

syndrome, may be treated successfully with physical therapy and trigger point injections. If symptoms persist, more aggressive treatment, such as serial sympathetic blockade, is indicated to avoid a chronic, irreversible pain syndrome.

This case demonstrates that relapse may occur and that symptoms of reflex sympathetic dystrophy may spread diffusely. Spread of reflex sympathetic dystrophy to a bilateral or ipsilateral extremity is common, but our patient represents an apparently unique case of total-body involvement.

#### REFERENCES

1. Bonica JJ: Causalgia and other reflex sympathetic dystrophies. *Postgrad Med* 53:143-148, 1973
2. Kiaer S: Remarks on the prognosis of the post-traumatic dystrophy of the extremities. *Acta Orthop Scand* 17:253-269, 1947
3. Pak TJ, Martin GM, Magness JM, et al: Reflex sympathetic dystrophy: Review of 140 cases. *Minn Med* 53:507-512, 1970
4. Kozin F, McCarty DJ, Sims J, et al: The reflex sympathetic dystrophy syndrome. I. Clinical and histologic studies: Evidence for bilaterality, response to corticosteroids and articular involvement. *Am J Med* 60:321-331, 1976
5. Kozin F, Genant JK, Bekerman D, et al: The reflex sympathetic dystrophy syndrome. II. Roentgenographic and scintigraphic evidence of bilaterality and of periarticular accentuation. *Am J Med* 60:332-338, 1976
6. Ochs S: Systems of material transport in nerve fibers (axoplasmic transport) related to nerve function and trophic control. *Ann NY Acad Sci* 228:202-223, 1974
7. Doupe J, Cunnen CH, Chance GQ: Post-traumatic pain and the causalgic syndrome. *J Neurol Neurosurg Psychiatr* 7:33-48, 1944
8. Livingston WK: Pain Mechanism: A Physiologic Interpretation of Causalgia and Its Related States. New York, Macmillan, 1943, pp 224-235
9. Melzack R, Loeser JD: Phantom body pain in paraplegics: Evidence for a central "pattern generating mechanism" for pain. *Pain* 4:195-210, 1978
10. Melzack R, Dennis SG: Pain mechanisms: Theoretical approaches, Mechanisms of Pain and Analgesic Compounds. Edited by RF Beers Jr, EG Bassett. New York, Raven Press, 1979, pp 190-191
11. Sternschein MJ, Meyers SJ, Frewin DB, et al: Causalgia. *Arch Phys Med Rehabil* 56:58-63, 1975
12. Editorial: Algodystrophy. *Br Med J* 1:461-462, 1978

Anesthesiology  
53:257-258, 1980

## The Effective Use of Epidural Morphine Sulfate for Postoperative Orthopedic Pain

JOHN EBERT, D.O.,\* AND PAMELA DUNCAN VARNER, M.D.†

Postoperative pain following arthrotomy of the knee is usually intense. Patient cooperation with physical therapy after this procedure is usually less than optimal when standard methods of producing analgesia, such as parenteral administration of narcotics, are used. We report the successful use of morphine, given epidurally, for analgesia following a knee arthrotomy. The medical management of this case took place in the Jefferson Tower of the University Hospitals in Birmingham, Alabama.

#### REPORT OF A CASE

A 29-year-old Caucasian woman was admitted for an arthrotomy and repair of a traumatic disruption of the right knee. Medical, surgical, and previous anesthetic histories prior to this injury were unremarkable. Her only medication was norethindrone acetate,

for treatment of pelvic endometriosis. Premedication one hour before the surgical procedure consisted of diazepam, 10 mg, orally, and 30 ml of an antacid containing aluminum hydroxide, magnesium hydroxide, and simethicone. The patient had not received narcotics for more than 27 hours prior to operation.

The surgical procedure was conducted using continuous lumbar epidural analgesia with .75 per cent bupivacaine with 1:200,000 epinephrine, 20 ml. The sensory level of anesthesia was T9 bilaterally. During the first 24 hours postoperatively pain relief was achieved with 0.25 per cent bupivacaine with 1:200,000 epinephrine, 5 ml given epidurally every two hours to maintain adequate analgesia. Narcotics were not needed during this period.

Twenty-four hours after the operation the analgesia induced by bupivacaine was allowed to dissipate, and intense knee pain occurred. Morphine sulfate crystals, 10 mg, were dissolved in 10 ml of 0.9 per cent NaCl and passed through a Millipore® filter as specified by the hospital pharmacy.\* Five milliliters of this preparation (morphine sulfate, 5 mg) were then administered through the epidural catheter following a negative aspiration for blood. Pain gradually decreased over the first hour, but was not completely relieved by two hours. A second dose of morphine sulfate, 4 mg (due to the loss of 1 ml, or 1 mg, in the Millipore filter), was then administered epidurally, after which the catheter was removed. Analgesia was complete an hour after the second dose,

\* Assistant Professor.

† Resident.

Received from the Department of Anesthesiology, University of Alabama in Birmingham, School of Medicine, Birmingham, Alabama 35294. Accepted for publication May 1, 1980.

Address reprint requests to Dr. Ebert.

Key words: Analgesics, narcotic: morphine. Anesthetic techniques, epidural. Surgery, orthopedic.

\* The use of epidural morphine has been approved by the Human Use Committee of the University of Alabama in Birmingham. Informed consent was obtained from the patient.

and lasted 40 hours. There was no sign of central nervous system depression or sympathetic or motor block during this time. Administration of additional narcotics or non-narcotic analgesics was not necessary during the hospitalization; sleep was uninterrupted by pain, and aggressive physical therapy could be tolerated without discomfort. On the third postoperative day, the day of discharge, the pain gradually reappeared, and orally administered narcotic analgesics were needed every four hours for pain relief. The narcotic requirement was tapered, and narcotic administration was discontinued on the seventh postoperative day.

#### DISCUSSION

The autoradiographic localization of opiate receptors<sup>1,2</sup> in both the mesencephalic central gray matter of the brain and the substantia gelatinosa of the posterior horn cells of the spinal cord has led to important research in the area of analgesia with narcotics. Analgesia has been produced by intracerebroventricular injection of morphine in rats and mice.<sup>3</sup> Morphine also affects<sup>5</sup> opiate receptors in the spinal cord, resulting in modulation of afferent nociceptive information,<sup>4-6</sup> without causing adverse reactions in cord tissue.<sup>7</sup> Spinal serotonin and norepinephrine terminals may mediate this spinal antinociceptive effect of morphine.<sup>8</sup>

The use of commercially available narcotic preparations in the epidural space has been criticized because these solutions contain preservatives and stabilizers that are presumed toxic to nerve tissue, and may also have anesthetic properties of their own.<sup>9</sup> Morphine sulfate in its liquid form is unstable, and will lose considerable potency over a short period (<eight hours). For these reasons, morphine sulfate in its crystallized form was freshly reconstituted and passed through a Millipore filter immediately prior to epidural injection.

The efficacious use of morphine administered in the epidural<sup>10</sup> or subarachnoid<sup>11</sup> space in the treatment of chronic and intractable pain in man has been well described. The practical use of epidural narcotics in labor has been less encouraging.<sup>12</sup> The beneficial effect of epidurally administered morphine in the treatment of postoperative orthopedic pain has not been described. In this case, the epidural administration of morphine sulfate in a total dosage of 9 mg provided complete pain relief for 40 hours and permitted pain-free rehabilitation therapy that is usually

not well tolerated. Although the systemic absorption of the epidurally administered morphine may have accounted for the analgesia, we feel it is unlikely that 9 mg of morphine sulfate could have provided 40 hours of relief of an intensely painful condition such as this. The lack of central nervous system depression with simultaneous relief of acute, intense postoperative pain is additional evidence that this represented a spinal antinociceptive effect of morphine. The absence of sympathetic and motor block with this technique is also considered beneficial. Physical therapy may proceed uninhibited by pain or decreased muscular tone, while the risks of venous pooling during ambulation are reduced. It is hoped that this brief case report will stimulate investigators to conduct controlled and clinical studies in an effort to establish the efficacy of epidurally administered narcotics in the relief of postoperative pain.

#### REFERENCES

1. Pert CB, Kuhar JM, Snyder SH: Opiate receptor: Autoradiographic localization in rat brain. *Proc Natl Acad Sci USA* 73:3729-3733, 1976
2. Snyder SH: Opiate receptors and internal opiates. *Nature* 257:185-189, 1975
3. Sewell RD, Spencer PS: Antinociceptive activity of narcotic agonist and partial agonist analgesics and other agents in the tail-immersion test in mice and rats. *Neuropharmacology* 15:683-688, 1976
4. Yaksh TL, Rudy TA: Studies on the direct spinal action of narcotics in the production of analgesia in the rat. *J Pharmacol Exp Ther* 202:411-428, 1977
5. Yaksh TL, Frederickson RCA, Hwang SP, et al: In vivo comparison of the receptor populations acted upon in the spinal cord by morphine and pentapeptides in the production of analgesia. *Brain Res* 148:516-520, 1978
6. Yaksh TL: Analgetic actions of intrathecal opiates in cat and primate. *Brain Res* 153:205-210, 1978
7. Wang JK: Analgesic effect of intrathecally administered morphine. *Regional Anesth* 2:3, 1977
8. Yaksh TL: Direct evidence that spinal serotonin and noradrenalin terminals mediate the spinal antinociceptive effects of morphine in the periaqueductal gray. *Brain Res* 160:180-185, 1979
9. Mathews E: Epidural morphine (letter). *Lancet* 1:673, 1979
10. Behar M, Olshwang D, Magora F, et al: Epidural morphine in treatment of pain. *Lancet* 1:527-528, 1979
11. Wang JF, Nauss LA, Thomas JE: Pain relief by intrathecally applied morphine in man. *ANESTHESIOLOGY* 50:149-151, 1979
12. Periss BW: Epidural opiates in labour (letter). *Lancet* 2:422, 1979