



**BlueCross BlueShield
of Alabama**

Name of Blue Advantage Policy:
Gastric Electrical Stimulation

Policy #: 148
Category: Surgery

Latest Review Date: February 2020
Policy Grade: A

BACKGROUND:

Blue Advantage medical policy does not conflict with Local Coverage Determinations (LCDs), Local Medical Review Policies (LMRPs) or National Coverage Determinations (NCDs) or with coverage provisions in Medicare manuals, instructions or operational policy letters. In order to be covered by Blue Advantage the service shall be reasonable and necessary under Title XVIII of the Social Security Act, Section 1862(a)(1)(A). The service is considered reasonable and necessary if it is determined that the service is:

- 1. Safe and effective;*
- 2. Not experimental or investigational*;*
- 3. Appropriate, including duration and frequency that is considered appropriate for the service, in terms of whether it is:*
 - Furnished in accordance with accepted standards of medical practice for the diagnosis or treatment of the patient's condition or to improve the function of a malformed body member;*
 - Furnished in a setting appropriate to the patient's medical needs and condition;*
 - Ordered and furnished by qualified personnel;*
 - One that meets, but does not exceed, the patient's medical need; and*
 - At least as beneficial as an existing and available medically appropriate alternative.*

Routine costs of qualifying clinical trial services with dates of service on or after September 19, 2000 which meet the requirements of the Clinical Trials NCD are considered reasonable and necessary by Medicare. Providers should bill **Original Medicare for covered services that are related to **clinical trials** that meet Medicare requirements (Refer to Medicare National Coverage Determinations Manual, Chapter 1, Section 310 and Medicare Claims Processing Manual Chapter 32, Sections 69.0-69.11).*

POLICY:

Effective for dates of service on or after June 30, 2009:

Blue Advantage will treat **Gastric Electrical Stimulation** as a **covered** benefit for the treatment of nausea and vomiting from **chronic gastroparesis that is refractory for medical management** when **all** of the following criteria are met:

- Diagnosis of delayed gastric emptying has been made; **AND**
- Patient is refractory or intolerant of prokinetic medications and antiemetic medications; **AND**
- Nutritional status is poor, and either enteral tube feedings or total parental nutrition is medically necessary.

Blue Advantage will treat **Gastric Electrical Stimulation** as a **non-covered** benefit and as **investigational** for all other indications including but not limited to initial treatment of gastroparesis and treatment of obesity.

Blue Advantage does not approve or deny procedures, services, testing, or equipment for our members. Our decisions concern coverage only. The decision of whether or not to have a certain test, treatment or procedure is one made between the physician and his/her patient. Blue Advantage administers benefits based on the members' contract and medical policies. Physicians should always exercise their best medical judgment in providing the care they feel is most appropriate for their patients. Needed care should not be delayed or refused because of a coverage determination.

DESCRIPTION OF PROCEDURE OR SERVICE:

Gastric Electrical stimulation (GES) is performed using an implantable device designed to treat chronic drug-refractory nausea and vomiting secondary to gastroparesis of diabetic, idiopathic or post-surgical etiology. Gastric electrical stimulation has also been investigated as a treatment of obesity. The device may be referred to as a gastric pacemaker.

Treatment

Gastroparesis

Gastric electrical stimulation, also referred to as gastric pacing, using an implantable device, has been investigated primarily as a treatment for gastroparesis. Currently available devices consist of a pulse generator, which can be programmed to provide electrical stimulation at different frequencies, connected to intramuscular stomach leads that are implanted during laparoscopy or open laparotomy.

Obesity

Gastric electrical stimulation has also been investigated as a treatment of obesity. It is used to increase a feeling of satiety with subsequent reduced food intake and weight loss. The exact mechanisms resulting in changes in eating behavior are uncertain but may be related to neuro-hormonal modulation and/or stomach muscle stimulation.

KEY POINTS:

The most recent literature review was performed through December 9, 2019. The following is a summary of the key findings to date.

Summary of Evidence

For individuals with gastroparesis who receive GES, the evidence includes randomized controlled trials (RCTs), non randomized studies, and case series. Relevant outcomes are symptoms and treatment related morbidity. Several crossover RCTs have been published. A 2017 meta-analysis of these 5 RCTs did not find a significant benefit of GES on the severity of symptoms associated with gastroparesis. However, numerous case series and uncontrolled studies on GES have been published. These studies generally report improvements in symptoms following treatment. Patients who have refractory, chronic gastroparesis, gastric electrical stimulation may provide relief of symptoms for this specific population.

For individuals who have obesity who receive GES, the evidence includes one published randomized study as well as small case series and uncontrolled trials. Relevant outcomes are change in disease status and treatment-related morbidity. The published RCT (the SHAPE trial) did not show significant improvement in weight loss with gastric electrical stimulation compared with sham stimulation. The evidence is insufficient to determine the effects of the technology on health outcomes.

Practice Guidelines and Position Statements

National Institute for Health and Care Excellence

In 2014, the National Institute for Health and Care Excellence issued guidelines on gastroelectrical stimulation for gastroparesis that made the following recommendations:

- Current evidence on the efficacy and safety of gastric electrical stimulation for gastroparesis is adequate to support the use of this procedure with normal arrangements for clinical governance, consent, and audit.
- Clinicians should inform patients considering gastric electrical stimulation for gastroparesis that some patients do not get any benefit from it. They should also give patients detailed written information about the risk of complications, which can be serious, including the need to remove the device.
- Patient selection and follow-up should be done in specialist gastroenterology units with expertise in gastrointestinal motility disorders, and the procedure should only be performed by surgeons working in these units.

American College of Gastroenterology

The American College of Gastroenterology published a clinical practice guideline on management of gastroparesis in 2013. The recommendation was that:

“GES may be considered for compassionate treatment in patients with refractory symptoms, particularly nausea and vomiting. Symptom severity and gastric emptying have been shown to improve in patients with DG, but not in patients with IG or PSG. [Conditional recommendation (there is uncertainty about trade-offs), moderate level of evidence (further research would be likely to have an impact on the confidence in the estimate of effect).]”

An update is in progress from the American College of Gastroenterology.

U.S. Preventive Services Task Force Recommendations

Gastric electrical stimulation is not a preventive service.

KEY WORDS:

Gastric electrical stimulation (GES), gastroparesis, Enterra™ Therapy System, gastric pacemaker, gastric pacing

APPROVED BY GOVERNING BODIES:

In 2000, the Gastric Electrical Stimulator (GES) system (now called Enterra™ Therapy System), manufactured by Medtronic, Inc. (Minneapolis, MN) was approved by the U.S. Food and Drug Administration through the humanitarian device exemption process. The GES system consists of four components: the implanted pulse generator, two unipolar intramuscular stomach leads, the stimulator programmer, and the memory cartridge. With the exception of the intramuscular leads, all other components have been used in other implantable neurologic stimulators, such as spinal cord or sacral nerve stimulation. The intramuscular stomach leads are implanted either laparoscopically or during a laparotomy and are connected to the pulse generator, which is implanted in a subcutaneous pocket. The programmer sets the stimulation parameters, which are typically set at an “on” time of 0.1 second alternating with an “off” time of 5.0 second.

There are no gastric electrical stimulation devices approved by the U.S. Food and Drug Administration (FDA) for the treatment of obesity. The Transcend® Implantable Gastric Stimulation device, manufactured by Transneuronix and acquired by Medtronic in 2005, is currently available in Europe for treatment of obesity.

BENEFIT APPLICATION:

Coverage is subject to member’s specific benefits. Group specific policy will supersede this policy when applicable.

CURRENT CODING:

CPT Codes:

Laparoscopic procedures related to gastric stimulation electrodes for **morbid obesity** should be reported using code **43659 (unlisted laparoscopy procedure, stomach)**, and *laparotomy* procedures related to gastric stimulation electrodes for **morbid obesity** should be reported using **43999 (unlisted procedure, stomach)**.

43647	Laparoscopy, surgical; implantation or replacement of gastric neurostimulator electrodes, antrum
43648	Laparoscopy, surgical; revision or removal of gastric neurostimulator electrodes, antrum
43659	Unlisted laparoscopy procedure, stomach
43881	Implantation or replacement of gastric neurostimulator electrodes, antrum, open
43882	Revision or removal of gastric neurostimulator electrodes, antrum, open
43999	Unlisted procedure, stomach
64590	Insertion or replacement of peripheral or gastric neurostimulator pulse generator or receiver, direct or inductive coupling
64595	Revision or removal of peripheral or gastric neurostimulator pulse generator or receiver
95980	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 measurements) gastric neurostimulator pulse generator/transmitter; intraoperative, with programming
95981	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 measurements) gastric neurostimulator pulse generator/transmitter; subsequent, without reprogramming
95982	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 measurements) gastric neurostimulator pulse generator/transmitter; subsequent, with reprogramming

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POLICY HISTORY:

Adopted for Blue Advantage, March 2005
 Available for comment May 1-June 14, 2005
 Medical Policy Group, March 2006
 Available for comment March 23-May 8, 2006
 Medical Policy Group, November 2006
 Available for comment November 3-December 17, 2006
 Medical Policy Group, June 2009
 Available for comment June 30-August 13, 2009
 Medical Policy Group, October 2010
 Medical Policy Group, December 2011
 Medical Policy Group, January 2013
 Medical Policy Group, August 2013
 Medical Policy Group, August 2014
 Medical Policy Group, December 2015
 Medical Policy Group, February 2017
 Medical Policy Group, February 2018
 Medical Policy Group, February 2019
 Medical Policy Group, February 2020

This medical policy is not an authorization, certification, explanation of benefits, or a contract. Eligibility and benefits are determined on a case-by-case basis according to the terms of the member's plan in effect as of the date services are rendered. All medical policies are based on (i) research of current medical literature and (ii) review of common medical practices in the treatment and diagnosis of disease as of the date hereof. Physicians and other providers are solely responsible for all aspects of medical care and treatment, including the type, quality, and levels of care and treatment.

This policy is intended to be used for adjudication of claims (including pre-admission certification, pre-determinations, and pre-procedure review) in Blue Cross and Blue Shield's administration of plan contracts.