



MEDICAL POLICY STATEMENT

Nevada Marketplace

| Policy Name & Number | Date Effective |
|---|----------------|
| Peripheral Nerve Blocks for Treatment of Pain-NV MP-MM-1772 | 05/01/2026 |
| Policy Type | |
| MEDICAL | |

Medical Policy Statements are derived from literature based on and supported by clinical guidelines, nationally recognized utilization and technology assessment guidelines, other medical management industry standards, and published MCO clinical policy guidelines. Medically necessary services include, but are not limited to, those health care services or supplies that are proper and necessary for the diagnosis or treatment of disease, illness, or injury and without which the patient can be expected to suffer prolonged, increased, or new morbidity, impairment of function, dysfunction of a body organ or part, or significant pain and discomfort. These services meet the standards of good medical practice in the local area, are the lowest cost alternative, and are not provided mainly for the convenience of the member or provider. Medically necessary services also include those services defined in any Evidence of Coverage or Certificate of Coverage documents, Medical Policy Statements, Provider Manuals, Member Handbooks, and/or other plan policies and procedures.

Medical Policy Statements do not ensure an authorization or payment of services. Please refer to the plan contract (often referred to as the Evidence of Coverage or Certificate of Coverage) for the service(s) referenced in the Medical Policy Statement. Except as otherwise required by law, if there is a conflict between the Medical Policy Statement and the plan contract, then the plan contract will be the controlling document used to make the determination.

According to the rules of Mental Health Parity Addiction Equity Act (MHPAEA), coverage for the diagnosis and treatment of a behavioral health disorder will not be subject to any limitations that are less favorable than the limitations that apply to medical conditions as covered under this policy.

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A. Subject

Peripheral Nerve Blocks for Treatment of Pain

B. Background

Peripheral nerve blocks are injections of medication into a specific area of the body where nerves cause pain to a specific organ or body region. Nerve blocks cause the temporary interruption of impulse conduction in peripheral nerves or nerve trunks and may or may not contain a steroid, which can be used to treat pain. Various areas of pain require different types of nerve blocks that can be administered in numerous parts of the body with some of the most common blocks being sympathetic, peripheral, and occipital.

Sacroiliac and facet joint interventions, epidural steroid injections, and trigger point injections are addressed in other policies.

C. Definitions

- **Acute Pain** – Pain lasting less than 4 weeks.
- **Ambulatory Surgery** – Surgery performed in a hospital-based or freestanding ambulatory surgery center (ASC) with patient discharge the same day.
- **Chronic Pain** – Pain lasting more than 3 months, which is considered beyond normal healing time.
- **Conservative Therapy** – A multimodality plan of care including both active and inactive conservative therapies.
 - **Active Conservative Therapies** – Action or activities that strengthen muscle groups and target key spinal structures, including physical therapy, occupational therapy, a physician-supervised home exercise program (HEP), and/or chiropractic care.
 - **HEP** – A 6-week program requiring an exercise prescription and/or plan and a follow-up documented in the medical record after completion, or documentation of the inability to complete due to a stated physical reason (ie, increased pain, inability to physically perform exercise). Patient inconvenience and/or noncompliance without explanation does not constitute an inability to complete.
 - **Inactive Conservative Therapies** – Passive activities that aid in treating symptoms associated with pain, including rest, ice, heat, medical devices, acupuncture, TENS unit, and/or pharmacotherapy (prescription or over the counter [non-steroidal anti-inflammatory drugs, acetaminophen]).
 - **Transcutaneous Electrical Nerve Stimulator (TENS)** – A device that utilizes electrical current delivered through electrodes placed on the surface of the skin to decrease the patient's perception of pain by inhibiting the transmission of afferent pain nerve impulses and/or stimulating the release of endorphins. Use, frequency, duration, and start dates must be documented in the medical record.

The MEDICAL Policy Statement detailed above has received due consideration as defined in the MEDICAL Policy Statement Policy and is approved.

- **Emergent** – Medically necessary care, which is immediately needed to preserve life, prevent serious impairment to bodily functions, organs, or parts, or prevent placing the physical or mental health of a patient in serious jeopardy.
- **Low-Risk Procedure** – Procedures associated with minimal physiologic effect and excludes any intrathoracic, intra-abdominal, vascular, or orthopedic procedures.
- **Sub-Acute Pain** – Pain lasting between 4 to 12 weeks.

D. Policy

- I. CareSource considers peripheral nerve blocks (PNB), single injection, medically necessary when appropriate documentation for the treatment of acute pain or chronic pain are included only as part of an active component of a comprehensive pain management program. CareSource uses MCG Health guidelines to address criteria for specific nerve blocks. Documentation must include indications that **ALL** the following criteria are met:
 - A. PNB is performed as an ambulatory or outpatient procedure that (1) is not treating an emergent condition, (2) is a low risk procedure, and (3) requires no inpatient care for a preoperative disease or condition (eg, altered mental status, hypotension, hypoxemia, tachycardia).
 - B. Acute, sub-acute or chronic, neuropathic or radicular pain, as indicated by **1 or more** of the following:
 1. cancer-related pain
 2. complex regional pain syndrome (CRPS)
 3. peripheral neuropathy with pain that limits activities of daily living, excluding diabetic neuropathy
 4. peripheral vascular disease with rest pain
 5. acute herpes zoster of face or neck and prevention of postherpetic neuralgia
 6. pancreatic pain, pelvic pain, or abdominal pain related to malignancy
 7. chronic, relapsing pancreatitis
 - C. Symptoms are poorly controlled by maximum medical therapy or intolerable side effects to such therapy.
 - D. Non-invasive treatment(s) have failed (eg, non-steroidal anti-inflammatory drugs (NSAIDs), exercise, physical therapy, spinal manipulation therapy).
 - E. No coagulopathy or thrombocytopenia is present.
 - F. No infection is present at or underlying the injection site.
- II. Acute or Sub-Acute Pain
PNB may provide means of analgesia for acute pain in the following (not an all-inclusive list):
 - A. Patients at risk of respiratory depression related to systemic or neuraxial opioids (eg, obstructive sleep apnea, severe obesity, underlying pulmonary disease, advanced age).
 - B. Patients with another indication to minimize opioid use (eg, chronic opioid use, intolerance to opioids).
 - C. Patients with acute, severe pain poorly managed with systemic medication.

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- D. Patients who cannot tolerate chiropractic or other physical and/or manipulative therapies.

III. Chronic Pain

CareSource considers PNB, single injection, medically necessary when appropriate documentation for the treatment of chronic pain is included only as part of an active component of a comprehensive pain management program when there has been a failure of conservative therapy, as evidenced by **ALL** the following:

- A. Documentation in the medical record of at least 6 weeks of active conservative therapy within the past 6 months **OR** inability to complete active conservative therapy due to contraindication, increased pain, or intolerance.
- B. Documentation in the medical record of at least 6 weeks inactive conservative therapy within the past 6 months.

Insufficient evidence supports the following uses of PNB for **chronic** pain and is not considered medically necessary:

- A. genicular nerve or branches for chronic knee pain
- B. cluneal nerve injections or blocks for chronic low back pain or pelvic pain
- C. pudendal blocks for chronic pelvic pain conditions

IV. Peripheral Radiofrequency Ablation (RFA) or Neurotomy

Radiofrequency ablation and/or neurotomy are considered experimental and investigational, or unproven for any indication, including but not limited to the treatment of acute or chronic pain due to insufficient evidence of efficacy in the peer reviewed literature.

V. Limitations and Exclusions

- A. A member can receive a maximum of 6 injections per area and anatomical side in a calendar year.
- B. Up to 2 anatomic sites (eg, specific nerve, plexus, or branch as defined by CPT code description) can be injected at any 1 session.
- C. Nerve blocks used as part of a surgical procedure or other medical procedure are not separately reimbursable but an inclusive component of that procedure. These injections will not be compensated separately or unbundled for coverage.
- D. Any procedure submitted for payment with an incorrect CPT code or description will be denied. It is the responsibility of the submitting provider to submit the most accurate and appropriate CPT/HCPCS code(s) for the product or service that is being provided. If requesting a block to a specific part of the body, coding to the highest level of specificity should be used.
- E. Exclusions
 1. Treatment of peripheral neuropathy due to diabetes.
 2. Use of nerve blocks with or without use of electrostimulation for treatment of multiple neuropathies or peripheral neuropathies caused by underlying systemic diseases as medical management using systemic medications is clinically indicated for the treatment of these conditions.

E. Conditions of Coverage
N/A

- F. Related Policies/Rules
Epidural Steroid Injections
Experimental or Investigational Item or Service
Facet Joint Interventions
Sacroiliac Joint Procedures
Trigger Point Injections

G. Review/Revision History

| | DATE | ACTION |
|-----------------------|------------|--------------------------------------|
| Date Issued | 06/18/2025 | Approved at Committee. |
| Date Revised | 01/28/2026 | Annual review; Approved at Committee |
| Date Effective | 05/01/2026 | |
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