Enterra® Therapy

Patient Ambassador Program

If you are considering Enterra Therapy to manage your nausea and vomiting due to gastroparesis, you may want to hear from others who have received Enterra Therapy.

Enterra Therapy Patient Ambassadors understand gastroparesis can be confusing and lonely, and generously volunteer to share their experiences with Enterra Therapy.

Questions often asked include:

- What are you able to do now that you couldn't before receiving Enterra Therapy?
- Can you feel the stimulation from the therapy?
- How has Enterra Therapy changed your life?

Connect with a Patient Ambassador

To schedule a call with an Enterra Therapy Patient Ambassador scan the QR code or call us at 1-855-204-0455.

You can also visit
enterramedical.com/patient-ambassador-program

Hear from others

Find real stories from people who have received Enterra Therapy.

To watch videos and read testimonials about how Enterra Therapy impacted their lives visit enterramedical.com/hear-from-real-patients

Patient Ambassadors are volunteers and do not receive a financial payment for participating in the Patient Ambassador Program.
Individual results may vary

Learn more about Enterra Therapy at **www.enterramedical.com**

Enterra Medical is dedicated to helping more people with chronic gastroparesis live better lives through advancing technology, bolstering clinical science, and accelerating patient access to Enterra Therapy.

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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