not expect either of these last two scenarios to be routine or frequent practices, but the beneficiary would have the option of signing the ABN and accepting liability in these situations.

- 7. Blank (G) Options:** Blank (G) contains the following three options:

- **OPTION 1.** I want the (D) _____ listed above. You may ask to be paid now, but I also want Medicare billed for an official decision on payment, which is sent to me on a Medicare Summary Notice (MSN). I understand that if Medicare doesn't pay, I am responsible for payment, but I can appeal to Medicare by following the directions on the MSN. If Medicare does pay, you will refund any payments I made to you, less co-pays or deductibles.

This option allows the beneficiary to receive the items and/or services at issue and requires the notifier to submit a claim to Medicare. This will result in a payment decision that can be appealed.

Suppliers and providers who don't accept Medicare assignment may make modifications to Option 1 only as specified below under "**H. Additional Information.**"

*** Special guidance for people who are dually enrolled in both Medicare and Medicaid, also known as dually eligible individuals (has a Qualified Medicare Beneficiary (QMB) Program and/or Medicaid coverage) ONLY:**

Dually Eligible beneficiaries must be instructed to check **Option Box 1** on the ABN in order for a claim to be submitted for Medicare adjudication.

Strike through **Option Box 1** as provided below:

OPTION 1. I want the (D)_____ listed above. ~~You may ask to be paid now, but I also want Medicare billed for an official decision on payment, which is sent to me on a Medicare Summary Notice (MSN). I understand that if Medicare doesn't pay, I am responsible for payment, but I can appeal to Medicare by following the directions on the MSN.~~

These edits are required because the provider cannot bill the dual eligible beneficiary when the ABN is furnished. Providers must refrain from billing the beneficiary pending adjudication by both Medicare and Medicaid in light of federal law affecting coverage and billing of dual eligible beneficiaries. If Medicare denies a claim where an ABN was needed in order to transfer financial liability to the beneficiary, the claim may be crossed over to Medicaid or submitted by the provider for adjudication based on State Medicaid coverage and payment policy. Medicaid will issue a Remittance Advice based on this determination.

Once the claim is adjudicated by both Medicare and Medicaid, providers may only charge the patient in the following circumstances:

- If the beneficiary has QMB coverage without full Medicaid coverage, the ABN could allow the provider to shift financial liability to the beneficiary per Medicare policy.
- If the beneficiary has full Medicaid coverage and Medicaid denies the claim (or will not pay because the provider does not participate in Medicaid), the ABN could allow the provider to shift financial liability to the beneficiary per Medicare policy, subject to any state laws that limit beneficiary liability.

Note: These instructions should only be used when the ABN is used to transfer potential financial liability to the beneficiary and not in voluntary instances. More information on dual eligible beneficiaries may be found at: https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNProducts/downloads/Medicare_Beneficiaries_Dual_Eligibles_At_a_Glance.pdf

- **OPTION 2.** I want the (D)_____listed above, but do not bill Medicare. You may ask to be paid now as I am responsible for payment. **I cannot appeal if Medicare is not billed.**

This option allows the beneficiary to receive the non-covered items and/or services and pay for them out of pocket. No claim will be filed and Medicare will not be billed. Thus, there are no appeal rights associated with this option.

- **OPTION 3.** I don't want the (D)_____listed above. I understand with this choice I am not responsible for payment, and I cannot appeal to see if Medicare would pay.

This option means the beneficiary does not want the care in question. By checking this box, the beneficiary understands that no additional care will be provided; thus, there are no appeal rights associated with this option.

The beneficiary or his or her representative must choose only one of the three options listed in Blank (G). Unless otherwise instructed to do so according to the specific guidance provided in these instructions, the notifier must not decide for the beneficiary which of the 3 checkboxes to select. Pre-selection of an option by the notifier invalidates the notice. However, at the beneficiary's request, notifiers may enter the beneficiary's selection if he or she is physically unable to do so. In such cases, notifiers must annotate the notice accordingly.

If there are multiple items or services listed in Blank (D) and the beneficiary wants to receive some, but not all of the items or services, the notifier can accommodate this request by using more than one ABN. The notifier can furnish an additional ABN listing the items/services the beneficiary wishes to receive with the corresponding option.

If the beneficiary cannot or will not make a choice, the notice should be annotated, for example: "beneficiary refused to choose an option."

- 8. Blank (H) Additional Information:** Notifiers may use this space to provide additional clarification that they believe will be of use to beneficiaries. For example, notifiers may use this space to include:

- A statement advising the beneficiary to notify his or her provider about certain tests that were ordered, but not received;
- Information on other insurance coverage for beneficiaries, such as a Medigap policy, if applicable;
- An additional dated witness signature; or
- Other necessary annotations.

Annotations will be assumed to have been made on the same date as that appearing in Blank J, accompanying the signature. If annotations are made on different dates, those dates should be part of the annotations.

***Special guidance for non-participating suppliers and providers (those who don't accept Medicare assignment) ONLY:**

Strike the last sentence in the Option 1 paragraph with a single line so that it appears like this: ~~If Medicare does pay, you will refund any payments I made to you, less co-pays or deductibles.~~

This single line strike can be included on ABNs printed specifically for issuance when unassigned items and services are furnished. Alternatively, the line can be hand-penned on an already printed ABN. The sentence must be stricken and can't be entirely concealed or deleted. There is no CMS requirement for suppliers or the beneficiary to place initials next to the stricken sentence or date the annotations when the notifier makes the changes to the ABN before issuing the notice to the beneficiary.

When this sentence is stricken, the supplier should include the following CMS-approved unassigned claim statement in the (H) Additional Information section:

“This supplier doesn't accept payment from Medicare for the item(s) listed in the table above. If I checked Option 1 above, I am responsible for paying the supplier's charge for the item(s) directly to the supplier. If Medicare does pay, Medicare will pay me the Medicare-approved amount for the item(s), and this payment to me may be less than the supplier's charge.”

This statement can be included on ABNs printed for unassigned items and services, or it can be handwritten in a legible 10 point or larger font.

An ABN with the Option 1 sentence stricken must contain the CMS-approved unassigned claim statement as written above to be considered valid notice. Similarly, when the unassigned claim statement is included in the “Additional Information” section, the last sentence in Option 1 should be stricken.

Signature Box:

Once the beneficiary reviews and understands the information contained in the ABN, the Signature Box is to be completed by the beneficiary (or representative). This box cannot be completed in advance of the rest of the notice.

9. **Blank (I) Signature:** The beneficiary (or representative) must sign the notice to indicate that he or she has received the notice and understands its contents. If a representative signs on behalf of a beneficiary, he or she should write out “representative” in parentheses after his or her signature. The representative's name should be clearly legible or noted in print.
10. **Blank (J) Date:** The beneficiary (or representative) must write the date he or she signed the ABN. If the beneficiary has physical difficulty with writing and requests assistance in completing this blank, the date may be inserted by the notifier.

Disclosure Statement: The disclosure statements in the footer of the notice are required to be included on the document.

CMS will work with its contractors to ensure consistency when determining validity of the ABN in general. In addition, contractors will provide ongoing education to notifiers as needed to ensure proper notice delivery. Notifiers should contact the appropriate CMS regional office if they believe that a contractor inappropriately invalidated an ABN.

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Check for Updates