Possible Pulmonary Gas Embolism Associated with Localized Thermal Therapy of the Liver

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EXTERNAL heating by conventional electric cautery or by laser provides coagulation of the tissue by conducting the heat from the surface to the inside of the tissue. Conversely, internal heating is the method for necrotizing tumor and hemostasis by means of a dielectric heating electrode, which is inserted into the tissue or tumor. The two methods of administering localized internal thermal therapy of the liver are microwave coagulation therapy (MCT)1 and radiofrequency ablation (RFA).2,3

A venous air embolism, a potential complication of surgery in the sitting position4 or of laparoscopy,5,6 is unlikely to occur during laparotomy in a horizontal position. However, we experienced a case with abrupt decreases in end-tidal carbon dioxide partial pressure (PETCO2) and hemoglobin oxygen saturation (SpO2) during laparotomic MCT in the left lateral recumbent position, probably due to pulmonary gas emboli. We believe that the case discussed below is the first detailed report of a pulmonary gas embolism caused by localized thermal therapy of the liver.

Case Report

A 71-yr-old man (weight, 44 kg) was scheduled to undergo right segmentectomy of the liver for the removal of a single hepatoma nodule (2 × 3 cm) under general anesthesia. The patient, classified as American Society of Anesthesiologists (ASA) physical status 2 because of emaciation, had no contributory past or family history, and the preoperative laboratory data were within normal ranges. After premedication with atropine and hydroxyzine and placement of both standard ASA monitors and an intraarterial catheter, an epidural catheter was inserted at T8–T9. Anesthesia was induced with fentanyl, thiamylal, and vecuronium, a four-chamber view of the heart was monitored using a 5 MHz transducer (PEF-507SB®; Toshiba Medical, Tokyo, Japan) for a transesophageal echocardiogram. Anesthesia was maintained with inhalation of isoflurane or sevoflurane in a 1:1 gas mixture of oxygen and nitrous oxide. The output/frequencies of MCT and RFA were 60 W/2,450 MHz and 70–90 W/350–450 KHz, respectively.

In all eight cases, MCT or RFA resulted in gas bubbles in the right atrium or right ventricle. Bubbles were identified as highly echogenic dots with high mobility. A gas-bubble-like image also appeared in the left atrium and left ventricle in one patient undergoing MCT (fig. 1). The occurrence of gas emboli continued throughout the thermal intervention, then gradually decreased after the end of heating. There were, however, no remarkable changes in the other variables, including PETCO2 during MCT or RFA.

Discussion

Venous gas emboli can occur during several types of hepatic interventions, such as electrocauterization,7 argon-enhanced coagulation,8,9 and water jet dissection.10 The present report showed the generation of gas emboli detected by transesophageal echocardiogram during hepatic surgery using MCT or RFA.

MCT has been applied clinically to obtain hemostasis during transection of the hepatic parenchyma and in the treatment of nonresectable liver cancer. Microwave electrodes induce a rapidly alternating electromagnetic field. As water molecules follow the changing polarity of the field, heat is generated within the tissue, resulting in...
agitation, which results in frictional heat production within the tissue.

The etiology of gas emboli during these interventions is unclear. Although the exact temperature around the electrode is unknown, the electrode tip temperature will reach more than 80°C. We speculate that evaporation of gases, including nitrous oxide dissolved in the blood and tissue, is a cause of gas bubbles produced during MCT or RFA. Also, the entrainment of atmospheric air from the opened venous sinuses of the liver may exacerbate gas embolism during laparotomic MCT or RFA.

There have been no clinical reports on harmful embolisms caused by MCT or RFA. However, we make the following recommendations for these procedures: (1) The minimal duration of MCT or RFA application required to achieve tissue necrosis should be selected. (2) Precautions should be taken for harmful gas embolism, particularly in patients with the potential for intracardiac right to left shunt, such as persistent foramen ovale. (3) Appropriate monitors, including transesophageal echocardiography, should be used in cases in which extensive therapy is planned.

References

Neuraxial Opioids and Koebner Phenomenon: Implications for Anesthesiologists

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NEURAXIAL opioids are widely used in clinical practice. However, their use is associated with various adverse effects, pruritus being prominent among them.1 This can be distressing to the patient. In most cases pruritus has no adverse consequences. However, in patients with preexisting dermatologic disease, pruritus may lead to an exacerbation of the skin disease.2,3

Case Reports

Case 1

A 42-yr-old man weighing 76 kg with a 20-yr history of psoriasis vulgaris presented for open reduction and internal fixation of a femur fracture. His dermatologic history was suggestive of remissions and relapses. His multiple therapies included topical anthralin, tar, steroids, systemic methotrexate, and phototherapy. Currently, he was receiving oral and topical steroids and anthralin. Care was taken during surgery to avoid pressure and trauma during positioning, placement of monitoring devices, securing of venous access, and administration of regional anesthesia. A combined spinal–epidural approach was used. Spinal anesthesia was performed with 3.0 ml intrathecal 0.5% heavy bupivacaine. At the end of a 90-min procedure, epidural catheter placement was validated by a preliminary dose of 3 ml lignocaine 2% with adrenaline 1:200,000. Once confirmed, 4 mg morphine in 10 ml 0.125% bupivacaine was administered into the epidural space. The patient reported good postoperative analgesic effects. Two hours after administration of epidural morphine, the patient complained of itching of his trunk, face, and legs. He also reported having nausea. Although the nausea responded to intravenous metoclopramide 10 mg, the patient complained of itching of his trunk, face, and legs. The rest of the postoperative period was uneventful. Intravenous naloxone in titrated doses of 1–2 µg/kg, to a total of 0.3 mg, relieved the itching. The rest of the postoperative period was uneventful. Patient was discharged on 6th postoperative day. However, the patient presented to the dermatology outpatient department after 10 days with exacerbation of the disease. She was found to have developed fresh lichen planus papules in a linear array, which were localized in the initial areas of pruritus; i.e., groin, anterior aspect of the trunk, and medial side of the thighs. The epidural infusion was stopped. Intravenous pheniramine was of no avail, and the patient was in great distress. Intravenous naloxone in titrated doses of 1–2 µg/kg, to a total of 0.3 mg, relieved the itching. The rest of the postoperative period was uneventful. Patient was discharged on 6th postoperative day. However, the patient presented to the dermatology outpatient department after 10 days with exacerbation of the disease. She was found to have developed fresh lichen planus papules in a linear array, which were localized in the initial areas of pruritus; i.e., groin, anterior aspect of the trunk, and medial side of the thighs. The patient was reassured and was advised to continue taking the topical steroids.

Discussion

Pruritus after neuraxial administration of opioids is a well-known adverse effect with a wide reported range of incidence of 0–100%.4,5 The incidence is higher with spinal opioids as compared with epidural opioids (46% vs. 8%).6 This compares with a very low incidence of only 1% after the systemic administration of morphine.1,6 Neuraxial morphine has been consistently associated with a higher incidence of pruritus than other opioids.7 Pruritus after neuraxial administration of opioids is unpleasant and difficult to manage. It responds poorly to histamine (H1) blockers and other conventional treatments.8 Naloxone and propofol are two drugs that have been found to be effective against opioid-induced pruritus.9,10

Isomorphic, or Koebner, phenomenon was described originally by Henrich Koebner in 1876.2 In persons with certain skin diseases, trauma is followed by new lesions in the traumatized but otherwise normal skin, and these new lesions are identical to those in the diseased skin. Although best known in psoriasis, it may also occur in other skin diseases, notably lichen planus, lichen nitidus, pityriasis rubra pilaris, vitiligo, and Darier’s disease. Koebner phenomenon usually begins 8–10 days after the injury.3 However, it may appear within 3 days or may be delayed as long as 18 days.3 Those who develop psoriasis at an early age and require multiple therapies to control their disease are more likely to develop Koebner phenomenon.11 When Koebner phenomenon follows medical therapy, it may have medicolegal implications.3

Abbreviations and Acronyms

DM: diabetes mellitus
NMM: neuropathic mood disorder
H1: histamine

References


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In addition to skin trauma, various other forms of skin irritation, such as stripping of the horny layer by adhesive tape, wounding, pressure, shaving, incision, and surgery itself can provoke this phenomenon. Any of the above-mentioned factors could have led to exacerbation of the disease in our patients, with the appearance of new lesions. However, we took due precautions to avoid trauma to the skin and mucous membranes, and we avoided the use of strapping and adhesive electrodes. Although surgical stress can provoke Koebner phenomenon, the peculiar location of fresh lesions at the site of scratching and the linear arrangement along the scratch marks in our patients tacitly implied the itching after administration of neuraxial opioids as a causative factor.

Opioids per se have not been implicated to induce Koebner phenomenon. The exacerbation of the disease in our cases can be attributed to the scratching, opioids being coincidentally the cause of pruritus and, hence, scratching. This may not be uncommon in usual clinical practice. However, the late manifestation of Koebner phenomenon after the skin trauma, and the loss of contact between the anesthesiologist and the patient by this time in most of the cases, may have precluded this clinical problem being noted by the anesthesia fraternity. Furthermore, most cases of exacerbation of the disease may have been ascribed simply to various surgical factors and the stress of surgery, thereby overlooking pruritus due to the use of neuraxial opioids.

Considering the high incidence of pruritus with neuraxial opioids, as well as Koebner phenomenon in patients with psoriasis and lichen planus to the extent of 30–50%, anesthesiologists should be aware of this complication that might result from pruritus induced by neuraxial opioids.

References


Psoas Abscess Complicating Femoral Nerve Block Catheter

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CONTINUOUS three-in-one blockade is widely used for providing postoperative analgesia after knee surgery. Distinct serious complications have been described after femoral nerve block: neurologic injury as well as hematoma compressive and epidural anesthesia. We report a case of psoas abscess complicating a continuous three-in-one blockade.

Case Report

A 35-year-old woman, American Society of Anesthesiologists physical status 1, was admitted for a knee arthroscopic arthrolysis. Before

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course of antibiotics, after which a second computed tomographic scan showed no recurrence of the psoas mass.

Discussion

The main infectious complication reported after a regional nerve block technique is the uncommon but potentially deleterious epidural abscess. Recently, it was shown that after 48 h, 57% of femoral nerve catheters had positive bacterial colonization; however, no patient developed an abscess. In our patient, a culture was not performed on the catheter because on the day of removal the patient reported no discomfort. Yet *S. aureus* found with computed tomographic scan puncture is the most common causative organism cultured from epidural abscess after epidural anesthesia. Pyogenic psoas abscesses are most often associated with vertebral osteomyelitis or Crohn disease. In our patient, the abscess probably resulted from catheter colonization at a superficial site and subsequently wicked the infection from the skin to the psoas space.

This case illustrates the importance of the golden aseptic rules during puncture and catheter insertion for regional anesthesia. In any patient who shows an infectious syndrome and has or has had a continuous nerve block, the possibility of a complication of regional anesthesia should be considered until proof of the contrary, even if the patient had no evidence of superficial infection at the catheter insertion site.

References