Introduction

The brain communicates with the body by sending electrical signals along nerves. When it comes time to go to the bathroom, the brain sends signals to specific nerves that travel through the lower back to the muscles that control the opening and closing of the bladder and bowel. Weak electrical signals may be used to address certain kinds of bowel and bladder problems that have not responded to other treatments. This procedure is known as sacral nerve neuromodulation. Another name for it is sacral nerve stimulation. This procedure involves implanting a small device under the skin in the lower back area. Small wires are also implanted so that the electric current activates the nerve important to either bladder or bowel function. This treatment usually is done in two steps. The first is a temporary placement to find out if sacral nerve stimulation works. The second is surgery to place the permanent implant. This policy describes when sacral nerve stimulation may be considered medically necessary.

Note: The Introduction section is for your general knowledge and is not to be taken as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals. It is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider also can be a place where medical care is given, like a hospital, clinic, or lab. This policy informs them about when a service may be covered.
**Note:** This policy addresses the InterStim® System (see **Regulatory Status** section).

<table>
<thead>
<tr>
<th>Service</th>
<th>Medical Necessity</th>
</tr>
</thead>
</table>
| **Urinary Incontinence and Non-obstructive Retention** | **Sacral nerve neuromodulation**  
A trial period of sacral nerve neuromodulation with either percutaneous nerve stimulation or a temporarily implanted lead may be considered medically necessary in patients who meet ALL of the following criteria:  
- There is a diagnosis of at least one of the following:  
  - Urge incontinence  
  - Urgency-frequency syndrome  
  - Nonobstructive urinary retention  
  - Overactive bladder (see **Definition of Terms**)  
  AND  
- There is documented failure or intolerance to at least 2 conventional conservative therapies (e.g., behavioral training such as bladder training, prompted voiding, or pelvic muscle exercise training, pharmacologic treatment for at least a sufficient duration to fully assess its efficacy, and/or surgical corrective therapy)  
  AND  
- The patient is an appropriate surgical candidate (see **Definition of Terms**)  
  AND  
- Incontinence is not related to a spinal cord injury or progressive, systemic neurologic condition (such as multiple sclerosis or diabetic neuropathy)  
 | **Permanent implantation, sacral nerve neuromodulation device**  
Permanent implantation of a sacral nerve neuromodulation device may be considered medically necessary in patients who meet ALL of the following criteria:  
- All of the criteria above are met  
  AND  
- A trial stimulation period demonstrates at least 50% improvement in symptoms over a period of at least 48 hours  
 | **Fecal Incontinence**  
| **Sacral nerve neuromodulation**  
A trial period of sacral nerve neuromodulation with either percutaneous nerve stimulation or a temporarily implanted lead  

<table>
<thead>
<tr>
<th>Service</th>
<th>Medical Necessity</th>
</tr>
</thead>
</table>
| lead may be considered medically necessary in patients who meet all of the following criteria: | • There is a diagnosis of chronic fecal incontinence of more than 2 incontinent episodes on average per week for more than 6 months, or for more than 12 months after vaginal childbirth <br> • There is documented failure or intolerance to conventional conservative therapy (eg, dietary modification, the addition of bulking and pharmacologic treatment) for at least a sufficient duration to fully assess its efficacy <br> • The patient is an appropriate surgical candidate (see Definition of Terms) <br> • The condition is not related to an anorectal malformation (eg, congenital anorectal malformation; defects of the external anal sphincter over 60 degrees; visible sequelae of pelvic radiation; active anal abscesses and fistulae) or chronic inflammatory bowel disease <br> • Incontinence is not related to a spinal cord injury or progressive, systemic neurologic condition (such as multiple sclerosis or diabetic neuropathy) <br> • The patient has not had rectal surgery in the previous 12 months, or in the case of rectal cancer, the patient has not had rectal surgery in the past 24 months | Permanent implantation, sacral nerve neuromodulation device may be considered medically necessary in patients who meet all of the following criteria: | • All of the criteria above are met <br> • A trial stimulation period demonstrates at least 50% improvement in symptoms over a period of at least 48 hours.
Other applications, urinary incontinence and nonobstructive retention

Other urinary/voiding applications of sacral nerve neuromodulation are considered investigational, including but not limited to treatment of stress incontinence or urge incontinence due to a neurologic condition (eg, detrusor hyperreflexia, multiple sclerosis, spinal cord injury or other types of chronic voiding dysfunction).

Other applications, chronic constipation or chronic pelvic pain

Sacral nerve neuromodulation is investigational in the treatment of chronic constipation or chronic pelvic pain.

Documentation Requirements

The patient’s medical records submitted for review should document that medical necessity criteria are met. The record should include clinical documentation of:

- Diagnosis/condition
- History and physical examination documenting the severity of the condition
- Conventional conservative therapies that have been tried and failed
- Any history of rectal surgery
- Any neurologic conditions or history of spinal cord injury
- If request is for permanent placement, results of trial

Coding

Sacral nerve neuromodulation involves several steps that are identified by the following codes.

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>CPT</td>
<td></td>
</tr>
<tr>
<td>64561</td>
<td>Percutaneous implantation of neurostimulator electrode array; sacral nerve (transforaminal placement)</td>
</tr>
<tr>
<td>64581</td>
<td>Incision for implantation of neurostimulator electrode array; sacral nerve (transforaminal placement)</td>
</tr>
<tr>
<td>HCPCS</td>
<td></td>
</tr>
<tr>
<td>A4290</td>
<td>Sacral nerve stimulation test lead, each</td>
</tr>
<tr>
<td>E0745</td>
<td>Stimulator electronic shock unit</td>
</tr>
<tr>
<td>Code</td>
<td>Description</td>
</tr>
<tr>
<td>---------</td>
<td>-----------------------------------------------------------------------------</td>
</tr>
<tr>
<td>L8679</td>
<td>Implantable neurostimulator, pulse generator, any type</td>
</tr>
<tr>
<td>L8680</td>
<td>Implantable neurostimulator electrode each (Note: Reported with 1-unit for each contact point on the implanted lead)</td>
</tr>
<tr>
<td>L8684</td>
<td>Radiofrequency transmitter (external) for use with implantable sacral root neurostimulator receiver for bowel and bladder management, replacement</td>
</tr>
<tr>
<td>L8685</td>
<td>Implantable neurostimulator pulse generator, single array, rechargeable, includes extension</td>
</tr>
<tr>
<td>L8686</td>
<td>Implantable neurostimulator pulse generator, single array, nonrechargeable, includes extension</td>
</tr>
<tr>
<td>L8687</td>
<td>Implantable neurostimulator pulse generator, dual array, rechargeable, includes extension</td>
</tr>
<tr>
<td>L8688</td>
<td>Implantable neurostimulator pulse generator, dual array, nonrechargeable, includes extension</td>
</tr>
</tbody>
</table>

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**Related Information**

**Definition of Terms**

**Inappropriate surgical candidates:** Includes patients with bleeding disorders, anatomical limitations, skin disease at risk for infection, psychiatric disease, or patients who are pregnant.\(^{30}\)

**Overactive bladder (OAB):** The International Continence Society has defined that overactive bladder syndrome as “urinary urgency, usually with increased daytime frequency and nocturia, with urinary incontinence or without urgency urinary incontinence, in the absence of urinary tract infection or other detectable disease” (available online at [https://www.ics.org/glossary/symptom/overactivebladderoaburgency syndrome](https://www.ics.org/glossary/symptom/overactivebladderoaburgency syndrome) Accessed June 2019).

**Evidence Review**
Description

Sacral nerve neuromodulation (SNM), also known as sacral nerve stimulation, involves the implantation of a permanent device that modulates the neural pathways controlling bladder or rectal function. This policy addresses use of SNM in the treatment of urinary or fecal incontinence, urinary or fecal nonobstructive retention, and chronic pelvic pain in patients with intact neural innervation of the bladder and/or rectum.

Background

Urinary and Fecal Incontinence

Urge incontinence is defined as leakage of urine when there is a strong urge to void. Urgency-frequency is an uncontrollable urge to urinate, resulting in very frequent, small volumes and is a prominent symptom of interstitial cystitis (also called bladder pain syndrome). Urinary retention is the inability to empty the bladder of urine completely. Fecal incontinence can arise from a variety of mechanisms, including rectal wall compliance, efferent and afferent neural pathways, central and peripheral nervous systems, and voluntary and involuntary muscles. Fecal incontinence is more common in women, due mainly to muscular and neural damage that may occur during vaginal delivery.

Treatment

Treatment using sacral nerve neuromodulation, also known as indirect sacral nerve stimulation, is one of several alternative modalities for patients with urinary or fecal incontinence (urge incontinence, significant symptoms of urgency-frequency, nonobstructive urinary retention) who have failed behavioral (eg, prompted voiding) and/or pharmacologic therapies.

The sacral nerve neuromodulation device consists of an implantable pulse generator that delivers controlled electrical impulses. This pulse generator is attached to wire leads that connect to the sacral nerves, most commonly the S3 nerve root. Two external components of the system help control the electrical stimulation. A control magnet, kept by the patient, is used to turn the device on or off. A console programmer is kept by the physician and used to adjust the settings of the pulse generator.

Before implantation of the permanent device, patients undergo an initial testing phase to estimate potential response to treatment. The first type of testing developed was percutaneous
nerve evaluation (PNE). This procedure is done with the patient under local anesthesia, using a test needle to identify the appropriate sacral nerve(s). Once identified, a temporary wire lead is inserted through the test needle and left in place for four to seven days. This lead is connected to an external stimulator, which is carried by patients in their pocket or on their belt. The results of this test phase are used to determine whether patients are appropriate candidates for the permanent device. If patients show a 50% or greater reduction in symptom frequency, they are deemed eligible for the permanent device.

The second type of testing is a 2-stage surgical procedure. In the first stage, a quadripolar-tined lead is implanted (stage 1). The testing phase can last as long as several weeks, and if patients show a 50% or greater reduction in symptom frequency, they can proceed to stage 2 of the surgery, which is permanent implantation of the neuromodulation device. The 2-stage surgical procedure has been used in various ways. They include its use instead of PNE, for patients who failed PNE, for patients with an inconclusive PNE, or for patients who had a successful PNE to refine patient selection further.

The permanent device is implanted with the patient under general anesthesia. The electrical leads are placed in contact with the sacral nerve root(s) via an incision in the lower back, and the wire leads are extended through a second incision underneath the skin, across the flank to the lower abdomen. Finally, a third incision is made in the lower abdomen where the pulse generator is inserted and connected to the wire leads. Following implantation, the physician programs the pulse generator to the optimal settings for that patient. The patient can switch the pulse generator between on and off by placing the control magnet over the area of the pulse generator for one to two seconds.

**Summary of Evidence**

For individuals with urinary incontinence who have failed conservative treatment who receive SNM, the evidence includes randomized controlled trials (RCTs), systematic reviews, and case series. The relevant outcomes are symptoms, morbid events, and treatment-related morbidity. Results from the RCTs and case series with long-term follow-up have suggested that SNM reduces symptoms of urge incontinence, urgency-frequency syndrome, nonobstructive urinary retention, and overactive bladder in selected patients. The evidence is sufficient to determine that the technology results in a meaningful improvement in the net health outcome.

For individuals with fecal incontinence who have failed conservative treatment who receive SNM, the evidence includes RCTs and systematic reviews. The relevant outcomes are symptoms, morbid events, and treatment-related morbidity. Although relatively small, the available trials
had a low risk of bias and demonstrated improvements in incontinence relative to alternatives. The evidence is sufficient to determine that the technology results in a meaningful improvement in the net health outcome.

For individuals with constipation who have failed conservative treatment who receive SNM, the evidence includes RCTs and systematic reviews. The relevant outcomes are symptoms, morbid events, and treatment-related morbidity. The available trials have not consistently reported improvements in outcomes with SNM. Additional studies are needed to demonstrate the health benefits of this technology. The evidence is insufficient to determine the effects of the technology on health outcomes.

For individuals with chronic pelvic pain who receive SNM, the evidence is limited to case series. The relevant outcomes are symptoms, morbid events, and treatment-related morbidity. The evidence is insufficient to determine the effects of the technology on health outcomes.

**Ongoing and Unpublished Clinical Trials**

Some currently unpublished trials that might influence this review are listed in Table 1.

**Table 1. Summary of Key Trials**

<table>
<thead>
<tr>
<th>NCT No.</th>
<th>Trial Name</th>
<th>Planned Enrollment</th>
<th>Completion Date</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ongoing</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NCT02434874</td>
<td>Sacral Nerve Stimulation to Treat Urgency Urinary Incontinence with Wireless Neuromodulation</td>
<td>60</td>
<td>Dec 2022 (ongoing)</td>
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<tr>
<td>NCT03261622</td>
<td>Sacral Nerve Stimulation for Fecal Incontinence – Placebo or Clinical Effective (SNS)</td>
<td>75</td>
<td>Nov 2020</td>
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<tr>
<td>NCT03139734</td>
<td>Sacral Neuromodulation for Pelvic Pain Associated with Endometriosis</td>
<td>50</td>
<td>May 2022</td>
</tr>
</tbody>
</table>

*denotes an industry-sponsored trial
NCT: national clinical trial
Clinical Input Received from Physician Specialty Societies and Academic Medical Centers

While the various physician specialty societies and academic medical centers may collaborate with and make recommendations during this process, through the provision of appropriate reviewers, input received does not represent an endorsement or position statement by the physician specialty societies or academic medical centers, unless otherwise noted.

In response to requests, input was received from 4 physician specialty societies and 2 academic medical centers while this policy was under review in 2012. Reviewers from 2 specialty societies and 2 academic medical centers provided opinions on the possible medical necessity of implantable leads for test stimulation, as part of a 2-stage process for device implantation. All four respondents supported the use of implantable leads for test stimulation as an alternative to percutaneous test stimulation for patients who had failed percutaneous test stimulation and/or for patients with inconclusive percutaneous test stimulation. Reasons for support included a longer period of interrupted treatment with stage-1 stimulation due to less lead migration and a higher rate of positive tests compared with percutaneous test stimulation.

Practice Guidelines and Position Statements

Urinary Disorders

American Urological Association

The American Urological Association (2014) updated its guidelines on the diagnosis and treatment of overactive bladder. The guidelines stated that sacral neuromodulation may be offered as a third-line treatment in carefully selected patients with severe refractory symptoms or in those who are not candidates for second-line therapy (eg, oral anti-muscarinics, oral β3-adrenoceptor agonists, transdermal oxybutynin) and are willing to undergo surgery.

American College of Obstetricians and Gynecologists

A practice bulletin on urinary incontinence from the American College of Obstetricians and Gynecologists (2005) considered sacral nerve neuromodulation to be beneficial for treating chronic voiding dysfunction. An updated 2015 practice bulletin on urinary incontinence from the College did not address sacral nerve stimulation (SNS).
Fecal Disorders

National Institute for Health and Care Excellence

The National Institute for Health and Care Excellence (2007) issued guidance on the management of fecal incontinence. The guidance was reviewed in 2014, and no changes were made. The guidance has recommended:

A trial of temporary sacral nerve stimulation should be considered for people with faecal incontinence in whom sphincter surgery is deemed inappropriate. All individuals should be informed of the potential benefits and limitations of this procedure and should undergo a trial stimulation period of at least 2 weeks to determine if they are likely to benefit. People with faecal incontinence should be offered sacral nerve stimulation on the basis of their response to percutaneous nerve evaluation during specialist assessment, which is predictive of therapy success.\(^{37}\)

American College of Gastroenterology

In its clinical guideline on the management of benign anorectal disorders, including fecal incontinence, the American College of Gastroenterology (2014) found "sacral nerve stimulation should be considered in [fecal incontinence] who do not respond to conservative therapy (strong recommendation, moderate quality of evidence)."\(^{38}\)

Pelvic Floor Society

The Pelvic Floor Society conducted a systematic review as the basis for practice recommendations on the use of SNS for the treatment of constipation.\(^{39}\) The systematic review assessed seven observational studies, all generally of poor quality due to inadequate description of methods. Due to inconsistent reporting on harms and treatment success, and heterogeneity in the patient populations, the Society could not recommend SNS.

Medicare National Coverage

The Centers for Medicare & Medicaid Services (2002) covers SNS for the “treatment of urinary urge incontinence, urgency-frequency syndrome, and urinary retention."\(^{40}\) SNS “involves both a
temporary test stimulation to determine if an implantable stimulator would be effective and a permanent implantation in appropriate candidates. Both the test and the permanent implantation are covered."

“The following limitations for coverage apply to all three indications:

- “Patients must be refractory to conventional therapy ... and be appropriate surgical candidates such that implantation with anesthesia can occur.

- “Patients with stress incontinence, urinary obstruction, and specific neurologic diseases ... that are associated with secondary manifestations ... are excluded.

- “Patients must have had successful test stimulation in order to support subsequent implantation. Before patients are eligible for permanent implantation, they must demonstrate a 50% or greater improvement through test stimulation. Improvement is measured through voiding diaries.”

**Regulatory Status**

In 1997, the Interstim® Sacral Nerve Stimulation system (Medtronic) was approved by the U.S. Food and Drug Administration (FDA) through the premarket approval process for the indication of urinary urge incontinence in patients who have failed or could not tolerate more conservative treatments. In 1999, the device received FDA approval for the additional indications of urgency-frequency and urinary retention in patients without mechanical obstruction.

In 2006, the Medtronic Interstim II® System (Medtronic) was approved by FDA through the premarket approval process for treatment of intractable cases of overactive bladder and urinary retention. The new device is smaller and lighter than the original and is reported to be suited for those with lower energy requirements or small stature. The device also includes updated software and programming options.

In 2011, the InterStim System was approved by FDA through the premarket approval process for the indication of chronic fecal incontinence in patients who have failed or could not tolerate more conservative treatments.

The InterStim® device has not been specifically approved by FDA for treatment of chronic pelvic pain.

FDA product code: EZW.
Note: This policy does not address pelvic floor stimulation, which refers to electrical stimulation of the pudendal nerve. Pelvic floor stimulation is addressed in a separate in policy (see Related Policies).

This policy does not address devices that provide direct SNS in patients with spinal cord injuries. The VOCARE® Bladder System/ FineTech Brindley Bladder Control System, a stimulator implanted in the sacral anterior nerve roots, is one device intended for patients with complete spinal cord injury and neurogenic bladder.

References


### History

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>03/02/99</td>
<td>Add to Surgery Section - New Policy</td>
</tr>
<tr>
<td>01/04/00</td>
<td>Replace Policy - Content updated; policy statement revised to include new FDA approved indications.</td>
</tr>
<tr>
<td>09/21/00</td>
<td>Replace Policy - Policy updated with reference to 2000 TEC assessment; policy statement unchanged.</td>
</tr>
<tr>
<td>04/09/02</td>
<td>Replace Policy - CPT code; policy statement unchanged.</td>
</tr>
<tr>
<td>05/13/03</td>
<td>Replace Policy - Policy updated; policy statement unchanged. New references added.</td>
</tr>
<tr>
<td>09/14/04</td>
<td>Replace Policy - Policy revised and expanded to address neuromodulation to treat fecal incontinence and constipation as investigational.</td>
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<tr>
<td>08/09/05</td>
<td>Replace Policy - Policy updated with literature search; no changes in policy statement.</td>
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<td>02/06/06</td>
<td>Codes updated - No other changes.</td>
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<tr>
<td>06/09/06</td>
<td>Disclaimer and Scope update - No further changes.</td>
</tr>
<tr>
<td>04/10/07</td>
<td>Cross Reference Update - No other changes.</td>
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<tr>
<td>11/13/07</td>
<td>Replace Policy - Policy updated with literature search; no changes in policy statement; references added; codes added</td>
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<tr>
<td>05/13/08</td>
<td>Code Updates - Code 787.6 added.</td>
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<td>Date</td>
<td>Comments</td>
</tr>
<tr>
<td>------------</td>
<td>------------------------------------------------------------------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>07/14/09</td>
<td>Replace Policy - Policy updated with literature search. Policy statement updated and clarified with selection criteria. References added.</td>
</tr>
<tr>
<td>08/10/10</td>
<td>Replace Policy - Policy updated with literature review through March 2010; references added and reordered. Policy statement on fecal incontinence has been changed from “investigational” to “may be medically necessary under specific conditions.”</td>
</tr>
<tr>
<td>07/12/11</td>
<td>Replace Policy - Policy updated with literature search through March 2011. References 7, 24, 26, 27, 29, 30, 34, and 36 added; other references reordered or removed. Policy statements unchanged. ICD-10 codes added to policy.</td>
</tr>
<tr>
<td>09/23/11</td>
<td>Related Policies updated; 2.01.27 added.</td>
</tr>
<tr>
<td>06/12/12</td>
<td>Replace policy. Policy updated with literature search through February 2012. Rationale and Policy Guidelines re-written. References 5, 13, 14 and 19-24 added; other references reordered or removed. Clinical input added. Implantable lead stimulation added as alternative stimulation method for eligible patients within medically necessary policy statements. Medically necessary policy statement for urinary incontinence changed to 2-part statement (has criteria for test stimulation and for permanent implantation). Material on methods of trial test stimulation added to Background and Rationale sections. Title changed to Sacral Nerve Neuromodulation/Stimulation. Code 64590 removed as it does not apply to this policy.</td>
</tr>
<tr>
<td>09/28/12</td>
<td>Update Coding Section – ICD-10 codes are now effective 10/01/2014. Add Related Policy 8.03.01.</td>
</tr>
<tr>
<td>06/10/13</td>
<td>Replace policy. Policy updated with literature search through March 13, 2013. Length of successful percutaneous test stimulation in medically necessary statements changed from at least 2 weeks to at least 1 week. Fecal incontinence policy statement separated into 2 statements; 1 on trial stimulation and 1 on permanent implantation. Edits made to statements so that criteria for fecal and urinary incontinence are similar, when applicable. References 12, 13, 16, 17 and 20 added; other references renumbered or removed.</td>
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<tr>
<td>01/28/14</td>
<td>Minor updated. CPT codes 95972 and 95973 listed within the Policy Guidelines extended to the Coding table to reflect consistency. Scope updated to indicate that this policy not part of Medicare Advantage.</td>
</tr>
<tr>
<td>03/21/14</td>
<td>Update Related Policies. Delete 7.01.106 and replace with 7.01.553.</td>
</tr>
<tr>
<td>06/09/14</td>
<td>Annual Review. Policy updated with literature review through March 5, 2014. References 5, 9, 12, 21, 30-31, and 34-35 added. Overactive bladder added to medically necessary statement on urinary incontinence. ICD-9 and ICD-10 diagnosis and procedure codes removed; this is performed inpatient.</td>
</tr>
<tr>
<td>06/27/14</td>
<td>Update Related Policies. Change title to 1.01.17.</td>
</tr>
<tr>
<td>09/08/14</td>
<td>Interim Update. Clarification added as #6 criteria under subheading “A” Fecal Incontinence. Specified the length of time after surgery when a trial of sacral nerve</td>
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<td>Date</td>
<td>Comments</td>
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<td>neurostimulation may be considered medically necessary. No new references added. Policy statement clarified as noted.</td>
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<td>04/17/15</td>
<td>Update Related Policies. Remove 7.01.553 as it was archived.</td>
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<tr>
<td>09/08/15</td>
<td>Annual Review. In both subheadings titled “B.2” the medically necessary statements that were changed from 1 week to “at least 48 hours” for the trial stimulation period. Policy updated with literature review through June 8, 2015; references 6, 16, 20, and 32 added. Policy statements revised as noted.</td>
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<tr>
<td>12/16/15</td>
<td>Update Related Policies. Remove 2.01.58 as it is archived.</td>
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<tr>
<td>01/29/16</td>
<td>Minor update. Added HCPCS code L8679.</td>
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<tr>
<td>06/01/16</td>
<td>Annual Review, approved May 10, 2016. Policy updated with literature review; reference added. No change to the policy statement. Updated the coding table.</td>
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<tr>
<td>03/01/17</td>
<td>Coding Update. Removed CPT code 95973 as it was deleted as of 01/01/2016.</td>
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<tr>
<td>08/25/17</td>
<td>Coding update, removed CPT codes 95970 and 95972. Moved into new format, no changes to policy statement.</td>
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<tr>
<td>07/01/18</td>
<td>Annual Review, approved June 22, 2018. Policy updated with literature review through February 2018; reference 40 added. Minor editorial changes to the Policy section; statements unchanged. Removed CPT codes 64585 and 64590.</td>
</tr>
<tr>
<td>04/01/19</td>
<td>Minor update, added Documentation Requirements section.</td>
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</tbody>
</table>

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. The Company adopts policies after careful review of published peer-reviewed scientific literature, national guidelines and local standards of practice. Since medical technology is constantly changing, the Company reserves the right to review and update policies as appropriate. Member contracts differ in their benefits. Always consult the member benefit booklet or contact a member service representative to determine coverage for a specific medical service or supply. CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). ©2019 Premera All Rights Reserved.

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  • Information written in other languages

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Email AppealsDepartmentinquines@Premera.com

You can also file a civil rights complaint with the U.S. Department of Health and Human Services, Office for Civil Rights, electronically through the Office for Civil Rights Complaint Portal, available at https://ocrportal.hhs.gov/ocr/portal/lobby.jsf, or by mail or phone at:
U.S. Department of Health and Human Services
200 Independence Avenue SW, Room S09F, HHH Building
Washington, D.C. 20201, 1-888-368-1019, 800-537-7697 (TDD)

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Appelez le 800-722-1471 (TTY: 800-842-5357).

Kreyol ayisyen (Creole):
Avi sila a gen Enfòmasyon Enpòtan Ilayadann. Avi sila a kapab genyen enfòmasyon enpòtan konsènan aplikasyon yon lan osawa konsènan kouvèti asirans lan atravé Premera Blue Cross. Kapab genyen dat ki enpòtan nan avi sila a. Ou ka gen pou pouan kék akson aven séten dat limit pou ka tenbe kouvèti asirans mante w la osaw pou yo ka ede w avèk depans yo. Se dwa w pou resewa enfòmasyon sa a ak asistans nan lang ou pale a, san ou pa gen pou peye pou sa. Rate nan 800-722-1471 (TTY: 800-842-5357).

Deutsche (German):

Hmoob (Hmong):

Ilokano (Ilocano):
Daytoy a Pakdaara ket naglaoon iti Napateg nga Impomarsa. Daytoy a pakdaara mabal ini nga adda ket naglaoon iti napateg nga impomarsa maianggepi ngi aklaysiyon nga coverage babaen iti Premera Blue Cross. Daytoy ket mabal ini daga importance a pelta iit daytoy a pakdaara. Mabalini nga adda rumbeng nga aramideng nga addang sakbay dagiti partikular a naituding nga adda talaw tapo napagtalaineyoo ti coverage ti salun-soy nga tungol kadagit gastos. Adda karbenganyo a mangala iti daytoy nga impomarsa ken tungol ti bukado a pagasaga nga awan ti bayadangay. Tumawag ti numero nga osawa 800-722-1471 (TTY: 800-842-5357).

Italiano (Italian):
Questo avviso contiene informazioni importanti. Questo avviso può contenere informazioni importanti sulla tua domanda o copertura attraverso Premera Blue Cross. Potrebbero essere date chiave in questo avviso. Potrebbe essere necessario un tuo intervento entro una scadenza determinata per consentirti di mantenere la tua copertura o sovvenzione. Hai il diritto di ottenere queste informazioni e assistenza nella tua lingua gratuitamente.
Chiamà 800-722-1471 (TTY: 800-842-5357).
Premera Blue Cross

Premera Blue Cross (TTY: 800-842-5357) 

This notice contains important information. This notice informs you of important dates for maintaining your health or assistance coverage through Premera Blue Cross. You have the right to receive this information and assistance in your language without cost. Call 800-722-1471 (TTY: 800-842-5357).

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