Use of an intravenous port catheter for the delivery of intrathecal morphine in a terminally ill cancer patient with pain

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ABSTRACT

We present a case of a terminally ill cancer patient with metastasis who was suffering from inadequate pain control despite high doses of systemic opioid administration. In addition, her quality of life was significantly impaired due to opioid-induced troublesome side effects. In the situation as above, intrathecal opioid administration has established as a safe and effective therapeutic option with a positive benefit–risk ratio. This case reports a novel technique, where an intravenous port catheter was used to administer intrathecal opioid.

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Many terminally ill patients with malignant pain, despite high doses of systemic opioids suffer from agony. Advanced analgesic strategy such as intrathecal drug delivery (ITDD) system is employed to deliver opioid directly to the spinal cord.1 This technique provides superior pain control, hence renders both, good quality of life and death to the patient. In a situation when ITDD system not available, alternate methods are used to deliver a drug to the CSF.

Case Report. Patient information and clinical examination. A 39-year-old terminally ill lady with metastatic breast cancer presented with moderately severe upper abdominal pain. The pain started about 4 months prior due to metastasis in her liver and spleen. Initially, the pain was well controlled with a transdermal fentanyl patch (TFP). Later, despite a high dose of fentanyl (TFP 200 µg per hour, superadded with frequent sublingual 100 µg fentanyl doses), her pain was not well controlled. In addition, she was having mild drowsiness fluctuating to almost a comatose state, moderately severe itching, nausea, and constipation. She also gave a history of orthopnea and felt less distress when sitting and in a left lateral position. She could walk a few steps with support and was continent of urine and stool. According to her oncologist, her life expectancy was less than 2 months. Upon a physical examination, she was 56 Kg, looked pale, cachectic, and distressed. She was well educated and orientated and expressed her complaints in broken sentences. Her abdomen was distended due to massive hepato-splenomegaly.

Diagnostic evaluation Blood hematology and biochemistry were within normal limits. Although the liver function tests were deranged, the prothrombin time was normal. A radiological investigation revealed massive hepato-splenomegaly and metastasis in the long bones but chest was clear.

Intervention for the relief of pain. She signed the informed consent for the insertion of a port catheter as
Intravenous as intrathecal port catheter … Anwari & Romdhane

A route for the administration of intrathecal morphine. With a fasting period of 6 hours for solid food, she was brought into the operating theatre and placed on an operating table in the left lateral position with 2 pillows under her head. Routine monitoring (such as ECG, BP, and SpO₂) was commenced, and intravenous access on the arm was secured, through which a prophylactic dose of ceftriaxone (1.0 G) was given. Both operators (JSA & KR) and the assisting nurse took strict aseptic precautions. A wide area of skin was double cleaned with 2% chlorhexidine and 70% alcohol, which was isolated with sterile drapes. The procedure was done under local anesthesia (lignocaine 1.0% with adrenaline 1:200,000). Sterile procedural instruments were place on the trolley, including a BardPort™, an implantable port with a 6.6 F catheter (BARD access system Inc. Utah USA) (Figure 1) and an epidural minipack of Portex® (Smith medical int. USA). The procedure was carried out under fluoroscopic guidance. An epidural Tuohy needle (18.G) was inserted between the L2 and L3 inter-spinous space and advanced, using the loss of resistance to air technique till reaching the epidural space. The Tuohy needle was further advanced for a few millimeters till a free flow of CSF appeared. The skin to subarachnoid space distance was noted. The Seldinger guide wire (GW) from the BardPort was inserted through the tapered end of a “J” straightener and through the Tuohy needle.

Before pushing the GW further to appear through the bevel opening, the patient was warned about current-like sensations in her lower limbs. In the fluoroscopic view, the GW appeared through the needle, which was further pushed about 4 cms in the subarachnoid space. The patient felt the paresthesia, but did not require any analgesic. Keeping the GW in place, the Tuohy needle was removed. Using a surgical scalpel blade No. 11, a stab incision was made from the exit point of the GW, 1.5 cms deep to the skin. Then, the operator took the BardPort™ vessel dilator and sheath as a unit, and a length equal to the previously measured skin–subarachnoid distance was marked with sterile ink. This unit was then advanced up to the mark over the exposed GW. While the sheath was held in place, the dilator and GW were removed together. The CSF appeared through the proximal end of the sheath. The BardPort™ catheter (6.6F) was inserted into the sheath and advanced 4 cms in the subarachnoid space. When the CSF appeared from the proximal end of the catheter and its tip position was fluoroscopically confirmed, the sheath was then peeled away from the catheter. A subcutaneous pocket for the port was created below the right costal margin over the anterior axillary line. Through the stab incision made earlier (at the catheter entry site), the tip of the tunneler was advanced to the port pocket site. The free end of the catheter was then attached to the tunneler barb, and the tunneler was completely pulled through to the pocket port and then detached from the catheter. The air from the port was removed by flushing with saline and placed in a subcutaneous pocket. The excess length of the catheter was cut and connected with the port. The port was secured with stitches, and the skin was sutured to close the port pocket. One ml of radio-opaque contrast (Iohexol 350) was injected trans dermally into the port, which confirmed its spread into the CSF (Figure 2). We used 2.0 ml of preservative-free (PF) morphine (0.5mg/
ml) mixed with 1.0 ml (50 µg) of fentanyl as the first intrathecal dose. The patient’s pain improved from 8/10 to 2/10 in less than 15 minutes.

Follow up and outcome. Subsequently, the patient received once daily intrathecal PF morphine (2.0 ml, or 1.0 mg) without fentanyl. She was allowed to take a fentanyl sublingual tablet (100 µg) for breakthrough pain. During subsequent days, her pain was well controlled with minimal side effects with the once a day intrathecal morphine. Throughout, she remained apyrexial with normal white cell count. On the 18th day after the insertion of the intrathecal BardPort™, she peacefully expired in the presence of her relative.

Discussion. Breast cancer is a common malignancy among Saudi females, with a prevalence of 21.8% and around 930 new cases of breast cancer diagnosed each year. Cancer pain is a global health issue. A survey conducted on adult cancer patients in an oncology unit of a teaching hospital in Riyadh found that 50% of patients reported moderate to severe cancer pain. The standard of care for cancer patients is set by the Joint Commission on Accreditation of Healthcare Organization, which states, “No cancer patient should live or die with unrelieved pain.” The expectations of terminal illness, their relatives, and care providers are akin to such a standard. A systemic review exploring the component of “good death” looked into 36 studies and revealed that all the stakeholders considered pain-free status a top priority. Eighty percent of patients with advanced cancer pain who received oral or transdermal opioid achieved adequate pain control. However, for the remaining 20% of patients, interventional treatment was required.

Although our patient reached the third step of the WHO analgesic ladder, her condition was miserable. Her pain, despite the high dose of transdermal fentanyl, was not well controlled. In addition, she suffered from many troublesome opioid side effects. Therefore, we decided to offer her the “fourth step.” One option was to perform an ablative procedure, such as a celiac plexus or a splanchnic nerve neurolysis. These procedures are usually carried out under fluoroscopic guidance and require the patient to be in a prone position. In our case, it was not possible for her to lie in the required position.

The second option was to directly administer the opioid into the spinal cord. In 1973, Tony Yaksh discovered that the opioid receptors in the spinal cord provided the scientific rationale for this form of treatment. A few years later, Wang et al. first reported the successful use of intrathecal morphine for cancer pain. Henceforth we opt for administering spinal opioid through an intrathecal route. As compared to the epidural, the intrathecal administration route offers lower doses and lower risks of catheter migration and infection. Moreover, as compared to a percutaneous intrathecal catheter, the risk of meningitis is further reduced with a totally implanted catheter with a subcutaneous injection port.

Such devices are commercially available (e.g., Codman 3000. Codman and Shurtleff, Inc., Raynham, MA, USA), but are not available at our institution. One of us (KR) has had good experience in using an implantable port pediatric intravenous access catheter (BardPort™. Bard Access Systems, Inc. UT, USA) as an intrathecal catheter for administering opioid in terminally ill cancer patients with pain. We decided to use this option. We searched Google and PubMed and could not find any report such as ours. Her pain was adequately controlled, and the fentanyl-induced side effects were significantly reduced with a daily administration of intrathecal morphine (1.0 mg). One week later, the morphine dose was increased to 1.5 mg daily. Throughout the period after the insertion of the intrathecal port catheter, she (and her family) acknowledged good analgesia, maintained good communication, and remained apyrexial, and the opioid-induced side effects were minimal.

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Describe statistical methods with enough detail to enable a knowledgeable reader with access to the original data to verify the reported results. When possible, quantify findings and present them with appropriate indicators of measurement error or uncertainty (such as confidence intervals). Avoid relying solely on statistical hypothesis testing, such as the use of *P* values, which fails to convey important information about effect size. References for the design of the study and statistical methods should be to standard works when possible (with pages stated). Define statistical terms, abbreviations, and most symbols. Specify the computer software used.