Ready To Find Personalized Pain Relief?For appropriate guidance, consult the instructions for use for these active devices.NM-787914-AA Indications for Use: The Boston Scientific Spinal Cord Stimulator Systems are indicated as an aid in the management of chronic intractable pain of the trunk and/or limbs including unilateral or bilateral pain associated with the following: failed back surgery syndrome, Complex Regional Pain Syndrome (CRPS) Types I and II, intractable low back pain and leg pain, Diabetic Peripheral Neuropathy of the lower extremities, radicular pain syndrome, radiculopathies resulting in pain secondary to failed back syndrome or herniated disc, epidural fibrosis, degenerative disc disease (herniated disc pain refractory to conservative and surgical interventions), arachnoiditis, multiple back surgeries.Indications for Use: The Superion(TM) Indirect Decompression System (IDS) is indicated to treat skeletally mature patients suffering from pain, numbness, and/or cramping in the legs (neurogenic intermittent claudication) secondary to a diagnosis of moderate degenerative lumbar spinal stenosis, with or without Grade 1 spondylolisthesis, having radiographic evidence of thickened ligamentum flavum, narrowed lateral recess, and/or central canal or foraminal narrowing.The Superion Indirect Decompression System (IDS) is contraindicated for patients who: have spinal anatomy that prevent implantation of the device or cause the device to be unstable in situ (i.e., degenerative spondylolisthesis greater than grade 1), Cauda equina syndrome, or prior decompression or fusion at the index level, scoliosis or spinous process fractures, osteoporosis, infection, allergy or reaction to any metal or implant or a high Body Mass Index.Patients implanted with the Precision Spectra(TM) or Spectra WaveWriter(TM) Spinal Cord Stimulator Systems with ImageReady(TM) MRI Technology are "MR Conditional" only when exposed to the MRI environment under the specific conditions defined in the applicable ImageReady(TM) MRI Head Only Guidelines for Precision Spectra(TM) or Spectra WaveWriter(TM) Spinal Cord Stimulator Systems. MR Conditional symbolBoston Scientific's ImageReady(TM) MRI Full Body Technology makes safe MRI scans possible. The Precision Montage(TM) MRI, WaveWriter Alpha(TM) and WaveWriter Alpha(TM) Prime SCS Systems with ImageReady(TM) MRI Full Body Technology are "MR Conditional" only when exposed to the MRI environment under the specific conditions defined in the applicable ImageReady(TM) MRI Full Body Guidelines for Precision Montage(TM) MRI or WaveWriter Alpha(TM) and WaveWriter Alpha(TM) Prime Spinal Cord Stimulator Systems.Indications for Use: The Boston Scientific Radiofrequency Generators, associated Radiofrequency Lesion Probes and RF Cannula are indicated for use in procedures to create radiofrequency lesions for the treatment of pain or for lesioning nerve tissue for functional neurosurgical procedures.The Boston Scientific Spectra WaveWriter(TM), WaveWriter Alpha(TM) and WaveWriter Alpha(TM) Prime SCS Systems are also indicated as an aid in the management of chronic intractable unilateral or bilateral low back and leg pain without prior back surgery.Refer to the Instructions for Use provided on www.vertiflex.com for additional Indications for Use, contraindications information and potential adverse effects, warnings, and precautions prior to using this product.For therapy that does not generate paresthesia (i.e. subperception therapy) it is less likely that sudden stimulation changes resulting in distraction could occur while having stimulation on when operating moving vehicles, machinery, and equipment.MR Conditional symbol Boston Scientific's ImageReady(TM) MRI Technology makes safe MRI head scans possible.