

Has gastroparesis taken your seat at the table?

Watch live and recorded educational series on gastroparesis & treatment options.



Scan the QR code to register for one of the upcoming live sessions or access a previously recorded session.



Register for one of our upcoming live educational discussions on gastroparesis and treatment options where you will:

- ✓ Learn about gastroparesis symptoms causes & diagnosis
- ✓ Understand treatment options, including gastric electrical stimulation
- ✓ Hear from an Enterra® Therapy recipient
- ✓ Find resources to support your gastroparesis journey

Or, watch one of our previously recorded discussions.

Panelists include healthcare providers who specialize in gastroparesis, an Enterra Therapy patient, and an Enterra Medical Patient Liaison.

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.



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

Enterra® is a registered trademark of Enterra Medical, Inc. in the US, EU, and other regions.

©2024 Enterra Medical, Inc. All rights reserved.

MKT-PM-01325, Rev B

