



Emerging-Solutions (ES) Percutaneous Electrical Nerve Stimulator (PENS)

- FDA cleared Class II Medical Device
- Offering relief from chronic pain and opioid withdrawal
- Covered by most insurance plans
- Results lasting up to 6 months

Contact

Emerging Solutions
15455 Dallas Parkway
Suite 600
Addison, TX 75001
United States
972-746-2722



Percutaneous Electrical Nerve
Stimulator
(PENS)



PENS Device

- PENS is a Programmable Class II Medical Device.
- FDA cleared in 2018 for the treatment of opioid withdrawal, typically associated with individuals suffering from chronic pain.
- Four strategically positioned, 2mm titanium needle electrode arrays implanted directly into the nerve endings on the ear based on the patient's diagnosis.
- Programing Technical Unit (PTU) generates an ear map allowing the provider to identify the appropriate location for implantation of the titanium needle electrode arrays, utilized by the Nerve Locator.
- Temporary, PENS device worn for 10-12 days with potential results offering up to 6 months of pain relief.



Our Services

- Predetermination/Preauthorization and billing process is facilitated by ES to mitigate denials by Commercial/Medicare insurance.
- Dedicated Billing Manager assigned to every facility.
- On site, one on one training provided by ES team.
- No up-front costs to the medical provider/medical facility.



Billing

In-office, surgical procedure; four titanium needle electrode arrays implanted under the skin

Must be performed by a MD, DO, PA, NP or CRNA. Specific states allow implantation to be performed by MD or DO.

Covered by Medicare, Workers Comp, PI and most commercial plans *NOT COVERED by MEDICAID.

Predetermination/preauthorization required by commercial insurance.

ES Portal updates the status of each preauthorization/predetermination and claim,

Improves Patient Care

Successful in mitigating pain for up to 6 months.

Decreases desire for pain medication; potentially aiding in opioid withdrawal treatment.

Improves sleep patterns as withdrawal systems are simultaneously reduced as a result of the device.

Potential to reduce anxiety and depression as pain is mitigated.

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Percutaneous Electrical
Nerve Stimulator
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Patients not eligible for the PENS device:

- Pacemaker
- Hemophilia
- Pregnancy
- Heart Monitor
- Deep Brain Stimulator
- Permanent Spinal Cord Stimulator – SCS.

Advantages of the Program

FDA cleared Class II Medical Device

Offering relief from chronic pain and opioid withdrawal

Covered by most insurance plans, including commercial, Medicare and Workers Compensation * Medicaid not covered

Pain mitigation is consistent with 70% of patients seeing significant reduction in pain with results lasting up to 2- 6 months

Patient Testimonials

- “I’ve felt happier, with zero anxiety, and on top of the world. It really helped my pain. I would recommend this to everyone that deals with chronic pain and any anxiety or depression.”
- “I would recommend the Neurostimulator for anyone with chronic pain, anyone who has trouble sleeping, high anxiety or wants to enjoy life the way it used to be before pain.”

