

Intelligent. Responsive. Relief.

- Therapy built on landmark evidence¹
- HFX AdaptivAI™ technology delivers responsive, personalised pain management^{*}
- Real-time app engagement is designed to improve therapy compliance and accelerate relief¹
- Advanced Proactive Remote Monitoring of patient data improves patient care and efficiency



HFX AdaptivAI™ puts patients in control of their pain relief while providing remote patient monitoring to clinicians.

Patient experiences with the HFX iQ™ or Senza Omnia™ spinal cord stimulation (SCS) system vary by individual, including the amount of pain relief. The occurrence of adverse effects associated with SCS implant surgery or use also varies by patient. **Indications for Use:** The HFX iQ, Senza Omnia, and HFX Trial SCS systems aid in the management of chronic intractable pain of the trunk and/or limbs, and, when programmed to 10 kHz, are indicated as aids in the management of chronic intractable pain of the lower limbs associated with diabetic neuropathy and the management of non-surgical refractory back pain. **Contraindications:** These include patients who are poor SCS surgical candidates, are unable to operate the SCS system and fail to receive effective pain relief during trial stimulation. **Warnings:** Interference with other implanted stimulators, may result in sensing problems or inappropriate responses. Sources of electromagnetic interference may result in unexpected changes in stimulation, serious patient injury and system damage. Energy from diathermy can cause tissue damage, resulting in severe injury or death. Senza implantable stimulators are MR conditional and scanning under different conditions may result in severe patient injury or device malfunction. Use of certain medical devices or procedures (electrocautery, radiation therapy, ultrasonic scanning) may result in device damage. Induced electrical currents from radiofrequency (RF) or microwave ablation may cause heating, resulting in tissue damage. **Precautions:** Avoid activities that put stress on the implanted components. Safety has not been established for Transcranial Magnetic Stimulation (TMS) or Electroconvulsive Therapy (ECT) in patients who have an implanted SCS system. Persistent discomfort or excessive redness may indicate infection. **Adverse Events:** May include hematoma, epidural hemorrhage, paralysis, pain at implant site, infection and other surgical risks. Device related adverse events may include loss of pain relief or paraesthesia, undesirable change in stimulation (uncomfortable, jolting or shocking sensation), tissue reaction or allergy to implanted materials. Refer to www.nevro.com/manuals for product manuals with complete indications, contraindications, warnings, precautions and potential adverse events.

^{*}HFX AdaptivAI™ is a set of additional features of iQ mode. It includes an advanced algorithm with an ability to interlace bipoles, reduce step size and pulse dosing. HFX iQ™ uses predefined rules and organised data to provide optimised treatment recommendations. It utilises direct patient input from assessments on pain and quality of life measures and machine-based inputs such as lead placement, electrodes, therapy utilisation and pain regions. HFX iQ is not a machine learning-based system. Big data is referring to Nevro's internal cloud database. [†]Customised programs are pre-defined parameters programmed by the physician. [‡]Calls made during business hours. [^]Coming in 2026.

1. Data on file. **2.** Kapural, L., et al. (2016). Comparison of 10-kHz high-frequency and traditional low-frequency spinal cord stimulation for the treatment of chronic back and leg pain: 24-month results from a multicenter, randomized, controlled pivotal trial. *Neurosurgery*, 79(5), 667-677. **3.** Patel, N.P., et al. (2023). Durable responses at 24 months with high-frequency spinal cord stimulation for nonsurgical refractory back pain. *J Neurosurg Spine*:1-11. **4.** Petersen, E., et al. (2023). Long-Term Efficacy of High-Frequency (10 kHz) Spinal Cord Stimulation for the Treatment of Painful Diabetic Neuropathy: 24-Month Results of a Randomized Controlled Trial. *Diabetes Research and Clinical Practice*, 110865. **5.** Data on file. Patients using the HFX App. **6.** Data on file. 3% PD 10 kHz 30 us 2.0 mA 20 secs 10mins 800 Ohms.

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Intelligent. Responsive.

Welcome to the relentless pursuit of **optimal relief**



The only SCS system with HFX AdaptivAI™ that combines Level 1 evidence and big data insights to deliver responsive relief for the ever-changing patient journey*

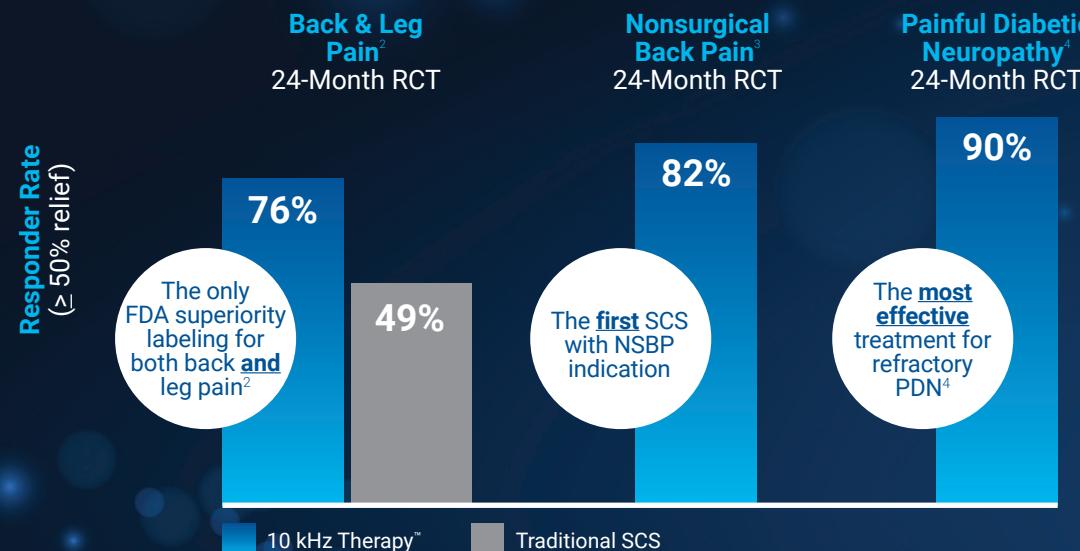
Now with Advanced Proactive Remote Monitoring[^]



HFX AdaptivAI™
powering **HFX iQ™**

Welcome to the relentless pursuit of optimal relief

Therapy built on landmark evidence¹



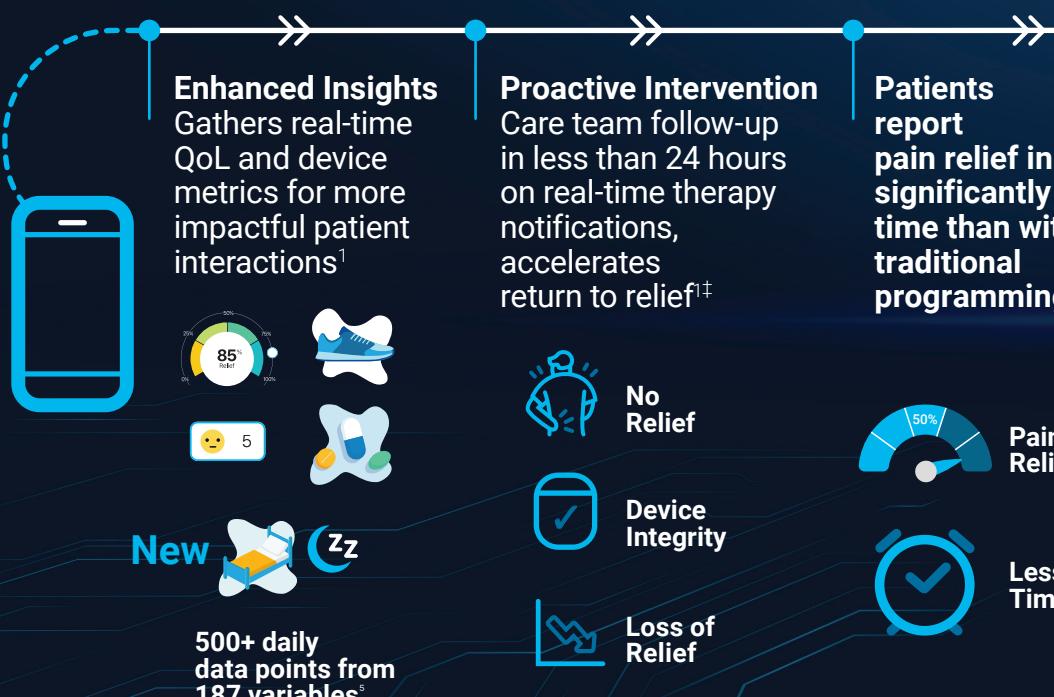
Responsive, personalised pain management via HFX AdaptivAI™ technology*

Starts patients with optimal settings based on their pain type, designed to get them to **relief faster**¹

Relief and Beyond
Engineered to maximise relief over time - even when relief over 50% has been achieved¹

Smart Power
With therapy optimisation, patients may only charge **6x per year**⁶

Real-time app engagement is designed to improve therapy compliance and accelerate relief¹



HFX AdaptivAI™ technology is built on indication-specific algorithms, enhanced by big data and proven outcomes*



Now with Advanced Proactive Remote Monitoring¹
Provides direct physician access to remotely monitor every patient's pain journey over time