Peripheral neuropathy/polyneuropathy overview

Definitions

**Neuropathy:** Nerve disease or disorder

**Peripheral neuropathy:** Disease or disorder of nerves located outside of the brain and spinal cord (peripheral nerves)

**Mononeuropathy:** Disease or disorder of a single nerve

**Polyneuropathy:** Disease or disorder of multiple nerves

**Mononeuritis multiplex:** A type of peripheral neuropathy in which there is damage to two or more different nerve areas

Three main types of nerves

- **Sensory:** Carry sensations from the body back to the brain and spinal cord
- **Motor:** Control muscles and movement
- **Autonomic:** Carry information to organs and glands and affect involuntary body functions, such as blood pressure, heart rate, perspiration, digestion, and bowel and bladder function

Signs and symptoms

- Gradual onset of numbness, prickling or tingling in feet or hands, which can spread upward into the legs and arms
- Sharp, jabbing, throbbing or burning pain
- Extreme sensitivity to touch
- Pain during activities that shouldn’t cause pain, such as pain in the feet when putting weight on them or when the feet are under a blanket
- Lack of coordination and falling
- Muscle weakness
- Patients may feel as if they are wearing gloves or socks when they are not
- Paralysis if motor nerves are affected

If autonomic nerves are affected, signs and symptoms might include:

- Heat intolerance
- Excessive sweating or not being able to sweat
- Bowel, bladder or digestive problems
- Drops in blood pressure that result in dizziness or lightheadedness

Causes

**Genetic:** Inherited or mutations

**Acquired:** Caused by another disorder or disease condition

Examples –

- Diabetes
- Autoimmune disorders
- Certain infections
- Tumors or growths
- Kidney and liver disorders
- Vitamin deficiencies
- Alcoholism
- Poisons and toxins
- Certain medications
- Physical injury

**Idiopathic:** Cause is unknown
Office note documentation – best practices

**Subjective** – Document the presence/absence of current symptoms related to peripheral neuropathy (e.g., numbness or tingling in upper or lower extremities, sensitivity to touch, glove or stocking sensation, etc.).

**Objective** – Include:
- Physical examination findings (e.g., muscle weakness, decreased sensation, results of monofilament and vibration testing, etc.)
- Other related diagnostic test results (e.g., EMG and nerve conduction studies, diagnostic imaging, laboratory tests, etc.)

**Assessment / final diagnostic statement**
- Describe current peripheral neuropathy to the highest level of specificity, using all applicable descriptors (diabetic, hereditary, autonomic, acute, chronic, etc.) and the exact site(s) or location(s).
- Spell out the diagnosis in full. Avoid use of abbreviations and acronyms.
- Clearly link peripheral neuropathy to the underlying cause, if known. Use linking terms such as “due to,” “secondary to,” “associated with,” “related to,” etc.
- Document the current status of peripheral neuropathy (stable, improved, worsening, etc.).

**Plan of treatment** – Document a clear and concise treatment plan.
- Clearly link peripheral neuropathy to medications being used to treat the condition.
- Include details of planned diagnostic testing and specialist referrals (e.g., neurologist, neurosurgeon).
- Include date of next appointment.

ICD-10-CM is a statistical classification; it is not a substitute for a healthcare provider’s final diagnostic statement. It is the provider’s responsibility to provide legible, clear, concise and complete documentation of each final diagnosis described to the highest level of specificity, which is then translated to a code for reporting purposes. It is not appropriate for providers to simply list a code number or select a code number from a list of codes in place of a written final diagnosis.

For electronic health records (EHRs) that insert diagnosis codes: The provider’s final statement of diagnosis should classify in ICD-10-CM to the EHR-inserted diagnosis code with description. Avoid mismatches between the two.

**Terms of uncertainty**
- For a confirmed diagnosis of peripheral neuropathy, do not use descriptors that imply uncertainty (such as “probable,” “apparently,” “likely” or “consistent with”).
- Do not document suspected peripheral neuropathy as if the diagnosis is confirmed. Rather, document the signs and symptoms in the absence of a confirmed diagnosis.

**Current versus historical**
- Do not describe current peripheral neuropathy as “history of.” In diagnosis coding, the phrase “history of” means the condition is historical and no longer exists as a current problem.

**Coding peripheral neuropathy/polyneuropathy**

To ensure accurate and specific diagnosis code assignment:
1. Review the entire medical record to verify the condition remains a current problem.
2. Note the exact diagnosis description documented in the medical record.
3. Search the alphabetic index for that specific diagnosis description and the corresponding code.
4. Verify the code in the tabular list, carefully following all instructional notes as applicable.

There are many different codes within the ICD-10-CM manual to represent the many different types and causes of peripheral neuropathy. Here are examples of some of the categories and subcategories under which various types of peripheral neuropathy and polyneuropathy are coded.

**Nerve, nerve root and plexus disorders (categories G50-G59)**
- G56 Mononeuropathies of upper limb
- G57 Mononeuropathies of lower limb
- G58 Other mononeuropathies
- G59 Mononeuropathy in diseases classified elsewhere

**Polyneuropathies and other disorders of the peripheral nervous system (categories G60-G65)**
- G60 Hereditary and idiopathic neuropathy
- G61 Inflammatory polyneuropathy
- G62 Other and unspecified polyneuropathies
- G63 Polyneuropathy in diseases classified elsewhere
- G65 Sequelae of inflammatory and toxic polyneuropathies

**Disorders of autonomic nervous system (category G90)**
- Subcategory G90.0- Idiopathic peripheral autonomic neuropathy