I. Policy: Sacral Nerve Stimulation

II. Purpose/Objective:
   To provide a policy of coverage regarding Sacral Nerve Stimulation

III. Responsibility:
   A. Medical Directors
   B. Medical Management

IV. Required Definitions

   1. Attachment – a supporting document that is developed and maintained by the policy writer or
department requiring/authoring the policy.
   2. Exhibit – a supporting document developed and maintained in a department other than the department
requiring/authoring the policy.
   3. Devised – the date the policy was implemented.
   4. Revised – the date of every revision to the policy, including typographical and grammatical changes.
   5. Reviewed – the date documenting the annual review if the policy has no revisions necessary.

V. Additional Definitions

Medical Necessity or Medically Necessary means Covered Services rendered by a Health Care Provider that the Plan
determines are:

   a. appropriate for the symptoms and diagnosis or treatment of the Member's condition, illness, disease or
      injury;
   b. provided for the diagnosis, and the direct care and treatment of the Member's condition, illness disease or
      injury;
   c. in accordance with current standards of good medical treatment practiced by the general medical
      community.
   d. not primarily for the convenience of the Member, or the Member's Health Care Provider; and
   e. the most appropriate source or level of service that can safely be provided to the Member. When applied
to hospitalization, this further means that the Member requires acute care as an inpatient due to the nature
of the services rendered or the Member's condition, and the Member cannot receive safe or adequate care
as an outpatient.

Medicaid Business Segment

Medically Necessary — A service, item, procedure, or level of care that is necessary for the proper treatment or
management of an illness, injury, or disability is one that:
• Will, or is reasonably expected to, prevent the onset of an illness, condition, injury or disability.
• Will, or is reasonably expected to, reduce or ameliorate the physical, mental or developmental effects of an illness, condition, injury or disability.
• Will assist the Member to achieve or maintain maximum functional capacity in performing daily activities, taking into account both the functional capacity of the Member and those functional capacities that are appropriate for Members of the same age.

DESCRIPTION:
A sacral nerve stimulator is composed of a pulse generator that transmits electrical impulses to the sacral nerves through an implanted wire electrode. These electrical impulses cause a contraction of the bladder muscles, giving the patient improved bladder control. After the electrodes are implanted, the member is provided with an external stimulator on a trial basis. If during the trial, it is determined that the stimulation is not effective or is unacceptable to the member, the electrodes are removed. If the trial proves successful, demonstrated by at least 50% decrease in incontinence symptoms, a stimulator and pulse generator are inserted subcutaneously and are connected to the implanted electrodes.

INDICATIONS:
• Urinary urge incontinence
• Urgency-frequency syndrome
• Urinary retention
• Fecal incontinence

CRITERIA FOR COVERAGE:

For urinary dysfunction: All must be met
• Medical documentation that the member’s urinary incontinence or retention is refractory to all conventional therapy (documented behavioral, pharmacologic and/or surgical corrective therapy) and the member is an appropriate surgical candidate such that implantation with anesthesia can occur.
• Demonstration of symptomatic improvement with a temporarily implanted electrode must precede permanent implantation.

For fecal incontinence: All must be met
Physician provided documentation of:
• Two or more fecal incontinence episodes per week for a minimum of 6 months, or for more than 12 months after vaginal childbirth; and
• Failure or intolerance of conventional therapy (e.g., Bowel re-training program, dietary modification, the addition of bulking and pharmacologic treatment); and
• Incontinence is not related to another neurologic condition such as peripheral neuropathy or complete spinal cord injury; and
• Fecal incontinence is not related to a treatable medical or surgical condition including, but not limited to congenital anorectal malformation, anal abscess and/or fistula, or visible sequelae of pelvic radiation; and
• the member is an appropriate surgical candidate such that implantation with anesthesia can occur.

For Neurogenic Bladder Secondary to Spinal Cord Injury: All must be met
• The Device must be FDA approved (e.g Vocare Bladder System / FineTech-Brindley Bladder Control System)
• patients who have clinically complete spinal cord lesions according to ASIA Classification (see: http://www.asia-spinalinjury.org/publications/59544_sc_Exam_Sheet_r4.pdf)
• intact parasympathetic innervations of the bladder
• skeletally mature and neurologically stable

Demonstration of at least 50% reduction in incontinent episodes with a temporarily implanted electrode must precede permanent implantation.

NOTE: Services related to component reimplantation or replacement in members previously approved for the implantation, or members having had the implantation prior to enrollment in the Plan, and who otherwise meet criteria for coverage, do not require prior authorization.

EXCLUSIONS:
There is insufficient evidence in the peer-reviewed, published medical literature to support the use of sacral nerve stimulation as a treatment of stress incontinence, urinary obstruction, and specific neurologic diseases (e.g., diabetes with
peripheral nerve involvement) which are associated with secondary manifestations of the above three indications. These applications are considered Experimental, Investigational, or Unproven and are NOT COVERED.

The Innova Feminine Incontinence Treatment System is considered Experimental, Investigational or Unproven and is NOT COVERED.

**Medicaid Business Segment:**
Any requests for services, that do not meet criteria set in the PARP, may be evaluated on a case by case basis.

Note: A complete description of the process by which a given technology or service is evaluated and determined to be experimental, investigational or unproven is outlined in MP 15 - Experimental Investigational or Unproven Services or Treatment.

**CODING ASSOCIATED WITH: Sacral Nerve Stimulation**

*The following codes are included below for informational purposes and may not be all inclusive. Inclusion of a procedure or device code(s) does not constitute or imply coverage nor does it imply or guarantee provider reimbursement. Coverage is determined by the member specific benefit plan document and any applicable laws regarding coverage of specific services. Please note that per Medicare coverage rules, only specific CPT/HCPCS Codes may be covered for the Medicare Business Segment. Please consult the CMS website at www.cms.gov or the local Medicare Administrative Carrier (MAC) for more information on Medicare coverage and coding requirements.*

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>64561</td>
<td>Percutaneous implantation of neurostimulator electrodes; sacral nerve (transforaminal placement)</td>
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<tr>
<td>64581</td>
<td>Open implantation of neurostimulator electrode array; sacral nerve (transforaminal placement)</td>
</tr>
<tr>
<td>64585</td>
<td>Revision or removal of peripheral neurostimulator electrodes</td>
</tr>
<tr>
<td>64590</td>
<td>Incision and subcutaneous placement of peripheral neurostimulator pulse generator or receiver, direct of inductive coupling</td>
</tr>
<tr>
<td>64595</td>
<td>Revision or removal of peripheral neurostimulator pulse generator or receiver</td>
</tr>
<tr>
<td>95970</td>
<td>Electronic analysis of implanted neurostimulator pulse generator system (e.g., rate, pulse amplitude and duration, configuration of wave form, battery status, electrode selectability, output modulation, cycling, impedance and patient compliance measurements); simple or complex brain, spinal cord, or peripheral (i.e., cranial nerve, peripheral nerve, autonomic nerve, neuromuscular) neurostimulator pulse generator/transmitter, without reprogramming</td>
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<tr>
<td>95971</td>
<td>Simple brain, spinal cord, or peripheral (i.e., peripheral nerve, autonomic nerve, neuromuscular neurostimulator pulse generator/transmitter, with intraoperative or subsequent programming</td>
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<tr>
<td>95972</td>
<td>Electronic analysis of implanted neurostimulator pulse generator system (e.g., rate, pulse amplitude, pulse duration, configuration of wave form, battery status, electrode selectability, output modulation, cycling, impedance and patient compliance measurements); complex spinal cord, or peripheral (i.e, peripheral nerve, sacral nerve, neuromuscular) (except cranial nerve) neurostimulator pulse generator/transmitter, with intraoperative or subsequent programming</td>
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<tr>
<td>A4290</td>
<td>Sacral nerve stimulation test lead, each</td>
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<tr>
<td>L8679</td>
<td>Implantable neurostimulator pulse generator, any type</td>
</tr>
<tr>
<td>L8680</td>
<td>Implantable neurostimulator electrode, each</td>
</tr>
<tr>
<td>L8683</td>
<td>Radiofrequency transmitter (external) for use with implantable neurostimulator radiofrequency receiver</td>
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<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>
<tr>
<td>L8689</td>
<td>External recharging system for battery (internal) for use with implantable neurostimulator, replacement only</td>
</tr>
<tr>
<td>C1767</td>
<td>Generator, neurostimulator (implantable), nonrechargeable</td>
</tr>
<tr>
<td>C1778</td>
<td>Lead, neurostimulator (implantable)</td>
</tr>
<tr>
<td>C1787</td>
<td>Patient programmer, neurostimulator</td>
</tr>
<tr>
<td>C1816</td>
<td>Receiver and/or transmitter, neurostimulator (implantable)</td>
</tr>
<tr>
<td>C1820</td>
<td>Generator, neurostimulator (implantable), non-high frequency, with rechargeable battery and charging system</td>
</tr>
<tr>
<td>C1822</td>
<td>Generator, neurostimulator (implantable), high frequency, with rechargeable battery and charging system</td>
</tr>
<tr>
<td>C1883</td>
<td>Adaptor/Extension, pacing lead or neurostimulator lead (implantable)</td>
</tr>
<tr>
<td>C1897</td>
<td>Lead, neurostimulator test kit (implantable)</td>
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LINE OF BUSINESS:
Eligibility and contract specific benefit limitations and/or exclusions will apply. Coverage statements found in the line of business specific benefit document will supersede this policy. For Medicare, applicable LCD’s and NCD’s will supersede this policy. For PA Medicaid Business segment, this policy applies as written.

REFERENCES:
Health Technology Trends, FDA Approves Implantable Electrical Stimulator for Patients with Urinary Incontinence, Vol. 9 No. 11; November 1997


Geisinger Clinic technology Assessment Committee, “Innova Feminine Incontinence Treatment System, April 27, 1994


American College of Obstetricians and Gynecologists (ACOG) FAQ Urinary Incontinence FAQ 081 February 2016

Centers for Medicare & Medicaid Services. National Coverage Determination (NCD) for Sacral Nerve Stimulation For Urinary Incontinence (230.18)


This policy will be revised as necessary and reviewed no less than annually.

Devised: 8/98

Revised: 7/99, 3/03; 3/04; 2/05; 5/10 (refs); 3/12 (added indication), 9/19 (Remove P/A)


Geisinger Health Plan may refer collectively to health care coverage sponsors Geisinger Health Plan, Geisinger Quality Options, Inc., and Geisinger Indemnity Insurance Company, unless otherwise noted. Geisinger Health Plan is part of Geisinger, an integrated health care delivery and coverage organization.

Coverage for experimental or investigational treatments, services and procedures is specifically excluded under the member's certificate with Geisinger Health Plan. Unproven services outside of an approved clinical trial are also specifically excluded under the member's certificate with Geisinger Health Plan. This policy does not expand coverage to services or items specifically excluded from coverage in the member's certificate with Geisinger Health Plan. Additional information can be found in MP015 Experimental, Investigational or Unproven Services.

Prior authorization and/or pre-certification requirements for services or items may apply. Pre-certification lists may be found in the member's contract specific benefit document. Prior authorization requirements can be found at https://www.geisinger.org/health-plan/providers/ghp-clinical-policies

Please be advised that the use of the logos, service marks or names of Geisinger Health Plan, Geisinger Quality Options, Inc. and Geisinger Indemnity Insurance Company on a marketing, press releases or any communication piece regarding the contents of this medical policy is strictly prohibited without the prior written consent of Geisinger Health Plan. Additionally, the above medical policy does not confer any endorsement by Geisinger Health Plan, Geisinger Quality Options, Inc. and Geisinger Indemnity Insurance Company regarding the medical service, medical device or medical lab test described under this medical policy.