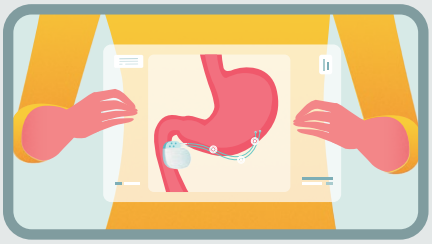
Are you living with gastroparesis? Take back your seat at the table.

IF YOU'RE STRUGGLING TO FIND RELIEF FROM **NAUSEA AND VOMITING CAUSED BY GASTROPARESIS,** IT MAY BE TIME TO CONSIDER A DIFFERENT KIND OF TREATMENT.

Meet Enterra® Therapy

Enterra Therapy is the first and only implantable device designed to help relieve the nausea and vomiting associated with gastroparesis through a unique kind of therapy called Gastric Electrical Stimulation (GES).

A different kind of treatment



MINIMALLY-INVASIVE

The Enterra neurostimulator is placed just beneath the skin, usually in the lower abdominal region.

CUSTOMIZABLE

Your doctor can non-invasively adjust your system to help find the level of stimulation that's right for you.

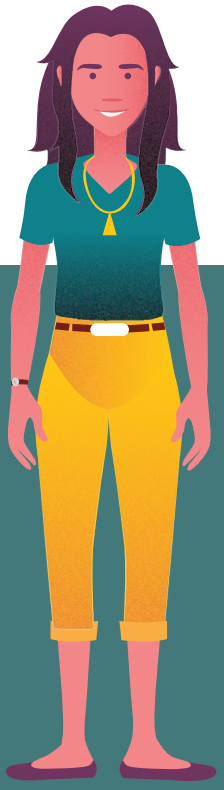
REVERSIBLE

If Enterra Therapy needs to be paused or is not right for you, your doctor can turn off or remove your system.

Is Enterra Therapy right for you?

- ☑ I have gastroparesis due to diabetes or an idiopathic (unknown) cause
- ☑ I experience nausea and/or vomiting that is not helped with medications
- ☑ I am 18-70 years old

If the above statements are true, you may want to talk to your doctor about the risks and probable benefits of Enterra Therapy.



Learn more

Visit www.enterramedical.com/introduction-to-enterra-therapy to watch a video about gastroparesis and Enterra Therapy.

Take the next step

Ask your doctor about Enterra Therapy or visit www.enterramedical.com.

Important Safety Information
Intended Use: The Enterra® Therapy System is an implanted device that provides gastric stimulation to treat chronic, intractable, nausea and vomiting that is not well treated by drugs or other means in patients aged 18 to 70 years caused by diabetes or an unknown origin.

Contraindications: Enterra Therapy is only for patients who are healthy enough for surgical procedures and/or anesthesia. Once implanted, patients need to avoid diathermy, which is deep heat treatment from electromagnetic energy, as it may cause injury or device failure.

Warnings: Enterra Therapy has not been studied in pregnant women, patients under the age of 18, or over 70. Issues may occur if the system interacts with other implanted devices such as pacemakers. Patient injury or device failure may be caused by other medical treatments such as electrocautery, defibrillation/cardioversion, therapeutic ultrasound, or radiofrequency (RF)/microwave ablation. Patient activities may cause shocking or jolting sensations.

MR The Enterra II System is MR Conditional. This means that patients with the Enterra II System can safely have MRI examinations of some body parts under certain conditions. Consult your doctor to determine if you are eligible for MRI examination.

Risks: Potential risks include infection, pain at the surgery site, allergic or immune system response, lead and bowel twist together, device wearing through the skin, bruising, bleeding, loss of therapeutic effect, jolting, shocking, burning sensation, gastrointestinal or stomach issues, loss of therapy due to component failure or battery wear out, or perforated stomach which may cause life-threatening blockage or infections that require immediate medical attention including surgery. Risks can be minimized by avoiding activities such as sudden, excessive, or repetitive bending, twisting, bouncing, or stretching.

Humanitarian Device: Authorized by Federal law for the intended use described above. The effectiveness of this device has not been demonstrated.

Always discuss potential risks and benefits of the device with your physician.
For further information, please contact Enterra Medical at info@enterramedical.com.
Rx Only.

www.enterramedical.com

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