

Nerve stimulation reduces pain and opioid use after orthopedic surgery

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A technique called percutaneous peripheral nerve stimulation yields "impressive" reductions in pain scores and opioid use during the first week after common orthopedic surgery procedures, concludes a



randomized clinical trial published Online First in *Anesthesiology*, the official peer-reviewed journal of the American Society of Anesthesiologists (ASA), today.

The benefits of postoperative <u>nerve</u> stimulation were "much greater than what we had anticipated, concurrently reducing <u>pain scores</u> by more than 50 percent and opioid consumption by 80 percent," according to the randomized trial report by Brian M. Ilfeld, M.D., MS, and colleagues. With further study, they believe that peripheral nerve stimulation could be a safe, effective, opioid reducing, non-drug adjunct for <u>pain</u> <u>management</u> after surgery.

In percutaneous peripheral nerve stimulation, a tiny electrical lead (insulated wire) is placed alongside a nerve and connected to a batterypowered pulse generator. The generator delivers mild electrical stimulation to the nerve, interrupting pain transmission. Although this 'neuromodulation' technique has been widely used for patients with <u>chronic pain</u>, the new study is the first randomized, controlled trial to evaluate its use for acute pain after surgery.

The study enrolled 65 patients undergoing common outpatient joint surgery, such as bunion surgery in the foot or rotator cuff repair in the shoulder. All patients underwent lead placement near the nerve(s) serving the surgical joint. Half of the patients were randomly assigned to active electrical stimulation, adjusted to achieve the desired sensory change—sometimes described as feeling like a "pleasant massage." The remaining patients received an inactive "sham" treatment, with a pulse generator that appeared to function normally, but did not deliver any electrical current. After one week, pain scores and <u>opioid use</u> were compared between groups. Evaluations were performed in "doublemasked" fashion: neither the researchers nor the patients knew which treatment the patient was receiving.



The results showed significantly lower pain ratings in the active nerve stimulation group. Mean pain score (on a 0 to 10 scale) was 1.1 in patients receiving active treatment, compared to 3.1 in the sham group.

Active nerve stimulation was also associated with a much lower use of opioids to control postoperative pain. Median opioid dose (oral morphine equivalents) for the entire first week after surgery was 5 milligrams in the active treatment group, compared to 48 milligrams in the sham group. That's an important benefit due to the possible adverse effects of postoperative opioids, such as nausea, constipation, sedation, and respiratory depression.

Although the trial was designed as a <u>pilot study</u>, the results "stand on their own and indicate that percutaneous peripheral nerve stimulation is highly effective for acute pain," Dr. Ilfeld and coauthors wrote. They outlined plans for a subsequent study that is currently enrolling a far larger number of patients and will include a more complete assessment of the treatment effect across patient subgroups.

More information: Brian M. Ilfeld et al. Percutaneous Peripheral Nerve Stimulation (Neuromodulation) for Postoperative Pain: A Randomized, Sham-controlled Pilot Study. *Anesthesiology* April 2021. doi: <u>doi.org/10.1097/ALN.00000000003776</u>

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