

Fundamentals of Neuromodulation

Peter Pryzbylkowski, MD

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Disclosure

Consultant: Nevro, Camber Spine, Vertos

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Learning Objectives

Explore the use of electrical signals to block pain

Review the theory of how neuromodulation works, MOA

• Explore the different products that are currently on the market

• Review the application of the devices in clinical practice

 $\hfill \ensuremath{\cdot}$ Review data supporting use of products and their role in decreasing opioid use

Discuss the process of trial and implantation of devices

History of Neuromodulation

- First used to treat pain in 1967
- Gate theory was published in 1965
- Became more mainstream in 1980s
- 1989 FDA approved use of devices to treat chronic pain from nerve damage in trunk, arms or legs
- Year after year the devices continue to improve upon earlier iterations

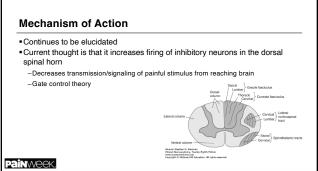
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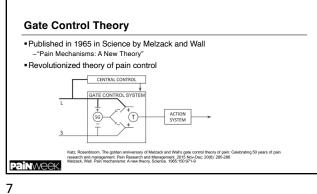
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Neuromodulation

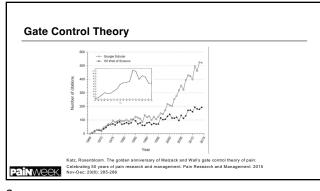
- So what is it?
- Application of electrical signals to lessen pain complaints
- Drug/medication = electricity
- Types of neuromodulators
- -Spinal cord stimulators, dorsal column stimulators, dorsal root ganglion stimulators, peripheral nerve stimulators

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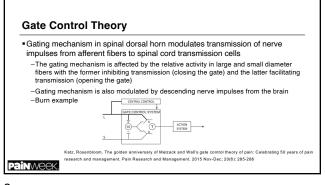














Neuromodulation and the Gate Control Theory



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Neuromodulation

FDA approved

-Alleviation of pain in trunk, arms or legs -Chronic regional pain syndrome • AKA RSD or causalgia

Most common indication/usage

Pacemaker companies

-Failed back surgery syndrome • Post laminectomy pain syndrome Chronic pain syndrome

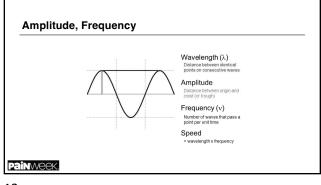


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Amplitude, Frequency, Pulse Width

- Parameters we can change with SCS devices
 -Frequency is how often device delivers charge and depolarization
- -Amplitude is relative strength of charge delivered -Pulse width is duration of charge delivery
- Tonic or low frequency
- -20-120Hz range
- -patients perceive individual pulses
- High frequency -pulses start to blend so no perception occurs



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Traditional vs High Frequency vs DRG

Traditional AKA "low frequency," "tonic"
 Tens unit sensation, paresthesia present, can go up to 1200Hz

• High frequency, paresthesia not present, 10,000Hz

DRG (dorsal root ganglion) stimulators
 Low frequency, used for focal pain locations

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Electrodes

• Typically 8 electrodes per lead with two leads typically used • Surgeons can place paddle leads with different configurations DRG 4 electrodes



Evidence for Neuromodulation

Kumar study

- RCT conventional medical management (CMM) vs SCS for neuropathic pain -Primary outcome was patients reporting 50% or more relief of leg pain
- -Secondary outcomes were improvement in back pain, QOL, functional capacity, use of medication, patient satisfaction Compared with CMM group the SCS group saw
- -Improved back and leg pain, better QOL, greater treatment satisfaction

Kumar, Taylor, Jacques et al. Spinal cord stimulation versus conventional medical management for controlled trial in patients with failed back surgery syndrome. Pain. 2007 Nov;132(1-2): 179-88 athic pain: a multic

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Evidence for Neuromodulation

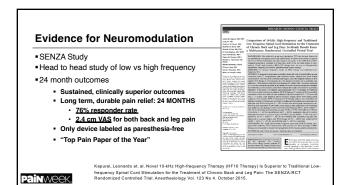
Deer study

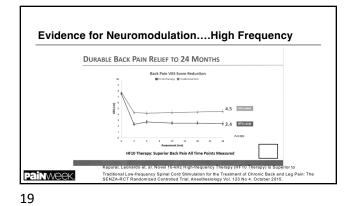
- Multicenter, prospective RCT
- -Following successful trial 100 patients were randomized after implant to receive 12 weeks of tonic stim followed by 12 weeks of burst
- -Primary endpoint assessed the noninferiorty of the within-subject difference between tonic and burst for mean daily VAS score

Burst stimulation is non inferior to tonic stim
 Significantly more subjects 70.8% preferred burst over tonic; preference was sustained over 1 year

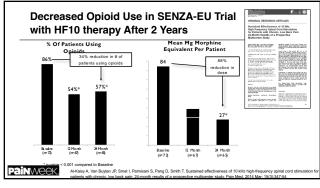
Deer, Slavin, Amirdelfan et al. Success using neuromodulation with burst (sunburst) study: results from a prospective randomized controlled trial using a novel burst waveform. Neuromodulation. 2018 Jan:21(1):56-66

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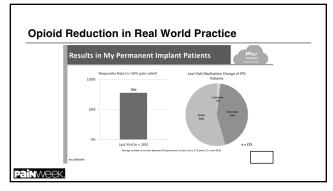




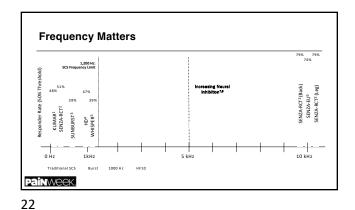














Procedure Overview (Trial)

- Only pain procedure that requires psychiatric/psychological clearance by insurance company -Patient is malingering or faking symptoms

 - -Patient will call if there is infection or issues with device -Most of these patients have undergone previous spine surgery
 - -Large scar present on back





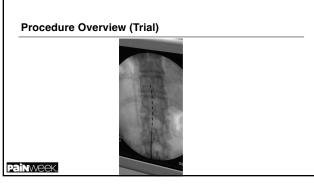
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Procedure Overview (Trial)

- Placement of percutaneous electrodes into epidural space
- Just like performing an epidural. Done under xray
- -Rather than injecting medication electrodes are placed
- -Trial leads stay in place for 5-7days and are connected to a battery $\,\cdot$ If >50% pain reduction Implant can be performed



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Procedure Overview (Permanent Implant)

Leads are again placed into epidural space and then tunneled under skin to a battery
 -Battery, which powers the device, is placed in the flank

Battery
 -Rechargeable vs Non-Rechargeable



Complications/Risks of Procedure

 Infection
 -Epidural abscess
 ·Paralysis Bleeding -Epidural hematoma • Paralysis Lead migration/lead fracture -Loss of efficacy

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Contraindications

- Severe uncontrolled psychological disorders
 -Schizophrenia, depression, bipolar disorder
- Bleeding disorder
 Use of blood thinners or NSAIDs
- Active infection
- Relative contraindication
- -Need for continued MRI studies Most newer devices have MRI approval

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