THE formation of cranial subdural hematomas after lumbar puncture is a well-known complication of spinal anesthesia, although rare in young patients.1–3 Pathogenesis of this condition has been related to rupture of bridging veins joining the cerebral surface and the dura mater. The precipitating event in these cases is the continuous leakage of cerebrospinal fluid through the dural hole. More unusual is the production of subdural hygromas in the intracranial cavity, which has not yet been described. Our aim is to describe the occurrence of a subdural hygromas complicating inadvertent lumbar puncture and to discuss the problems encountered in their diagnosis and management.

Case Report

A 26-yr-old nulliparous woman presented in active labor. She had a history of migraine, but was otherwise healthy. She requested epidural analgesia. The anesthesiologist identified the L2–L3 interspace and attempted to place a catheter using an 18-gauge Tuohy needle. During placement, an accidental dural puncture was noted, and the needle was withdrawn. A subsequent puncture was performed at the L3–L4 and the epidural catheter was successfully inserted. The labor elapsed uneventfully.

Six hours after delivery, the patient began to complain of an intense, throbbing, occipital headache that worsened when she was in the upright position. There was neither neck stiffness nor fever. A tentative diagnosis of postdural puncture headache was made. Treatment consisted of bed rest, hydration, and analgesics.4,5 At this time, the patient refused a recommended epidural blood patch and, after a 3-day hospital stay, was discharged home.

Fifteen days later, the patient returned to the hospital complaining of continued severe headaches. She did not describe diplopia, tinnitus, or vomiting. Except for mild neck rigidity, her neurologic examination was normal and there was no papilloedema. The patient was again offered an epidural blood patch, which she accepted. During aseptic conditions, an epidural injection of 7 ml autologous blood was performed over 1 min. After an overnight stay at the hospital, the patient became symptom free and was discharged home.

The patient was readmitted 12 days later because her headache had returned to its original intensity after a 4-day period of pain relief. She denied being confused, experiencing vomiting, or having other neurologic symptoms. Cranial computed tomography was performed, which

![Cranial computed tomography showing frontoparietal hygroma (black arrowheads).](Fig. 1)
showed bilateral frontoparietal subdural hygromas, with gyral effacement and small ventricles (fig. 1). The patient was also studied using magnetic resonance imaging venography of the brain that ruled out thrombosis of the cerebral venous sinuses.6 Gadolinium-enhanced and -unenhanced magnetic resonance imaging of the lumbosacral spine was also obtained,7 which failed to show dural enhancement, subdural collections of fluid, or the site of the cerebrospinal fluid fistula.8,9

A subsequent epidural blood patch was performed using 14 ml autologous blood, and adequate intravenous hydration and analgesia were administered. After this management regime, the patient's headaches decreased in severity and were controlled with the intake of mild oral analgesics. The patient was discharged from the hospital 11 days later (on day 39 postpartum). Two weeks after the second epidural blood patch, the patient was symptom free and subsequent cranial computed tomography showed complete resolution of the subdural collections (fig. 2).

Discussion

The occurrence of subdural hygromas after epidural procedures may be more common than has been recog-

ized previously. Subdural hygromas are most often the result of traumatic events, particularly head trauma. The mechanism involved is probably the continuous leakage of cerebrospinal fluid through the orifice made in the dura mater at the time of the lumbar puncture.10,11 If the brain shrinks because of brain atrophy, excessive dehydration, or decreased intracranial pressure, a subdural collection may develop by a passive effusion of fluid. Most subdural hygromas from other causes resolve when the brain is well-expanded. However, some instances of hygromas may evolve to chronic subdural hematomas if the conditions leading to their formation persist over several weeks.10

The diagnosis of subdural hygroma can be difficult on a clinical basis alone, especially if an obvious cerebrospinal fluid leak is not reported. We performed a review of the literature in regard to subdural hygromas associated with postdural puncture headache and have not found any previous references to this condition. We suggest that the existence of a subdural hygroma should be included in the differential diagnosis of the cause of a persistent headache presenting after a dural puncture, especially if first steps of management have already failed.

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Preoperative Use of Enoxaparin and Tirofiban: Possible Association with Increased Bleeding Postbypass

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THE purpose of this case report is to alert anesthesiologists and other medical and surgical staff as to the potential for increased postoperative bleeding in patients undergoing heart surgery after having received the recently introduced low-molecular-weight heparin (LMWH) preparations (e.g., enoxaparin) and new platelet inhibitors (e.g., tirofiban) in the setting of acute coronary syndromes or coronary interventional procedures.

Case Report

Four days after acute anterolateral myocardial infarction, a 75-yr-old, 56-kg man was scheduled for coronary arterial revascularization. His preoperative medications were 60 mg enoxaparin subcutaneously two times a day, aspirin and metoprolol orally, and intravenous infusion of the platelet glycoprotein (GP) IIb/IIIa receptor inhibitor tirofiban (Aggrastat, Merck, West Point, PA). The tirofiban infusion was discontinued at midnight before the day of operation.

Normal electrolyte and hematologic panels were obtained preoperatively: prothrombin and activated partial thromboplastin times (aPTT) of 13.9 (1.1 × control, international standardized index [ISI] 2) and 34.9 s (1.1 × control), respectively, a hematocrit of 48.1%, and a platelet count of 152,000/µl. Creatinine and blood urea nitrogen levels were normal as well (1.2 mg/dl and 10 mg/dl, respectively). During 90 min of extracorporeal support, with an ischemic time of 51 min, the surgical procedure involved anastomosis of the left internal thoracic artery to the left anterior descending coronary artery and a free reversed saphenous vein graft to the left circumflex coronary artery. During cardiopulmonary bypass (CPB), the triggers for additional heparin administration were an activated clotting time of less than 480 s and a whole blood heparin concentration less than the patients' pre-CPB reference level (4.2 U/ml) using an automated protamine titration method (Hepcon, Medtronic Blood Management, Parker, CO). After discontinuation of CPB, the total dose (25,000 IU) of heparin was neutralized with a total of 350 mg protamine sulphate, as guided by the automated protamine titration method. Nevertheless, the surgical wound appeared diffusely “wet,” and the subjective diagnosis of microvascular bleeding was made by the staff surgeon. Based on immediately available whole blood tests, which revealed a normal activated cloting time (> 120 s); no detectable heparin (automated protamine titration heparin concentration method); a hematocrit of 22%; a platelet count of 55,000/µl; and a whole blood prothrombin/PTT of 17.8 s (1.48 × control, ISI 2)/40.9 s (1.52 × control), respectively; 2 U of packed erythrocytes and one single donor pack (apheresis) of platelets were transfused. The patient remained normothermic (36.9°C) after discontinuation of bypass and a whole blood heparin concentration less than the patients’ pre-CPB reference level. Support was provided solely from institutional and/or departmental sources.

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Key words: Complications; low-molecular-weight heparin; platelet GP IIb/IIIa inhibitor.

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bin/aPTT values of 15.2 s (1.2 × control)/61.7 s (1.9 × control), as well as a single donor apheresis platelet unit (six-pack equivalent) for a platelet count of 82,000/μl. Persistent excessive CTD (660 ml) during the third 8-h period necessitated the administration of an additional six-pack of random donor platelets. Despite excessive CTD (3,750 ml over 2 days), necessitating the transfusion of 7 U erythrocytes, the patient was hemodynamically stable, did not require vasopressor therapy, and was extubated the morning after surgery (15 h postoperatively). No additional hematologic assays were performed. Neither thromboelastography nor anti-Xa assay are available in our institution, and a point-of-care platelet function test, although available, was not performed by the managing physicians, who were not aware of the potential for excessive postbypass bleeding in the setting of concurrent use of LMWH and tirofiban. The patient was discharged from the intensive care unit 2 days later and from the hospital after 5 days.

**Discussion**

Based on results of the ESSENCE trial, the combination of LMWH and aspirin is increasingly being used for patients with acute coronary syndromes. LMWHs, such as enoxaparin, are derived from unfractionated heparin (UFH) and produce their major anticoagulant effect by accelerating the binding of antithrombin III (AT III) to predominantly factor Xa (enoxaparin has an anti-Xa:anti-IIa activity ratio of 3.8:1.0). LMWHs generally have enhanced bioavailability, a longer half-life (two to four times as long as that of UFH), a dose-independent renal clearance, and a slightly decreased incidence of heparin-induced thrombocytopenia. LMWHs exhibit an antithrombotic effect that is assessed most accurately by measuring anti-Xa activity (which is often unavailable). The inability of routine doses of protamine to completely reverse the anti-Xa activity represents a significant limitation. In an animal study, the dose of protamine sulfate necessary to fully minimize bleeding was 10-fold greater for LMWH than for UFH. In addition, excessive CTD was observed, when enoxaparin was used as an alternative to UFH in patients with heparin-induced thrombocytopenia undergoing CPB.

Platelet aggregation has been implicated as one of the major causes in the development of unstable angina and acute myocardial infarction. Activation of the platelet IIb/IIIa GP receptor represents a crucial step in platelet aggregation, and 80–100% blockade of this receptor should completely suppress platelet aggregation. Therefore, potent platelet inhibitors, such as the intravenous GP IIb/IIIa inhibitors have been developed, and their clinical effectiveness is supported by several studies. Recent data showed that substantially increased blood
usage was associated with use of the GP IIb/IIIa inhibitor abciximab, especially if surgery was performed within 24 h after receiving the medication. However, these findings have not been universal. Platelet transfusions, which will reduce the drug’s receptor occupancy on platelet surfaces to less than the critical 50% threshold associated with prolongation of the bleeding time, is the most appropriate therapy.

Although postbypass bleeding can have a multifactorial cause, several important factors, such as prolonged duration of CPB and type of procedure, are associated with transfusion of hemostatic blood products and increased CTD. The postoperative CTD of our patient, 2,360 ml over the first 24 h and 1,380 during the second 24 h, far exceeded the previously reported mean 24-h cumulative chest tube output of 986 ± 581 ml. In the case presented, the CPB duration was rather short, bleeding from the two distal anastomoses should have been easily detected, and normothermia was maintained after CPB. The role of tirofiban with respect to excessive bleeding is uncertain based on its short half-life time of about 90 min, and the infusion was discontinued the night before. Use of this agent may be important because the patient also received enoxaparin on the day of the surgery. The peak anti-Xa activity after subcutaneous LMWH administration occurs within 3–4 h, and the anti-Xa levels are approximately 50% of peak levels 12 h later.

When patients who are receiving a LMWH preparation are scheduled to undergo a surgical procedure, such as coronary revascularization, it would be prudent to discontinue the LMWH at least 24 h before surgery or to replace the LMWH with an intravenous infusion of UFH. This approach may be even more important in the setting of concurrent administration of a potent platelet inhibitor, such as tirofiban. In the setting of concurrent use of LMWH and platelet inhibitors, routine coagulation results (prothrombin, aPTT, activated clotting time, platelet count) will most likely not be helpful in guiding appropriate therapy. Further studies are needed to evaluate the relation between concomitant use of these new antithrombotic agents and perioperative bleeding, as well as the use of new monitoring systems and heparin reversal agents. Future advances in coagulation monitoring may help in the evaluation of patients with residual anti-Xa activity, such as the HepTest anti-Xa (Hematest, Richmond, VA), may facilitate evaluation of the degree of residual platelet inhibition as well as the degree of CPB-related platelet dysfunction. Novel reversal agents, such as the [1+8RGD] protamine variant, which reverses conventional UFH and enoxaparin, may be clinically useful in reversing the anticoagulant effects of LMWH.

References

CASE REPORTS

Unanticipated Difficult Intubation as a Result of an Asymptomatic Vallecular Cyst

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Although vallecular cysts are often asymptomatic and harmless to the patient, discovery of a large vallecular cyst after induction of anesthesia is a potentially life-threatening problem for the patient and provides for the anesthesiologist a challenge in airway management. This report describes the treatment of a patient with an asymptomatic vallecular cyst that was discovered during rapid-sequence induction of general anesthesia.

Case Report

A 47-year-old woman presented for elective right knee arthroscopy and removal of painful orthopedic hardware of the lower extremity. Her medical problems included obesity and a history suggestive of gastroesophageal reflux disease and hypertension treated with enalapril. She had no known drug allergies. A year previously she received a knee arthroscopy. She had no known drug allergies. A year previously she received a knee arthroscopy.

She was premedicated with 10 mg metoclopramide intravenously and 30 mL sodium citrate orally. After application of routine monitoring devices, she was preoxygenated. Anesthesia and paralysis were induced with 200 mg propofol and 120 mg succinylcholine while cricoid pressure was applied. Direct laryngoscopy with a Macintosh 3 blade revealed a 2-cm cyst arising from the right side of the vallecula; it was pedunculated with a 5-mm stalk, making it somewhat mobile. The cyst completely obscured the view of the epiglottis and larynx, preventing intubation despite multiple attempts by two anesthesiologists using Macintosh 3 and Miller 2 blades, and increased neck extension and neck were observed. Chest auscultation revealed a mild expiratory wheeze.

She was repremedicated with 10 mg metoclopramide intravenously and 30 mL sodium citrate orally. After application of routine monitoring devices, she was preoxygenated. Anesthesia and paralysis were induced with 200 mg propofol and 120 mg succinylcholine while cricoid pressure was applied. Direct laryngoscopy with a Macintosh 3 blade revealed a 2-cm cyst arising from the right side of the vallecula; it was pedunculated with a 5-mm stalk, making it somewhat mobile. The cyst completely obscured the view of the epiglottis and larynx, preventing intubation despite multiple attempts by two anesthesiologists using Macintosh 3 and Miller 2 blades, and increased neck extension and thyroid pressure. Because of a decrease in oxygen saturation (SaO2) the patient was ventilated via mask as cricoid pressure was maintained. Despite insertion of an oropharyngeal airway, this proved to be difficult; laryngospasm began to develop as muscle relaxation waned. After a second 80-mg dose of succinylcholine, intubation was attempted using a Miller 2 laryngoscope blade and a styletted 7.0-mm endotracheal tube. By using the tube to push the cyst aside, intubation of the trachea was performed after a brief although limited view of the laryngeal inlet.

The remainder of the anesthetic proceeded uneventfully. While the patient remained anesthetized, an otolaryngologist was consulted and the cyst was surgically removed. Dexamethasone 10 mg was given as suggested by the otolaryngologist. Anesthesia was discontinued and 100% oxygen was administered; after the patient awakened, the pharynx was carefully suctioned and the trachea was extubated.

Her postoperative course was also uneventful and she was discharged to home on the same day without any airway-related problems apart from a sore throat. During further questioning in the postanes-

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Her postoperative course was also uneventful and she was discharged to home on the same day without any airway-related problems apart from a sore throat. During further questioning in the postanes-
thysia care unit she admitted to a several-year history of dysphagia. The pathology report confirmed the findings of a 2.2 × 2.2 × 0.5 cm, 0.3-cm thick membranous wall vallecular cyst.

Discussion

Cysts of the larynx are rare and usually follow a benign course. They can, however, present with acute airway obstruction by nature of their location. Laryngeal cysts have been previously reported in the anesthesia literature.1–4 However, this case presented the additional challenge of producing unanticipated upper airway obstruction during rapid-sequence induction general anesthesia with subsequent successful intubation of the trachea by manipulation of the cyst with a styletted endotracheal tube.

Laryngeal cysts have been extensively studied and classified by site, size, contents, and relation to the laryngeal mucosa. Ductal cysts or mucous retention cysts are the most common type of laryngeal cysts, comprising 75% of cases, and are formed by distention of obstructed collective glanular ducts. (Saccular cysts, which comprise the remaining 25%, arise from the saccul, an extension of the ventricle; they may cause respiratory symptoms by enlarging the aryepiglottic fold).5,6 Ductal cysts are usually small, approximately 1–5 mm in diameter and as such are often asymptomatic.5 They are usually only discovered incidentally at the time of routine otolaryngologic examination or at postmortem examination.1,7

As a result, the exact incidence of laryngeal cysts is unknown but is estimated to be quite low.8 There is no gender predominance and, although they may occur at any age, a greater prevalence in the fifth and sixth decades has been observed.5,6 The most common location of ductal cysts is on the true vocal cords, whereas the second most common site is in the vicinity of the epiglottis—on its lingual surface or in the vallecula itself.5–7 Although they are usually asymptomatic, they may occasionally cause symptoms of stridor and failure to thrive (particularly in the newborn),8–10 cough, dysphonia, foreign body sensation and dysphagia.6

In our case, a previously undiagnosed vallecular cyst presented as a difficult intubation. We had no reason to suspect any difficulties with intubation because the patient had no symptoms preoperatively—the patient’s history of dysphagia was only elicited on specific, symptom-directed questioning postoperatively. In view of her normal airway examination, previous uneventful general anesthesia and history of gastroesophageal reflux disease, we thought that rapid-sequence induction of anesthesia and tracheal intubation was appropriate for our patient. Had both tracheal intubation and mask ventilation been impossible, aspiration of the cyst via a spinal needle might have been an option. In contrast to previous case reports in which intubation was either abandoned or performed blindly with the aid of an gum elastic bougie,1–4 we were able to intubate the trachea by displacing the pedunculated vallecular cyst with the endotracheal tube. We elected to obtain an otolaryngology consultation while the patient was still anesthetized so the cyst could be removed before extubation. This avoided the risk of any airway obstruction after extubation caused by the potential “ball-valve” effect of the cyst. This case would certainly have been managed differently had we known that the cyst was present before induction of anesthesia. For example, regional anesthesia or awake fiberoptic intubation would have been reasonable alternatives.

In summary, there are several causes for an unanticipated difficult intubation during rapid-sequence induction of general anesthesia. This case report helps to highlight one such cause—an asymptomatic vallecular cyst—and its subsequent management, resulting in successful tracheal intubation.

References

Shivering Complicating the Treatment of Neurologically Impaired Surgical and Intensive Care Unit Patients

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NEW-ONSET rhythmic movement can confound diagnosis and complicate management in patients with central nervous system disease. Rhythmic movement may result from the nonconvulsive effects of drugs (e.g., etomidate), the convulsive effects of drugs (e.g., local and general anesthetics), epileptic seizures, or other causes. Regardless of the source, inadvertent patient motion may complicate awake neurosurgical procedures, especially if the surgery involves real-time assessment of neurologic function or the acquisition of fine electrical recordings.

Because movement disorders may reflect new or evolving brain injury or result from more benign causes, the identification of the source of the movement may lead to widely divergent patient care interventions. We describe two patients who experienced new-onset, nonconvulsive, rhythmic movement that interfered with their treatment. In both, the source of the movement was normothermic or near-normothermic shivering.

Case Reports

Case 1

A 29-yr-old man with a history of aqueductal stenosis was admitted to the neurologic intensive care unit after surgery to remove an infected right-sided ventriculoperitoneal shunt. Six hours later, generalized tonic-clonic convulsions occurred, and electroencephalography (EEG) confirmed status epilepticus. When initial therapy with diazepam, phenytoin, and phenobarbital failed to control the seizure activity, an amobarbital infusion was begun. The trachea was intubated, and the lungs were mechanically ventilated. Additionally, the patient was treated with 100 mg intravenous phenytoin four times per day.

After 18 h of amobarbital infusion, subsequent EEG was obtained while the patient was without seizures or rhythmic movement. This revealed 0.5–1 Hz δ activity with superimposed rhythmic 10–12 Hz θ activity. Also observed were sporadic or quasiperiodic sharp waves arising from the right frontotemporal region. These findings were consistent with residual barbiturate effect plus potentially epileptogenic components. The amobarbital infusion rate was reduced to permit further evaluation of the patient’s neurologic status. However, over the ensuing 4 h, the patient began experiencing rhythmic movement interpreted as clinical evidence for return of seizures. The amobarbital infusion was returned to the previous delivery rate, and a prolonged EEG recording was begun. The EEG data were consistent with general anesthesia; no seizure activity was observed.

A second attempt to wean the patient 30 h after the original administration of the barbiturate infusion resulted in intense, violent, rhythmic muscle activity, although the patient remained unresponsive to stimuli. The EEG findings were consistent with altered consciousness as a result of residual anesthetic; no clear seizure activity was identified. Instead, EEG tracings contained a rhythmic artifact that was represented in the scalp electrode recordings as 5- to 6-Hz waxing and waning deflections, consistent with a nonepileptic “movement disorder.” The patient’s core (bladder) temperature was 37.8°C. Based on the available data, a tentative diagnosis of near-normothermic shivering was made.

The patient was stripped of all clothing. The skin was irradiated using two 500 W infrared heat lamps (Emerson Equipment Model 96H; Emerson Electric, St. Louis, MO), with the light source 1 m from the skin surface. Within approximately 90 s, all rhythmic muscle activity ceased. Two minutes later, the light therapy was discontinued, and the rhythmic activity returned. When this sequence was repeated 5 min later with the same results, the diagnosis of near-normothermic shivering was confirmed. Throughout the test, the patient’s temperature was unchanged and no antipyretics were administered.

The amobarbital infusion was not reinitiated. Over the next 24 h, the patient awakened from the amobarbital anesthetic without EEG or clinical evidence of seizures. The trachea was extubated 2 days later. During this period, the patient’s skin was kept warm by the use of warm blankets, and the rhythmic muscle activity did not return. The patient was discharged from the intensive care unit 1 week after surgery.

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Key words: Convective warming; electroencephalography; hypothermia; movement disorders; neurosurgery; radiant warming; temperature; thermoregulation.
We describe two patients whose clinical course was complicated by shivering. In the first, the shivering interfered with the clinical and electrophysiologic diagnosis of seizures, thus extending the period in which the patient received an amobarbital anesthetic. In the second, intraoperative shivering again precluded accurate electrophysiologic monitoring and, additionally, increased the risk of bleeding. This patient experienced extreme anxiety associated with the shivering. Had the coupled shivering and anxiety not been treated, it would have been necessary to discontinue the surgery.

In patient 1, we had evidence that the shivering occurred during a period without core hypothermia. (A possible cause is that the hypothalamic “set point” for temperature regulation was increased, perhaps because a fever was developing.) In both patients, shivering ceased within minutes after initiating skin warming. Thus, the period of warming, that was sufficient to stop shivering, was insufficient to meaningfully alter core temperature.

Distinguishing between shivering and other forms of rhythmic motor activity can be difficult. One possible approach is to characterize and quantify the frequency of the motor activity and compare it to the reported properties of motor activity in different physiologic states. Electromyographic recordings at frequencies of 5–12 Hz can represent shivering, but there is overlap with a variety of physiologic and pathologic conditions. Although shivering also may exhibit a fairly characteristic 4- to 8-cycles/min waxing and waning pattern, this was not specifically identified as an artifact within the EEG in the first patient we described.

The effect of barbiturate infusion on the first patient’s rhythmic movements was believed to represent evidence for underlying seizure activity. In hindsight, it probably represented the influence of an anesthetic agent on the “interthreshold range” for thermoregulation (i.e., the 0.2°C range outside of which thermoregulatory mechanisms are initiated). For the same reason, the early use of propofol probably prevented shivering with hypothermia (i.e., bladder temperature 35.6°C) during the second patient’s second surgery. After we suspected a diagnosis of normothermic or near-normothermic shivering in our patients, the treatment was patterned after the reports of Sharkey et al. This therapy is based on the principle that reduction in skin temperature, independent of core hypothermia, can initiate shivering. Warming the skin in such patients will

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abruptly halt shivering. Sharkey et al.7 irradiated shivering patients with a warming light. The patients ceased shivering in 61 ± 10 s (mean ± standard deviation). However, after the warming light irradiation was discontinued, the patients resumed shivering in 43 ± 7 s. We used this principle of skin warming to effectively treat shivering in our patients and, in patient 1, to establish a diagnosis. In patient 2, direct skin irradiation was impossible because of space limitations caused by surgical draping and equipment. Instead, we used a forced-air warming blanket.9,10 In our patients, the rapid onset of the warming effect (patients 1 and 2) and the rapid offset (patient 1) were valuable in establishing the origin of the undesired movement.

Patient 2 had bouts of anxiety during both surgeries. In the first, when the patient shivered violently, the anxiety was debilitating. Although not relieved by the conventional approaches of conversation and sedation, the anxiety was halted by skin warming. Although core hypothermia plus input from thermal sensors within the skin may have directly affected the anxiety spells, the experience during the first surgery also suggests a correlation between muscle activity and the anxiety. Surface warming, which rapidly halted the shivering, also abolished the anxiety. These observations suggest that shivering per se, perhaps acting through muscle receptors (e.g., muscle spindles), served to modulate the anxiety attack. Such an interpretation is consistent with a large body of experimental evidence that shows that increased muscle afferent traffic (as would be expected during shivering) has the potential to desynchronize the EEG and produce alterations in mentation and behavior in awake and lightly anesthetized subjects.11

In summary, we present two cases in which the presence of shivering confounded the diagnosis and treatment of patients having central nervous system disease. Warming the skin, without altering core temperature, halted the shivering within minutes and, in one patient, also alleviated anxiety. Based on this experience, when patients are normothermic, we recommend the inclusion of shivering in the differential diagnosis of patients who experience new-onset rhythmic movements. Additionally, our report confirms the effectiveness of using heating lamps or forced-air warmers to prevent, diagnose, and treat shivering.

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Management of Post-thoracotomy Pseudoangina and Myofascial Pain with Botulinum Toxin

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Case Report

A 52-yr-old man with a 70% stenosis of the left main coronary artery underwent a single coronary artery bypass graft with a LIMA graft to the proximal left anterior descending coronary artery in 1988 at age 42. The coronary artery bypass graft was performed during balanced narcotic and inhalational anesthesia through a median sternotomy incision with the patient in the supine position with both arms adducted. Invasive vascular monitoring included a right radial artery catheter and a central venous catheter inserted via the right internal jugular vein. The LIMA was harvested over 25 min with exposure provided by a Favaloro retractor. The LIMA-to-left anterior descending anastomosis was conducted during 25 min of complete cardiopulmonary bypass and circulatory arrest. The immediate postoperative course was uneventful, and the patient was discharged on the sixth postoperative day. The patient was readmitted for a 24-h hospital stay 2 days after discharge for tube thoracostomy drainage of a spontaneous left pneumothorax.

One week after the second hospital discharge, the patient first noted left anterior chest wall pain during movement of the left arm with radiation to the left side of the neck, the left shoulder, and left-sided scapula. The pain was dysesthetic, nonlancinating, and pressing, and was initiated by exercise of the left arm, particularly lifting and carrying light items, such as books or a briefcase. The pain was relieved only by resting the left arm. Exercise electrocardiography showed no evidence of cardiac ischemia or myocardial infarction. Cardiac angiography showed no significant additional coronary artery disease or LIMA graft stenosis or occlusion. Cardiac ejection fraction and myocardial performance were assessed as normal, and oral analgesics for prolonged postoperative incisional pain were prescribed.

Between 1988 and 1997, the patient consulted innumerable specialists, including cardiovascular surgeons, physiatrists, cardiologists, and rheumatologists, and underwent five additional coronary arteriograms to assess “recurrent angina.” The patient was treated for presumed costochondritis with oral analgesics, nonsteroidal antiinflammatory agents, muscle relaxants, and trigger-point injections in the back. The patient also was prescribed long-term aspirin prophylaxis for coronary artery disease and lovastatin for hypercholesterolemia, and encouraged to take sublingual nitroglycerin for exertional “anginal”, i.e., left chest wall and upper arm pain and muscle spasm. The patient continued to exercise despite attacks of left-sided chest wall pain, participated in several marathon runs over the years, but gave up golf and weight-lifting because of severe pain and spasms in the left pectoralis major muscle precipitated by movement and weight-bearing exercise of the left upper extremity.

The patient was ultimately referred to the Louisiana State University Medical Center Multidisciplinary Pain Mastery Center for evaluation in November 1997, and, at physical examination, was found to have a 5-cm circumferential area of dysesthesia, muscular irritability, and spasm overlying the left-sided first and second parasternal intercostal spaces. The remainder of the physical examination, particularly the neuromuscular examination of the left upper extremity, was within normal limits. A magnetic resonance imaging study of the thorax in January 1998 showed no evidence of costochondritis, costochondral, or costotransverse junction separation or dislocation; rib fracture, callus, or pseudoarthrosis; sternal or xiphisternal abscess; pseudoar-
thorosis; or dehiscence. The mediastinum, hemidiaphragms, lungs, and hemithoraces were consistent with poststernotomy and otherwise normal. The hemidiaphragms were symmetrically shaped and positioned bilaterally, and there was no radiographic evidence of phrenic neuropathy. Phrenic nerve conduction studies were not indicated.

The patient received an intramuscular injection of 50 U botulinum A toxin suspended in 10 ml bupivacaine, 0.25%, into the dysesthetic trigger point described in the left pectoralis major muscle in January 1998, with dramatic reduction of symptoms. Supplemental therapy for breakthrough pain was initiated at the same time with 75 mg oral venlafaxine and 500 mg oral methocarbamol, both taken at bedtime. The patient remains on a regimen of semiannual intramuscular trigger-point injections with 50 U botulinum toxin supplemented orally with night-time venlafaxine, a selective norepinephrine-reuptake-inhibiting antidepressant with analgesic properties, and methocarbamol, a skeletal muscle relaxant. The patient continues to participate in jogging marathons and has restarted a weekly golf game.

Discussion

Brachial plexus and musculoskeletal injuries of the chest wall and upper extremities have been reported after cardiac surgery, with incidences ranging from 2% to 38%.1 Brachial plexus injuries have been associated with the median sternotomy technique, paramedian sternotomy, forced or prolonged sternal retraction with the Favoloro retractor, penetrating trauma from first-rib fractures, hyperabducted arm positioning, and needle trauma during insertion of internal jugular catheters.3 Musculoskeletal injuries have also been associated with rib fractures from forced sternal retraction, costochondral cartilage separations, and costotransverse rib disarticulations.1–3 In a prospective investigation of 162 patients undergoing median sternotomy for cardiac surgery, Roy et al.1 reported a significantly greater incidence (39%) of musculoskeletal complaints and neurologic dysfunction in patients undergoing internal mammary artery grafts for aortocoronary bypasses than in patients (17%) undergoing valvular and other cardiac operations without internal mammary artery graft harvesting.

Neurologic dysfunction from brachial plexus and peripheral nerve injuries after cardiac surgery often present with neuropathic pain and dermatomal sensorimotor disturbances that require diagnostic confirmation with nerve conduction studies and electromyography.2 Musculoskeletal complaints after cardiac surgery were described by Roy et al.1 as pain at rest and exercise and limitation of motion in the arms, shoulders, chest, and back. Postoperative musculoskeletal complaints were difficult to describe clinically and confirm radiographically in the absence of old fractures, cartilaginous separations, and rib disarticulations.1 Using either chest wall computed tomography or exploratory resternotomy, Shafir et al.5 identified inadvertent paramedian sternotomy as a cause of postoperative musculoskeletal pain, chronic thoracic fistulas, sternal osteomyelitis, and sternal dehiscence in 11 of 55 patients undergoing cardiac surgery. Shafir et al.5 recommended that when a painful paramedian sternotomy is diagnosed radiographically or surgically at resternotomy, sternectomy and myocutaneous flap wound closures be performed rather than simple sternal reclosures.

We reported a case of chronic myofascial pain in the left upper thorax and arm repeatedly misdiagnosed as angina pectoris and coronary artery spasm despite normal coronary angiography and left ventricular wall motion and ejection fraction (table 1). Although the patient requested sternal reexploration to assess faulty sternotomy, magnetic resonance imaging of the chest wall, mediastinum, and thorax was initially recommended to rule out sternal fistula, sternal osteomyelitis, occult sternal dehiscence, costochondral cartilage separations, old rib fractures or pseudoarthroses, costotransverse rib disarticulations, and pleuropneumopericardial defects or hernias. Chest magnetic resonance imaging was selected over chest radionuclide scanning, a better imaging technique to detect chronic infection and deep abscesses, for its superior soft tissue and bone density delineation and

<table>
<thead>
<tr>
<th>Date</th>
<th>Treatment or Procedure</th>
<th>Pain Type and Outcome</th>
</tr>
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<tbody>
<tr>
<td>November 1997</td>
<td>Open heart bypass surgery</td>
<td>Cardiac angina relieved; pseudoangina began; function restricted</td>
</tr>
<tr>
<td>January 1998</td>
<td>Oral nitrates</td>
<td>Pseudoangina unrelieved; function restricted</td>
</tr>
<tr>
<td></td>
<td>Thoracic MRI scan</td>
<td>Pseudoangina redefined as myofascial pain</td>
</tr>
<tr>
<td></td>
<td>Initial botulinum toxin (BoTox) injection</td>
<td>Myofascial pain relieved; nighttime breakthrough pain only; full function resumed</td>
</tr>
<tr>
<td>July 1998–Present</td>
<td>Subsequent BoTox injection (July 1998); oral adjuvant therapy with venlafaxine and methocarbamol</td>
<td>Myofascial pain relieved; nighttime breakthrough pain relieved; full function continued</td>
</tr>
</tbody>
</table>

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Myofascial pain syndromes may mimic angina pectoris after median sternotomy for cardiac surgery. Brachial plexus and musculoskeletal injuries may occur even more commonly after median sternotomy for cardiac surgery and have been correlated with harvesting of the ipsilateral internal mammary artery, usually the left, but not with intraoperative arm positioning. Precise diagnostic imaging techniques may be necessary to assess chronic chest wall pain after median sternotomy for cardiac or anterior mediastinal surgery and to eliminate faulty sternotomy techniques, cartilaginous injuries, and rib fractures or disarticulations. Nerve conduction studies may be necessary to assess sensorimotor dysfunction in brachial plexus injuries. Although usually reserved for painful spasticity disorders, botulinum A toxin injections may offer a new, safe, and effective technique to manage myofascial pain syndromes.

References


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Life-threatening Ventricular Dysrhythmias with Inadvertent Asynchronous Temporary Pacing after Cardiac Surgery

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POTENTIALLY lethal arrhythmias have long been recognized as a rare complication of cardiac pacing.1,2 The incidence of this catastrophic event is thought to be low even in cases of asynchronous ventricular pacing, presumably because the electrical stimuli provided by the pulse stimulator usually remain below the fibrillation threshold.3 Furthermore, the possibility of serious pacemaker-induced arrhythmias has been greatly reduced by the introduction of advanced demand modes of pacing into clinical practice. Nevertheless, this problem can still be encountered when pacemaker pulse generator is switched to asynchronous mode. We report a case in which a recently introduced and commonly used epicardial temporary pacemaker (Model 5388, Medtronic Inc., Minneapolis, MN) was accidentally switched several times to asynchronous mode, causing ventricular fibrillation in a patient after cardiac surgery.

Case Report

A 49-yr-old man with a history of aortic insufficiency of rheumatic origin and hypertension, treated with calcium channel blockers and angiotensin-converting enzyme inhibitors, was admitted to the hospital for elective aortic valve replacement. The preoperative coronary angiogram showed no signs of coronary artery disease. The electrocardiogram (ECG) demonstrated signs of left ventricular hypertrophy with strain pattern. Holter monitoring showed sinus rhythm with occasional premature atrial contractions and premature ventricular contractions (couplets and triplets).

The patient underwent aortic valve replacement with a 27-mm stentless porcine valve. No intraoperative problems were encountered. In the immediate postoperative hours, several episodes of premature ventricular contractions were noticed, which resolved after normalization of potassium balance (fig. 1A). Two hours later his heart rate slowed down. Second-degree atrioventricular block with a ventricular rate of 50–54 beats/min was apparent on the ECG trace. Dual chamber paced, dual chamber sensing, and dual response (DDD) pacing was started (Medtronic 5388 dual chamber temporary pacemaker) with an atrial sensitivity of 0.5 mV and output of 10 mA, ventricular sensitivity of 2 mV and output of 10 mA, and an atrioventricular interval of 170 ms. This resulted in ventricular rate of 100–102 beats/min (atrial sensing and ventricular pacing; fig. 1B). The patient was uneventfully extubated 6 h after surgery. Occasional premature ventricular contractions, present during the first night after surgery, resolved after removal of the pulmonary artery catheter on the morning of the first postoperative day. The parameters of the pacemaker were checked. The underlying rhythm at this time was third-degree atrioventricular block with a rate of ventricular escape of 50–60 beats/min. DDD pacing with the previous parameters was commenced, and the patient was transferred to the surgical ward on a telemetry bed. Before the transfer, the pacemaker was checked again, and settings were locked. Serum magnesium and potassium levels were normal.

Approximately 1 h after the transfer, asynchronous ventricular pacing at a rate of 80 beats/min was noticed on ECG telemetry monitor. Several minutes later, the patient suddenly lost consciousness. A decrease of pacing stimuli on the T wave with resultant ventricular fibrillation was observed on the ECG tracing (fig. 2). Cardiopulmonary resuscitation was initiated. Over an 8-min period, the patient received three external defibrillatory shocks of 300 J. After the second shock, slow ventricular rhythm of 50 beats/min was observed with rapid degeneration to polymorphic ventricular tachycardia. After the administration of 80 mg lidocaine, 2 g magnesium sulfate, and a third shock, stable rhythm with dual-chamber pacing of 100 beats/min was achieved. The patient regained consciousness and was transferred back to the intensive care unit. On arrival, the pacemaker was found to be in asynchronous mode with maximal atrial and ventricular output (20 and 25 mA, respectively). The underlying rhythm at this time was complete heart block with ventricular escape of 60–65 beats/min. The impulse generator was reprogrammed to DDD mode with the parameters described previously and was locked. The patient was stable and breathing spontaneously with a face mask. No neurologic deficit was found. After 24 h in the intensive care unit, the patient continued with DDD pacing with atrial tracking and a ventricular rate of 100–104 paced pulses/min. The underlying rhythm was complete heart block. At this point, the pacemaker was checked again because a short run of ventricular tachycardia was observed on the ECG monitor. The pacemaker was found to be in asynchronous mode again (dual chamber pacing with the previous parameters was commenced, and the patient was transferred to the surgical ward on a telemetry bed. Before the transfer, the pacemaker was checked again, and settings were locked. Serum magnesium and potassium levels were normal.

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Key words: Asynchronous pacing; pacemaker arrhythmias; ventricular fibrillation.
Fig. 1. Rhythm disturbances during the night after surgery. (A) Multiple multifocal premature ventricular contractions; (B) DDD pacing with atrial tracking.

Fig. 2. Initiation of ventricular fibrillation by pacing impulse, superimposed with T wave, and electrocardiogram after resuscitation.
paced, zero sensing, zero inhibition [DDO]: dual-chamber asynchronous pacing at 80 paced pulses/min, atrial output of 20 mA, and ventricular output of 25 mA). The impulse generator was replaced and checked in detail. On close examination it was found that this particular model of temporary pulse generator was very easily switched from DDD mode (atrial-triggered, ventricular-inhibited pacing) to dual-chamber asynchronous (DOO) pacing (the factory programmed default mode) by light pressure (single-finger touch) on the emergency button situated on the front panel of the device (fig. 3). This emergency DOO default mode overrides any preset (locked) pacing parameters and can be switched on even when the pacemaker is off.

The patient experienced no further dysrhythmias after changing to another atrioventricular sequential demand pulse generator (Model 5330, Medtronic Inc.). He remained in complete atrioventricular block and eventually required implantation of a permanent pacemaker.

Discussion

Reported cases of ventricular tachyarrhythmias caused by a decrease in pacemaker stimulus during the “vulnerable period” of cardiac cycle have occurred in settings of acute myocardial ischemia, electrolyte disturbances, and drug effects, when the ventricular vulnerability to fibrillation is lower.4 Our patient did not have demonstrable coronary artery disease. Potassium and magnesium concentrations before arrhythmias were normal. However, the potential for altered ventricular excitation threshold must have existed. Evidence for this was atrial and ventricular dysrhythmias on the patient’s preoperative Holter recording. The probable cause for this was cardiomyopathy secondary to long-standing ventricular volume overload.

This case report should alert clinicians and nursing staff to the possible danger of accidental asynchronous pacing with this model of temporary pulse generator. This could be atrial asynchronous (AOO), ventricular asynchronous (VOO), or DOO depending on whether the device was programmed AAI (atrial inhibited or demand), VVI (ventricular inhibited or demand) or DDD (also DDI; dual-chamber inhibited), respectively. Even light pressure on the emergency button, located on the front panel of the device, may trigger default mode of asynchronous pacing. This easily could happen if the patient or care provider mistakenly mishandled or touched the emergency button on the pulse generator (e.g., by picking up the pacemaker). Possible means to reduce this risk include eliminating this feature or requiring more than one depression or some other method to activate default-mode asynchronous pacing.
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