Severe Thrombocytopenia, Type 2B von Willebrand Disease and Pregnancy

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VON Willebrand’s Disease (vWD) is the most common inherited bleeding disorder in humans with an estimated incidence as high as 2–3% in the general population.1 Characterized by abnormal platelet interactions with the subendothelium or other platelets, the disease is caused by changes in the multimeric glycoprotein, von Willebrand’s Factor (vWF).3,4 Types 1 and 3 vWD are associated with relative or absolute quantitative defects in the protein, respectively. Type 1 vWD accounts for 70% of all cases and is likely to temporally improve in parturients as a result of an increase in vWF and factor VIII with pregnancy.2 Type 2 vWD has qualitative abnormalities and comprises 20–30% of all vWD diagnoses. A unique subtype, type 2B, accounts for less than 20% of all type 2 vWD and is characterized by an increased affinity of vWF for platelet glycoprotein Ib, resulting in spontaneous binding and clearance of both vWF and platelets.3,4 This feature allows type 2B, unlike most other vWD variants, to be exacerbated by pregnancy and to exhibit a poor or worsening response to desmopressin (DDAVP; Aventis, Bridgewater, NJ).5,6 We present the management of a parturient with type 2B vWD who had severe thrombocytopenia during the peripartum period.

Case Report

A 35-yr-old, 157-cm, 71-kg, uniparous woman, gravida 2 presented to our high-risk anesthesia consultation service at 36 weeks’ gestation. Although the patient reported no known drug allergies, the use of aspirin was associated with prolonged bleeding. Her history was notable for chronic hepatitis C, with no associated liver dysfunction, resulting from a blood transfusion and type 2B vWD, which was diagnosed as a child because of a strong family history (grandmother, mother, uncle, and brother) of type 2B vWD. Her obstetric history included a normal spontaneous vaginal delivery at 39 weeks’ gestation 2 yr previously without neuraxial analgesia. The platelet count at the time of her first delivery was 20,000 giga/l, and the patient was given antihemophilic factor (factor VIII) (Humate-P; Aventis, Bridgewater, NJ) 3,000 ristocetin cofactor units intravenously, and aminocaproic acid (Amicar; Immunex, Seattle, WA) 5 g per os loading dose followed by 3 g per os every 3 h.

The current pregnancy had a breech presentation, for which the patient was scheduled for a cesarean delivery. The platelet count at the time of consultation was 46,000 giga/l and the international normalized ratio 1.0; consultation with the hematologist included a plan to give factor VIII 4,000–4,500 ristocetin cofactor units followed by a unit of pheresis platelets starting 1 h before the procedure. On the day of the delivery, 2 weeks later, the initial platelet count was 56,000 giga/l and factor VIII and platelets were given. Within 30 min of the transfusion and before a planned spinal anesthetic technique, a repeat platelet count was drawn; the results demonstrated a platelet count of 30,000 giga/l. An additional unit of platelets was given in anticipation of bleeding during the cesarean delivery and the anesthetic plan was changed to a general anesthetic. Anesthesia was induced via rapid sequence with 250 mg sodium thiopental, 100 mg of succinylcholine, and 100 µg of fentanyl. A male infant with Apgar scores of 8 and 9 was delivered 3 min after the anesthetic induction. On closing the abdomen, no excessive intraoperative bleeding was observed, the total blood loss was estimated at 1,000 ml, and the patient had received a total of 2,800 ml of lactated Ringer’s solution. After tracheal extubation, an intraoperative sample sent for a platelet count was reported as 10,000 giga/l. The hematology consultant recommended additional factor VIII (5,000 units every 12 h for 3 days) to be commenced immediately. The obstetric team followed the recommendation and gave an additional unit of pheresis platelets as well. Six hours after delivery, serial platelet counts revealed progressive increases from 20,000 giga/l to 31,000 giga/l at 24 h, 42,000 giga/l at 48 h, and 55,000 giga/l at the time of discharge on postoperative day four. No postpartum hemorrhagic complications occurred and no additional thrombogenic products were needed. At time of hospital discharge, vWF ristocetin cofactor activity (vWF:RCO) and factor VIII concentrations were 45% and 258%, respectively.

Discussion

Previously defined by multimer analysis, the current classification scheme recognizes three general types of vWD based on molecular mechanisms.7 Types 1 and 3 represent partial and severe quantitative deficiencies of vWF and type 2 exhibits a qualitative deficiency. A characteristic gain-of-function phenotype, subtype 2B, may be the result of single amino acid substitutions within the A1 domain of vWF.4,8,9 These changes result in an increased interaction between vWF and the glycoprotein Ib-IX-V complex in platelets10 and an acceleration in binding and clearance of vWF and platelets; this interaction can be exacerbated by an increased production of vWF that can be produced with stressful clinical situations, of which pregnancy is well recognized.2,11,12 The resulting thrombocytopenia can be impressive, as observed in our case, and of sufficient concern as to allow

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some hematologists to support vWD testing for all women with thrombocytopenia in pregnancy.5

Neonatal thrombocytopenia may also occur with parturients with type 2B vWD,1,4 and in our case, the initial platelet count in the newborn was 35,000 giga/l. Antenatal testing for fetal vWD is possible but not often recommended; fetal morbidity and mortality may be associated with testing and the results are not always predictive of neonatal bleeding. The collection of cord blood at the time of delivery for vWD testing is preferable, and in our case, although the neonatal plasma vWF quantity was normal, the vWF activity and the amount of plasma factor VIII were markedly decreased. The possibility of neonatal vWD should not, however, prompt a cesarean delivery, as increased maternal bleeding may occur.13 Instead, a vaginal delivery with the avoidance of an episiotomy or instrumented delivery is preferable.2

The decision for cesarean delivery in our case was based on the breech presentation and an appropriate obstetric reluctance to perform either an external cephalic version or a vaginal breech delivery. All intramuscular injections were avoided in the mother and neonate, and a planned circumcision in the male infant was postponed.

Although desmopressin, a vasopressin analog that raises plasma vWF, is a common therapy for vWD, in type 2B vWD it can increase the abnormal vWF, resulting in further binding of platelets, depletion of high molecular weight multimers, and potentially greater bleeding.10,14–16 As such, the preferred therapy for this specific vWD subtype is factor VIII plasma concentrate (FVIII) (Humate-P or Alphanate; Grifols, Los Angeles, CA), which, although FDA approved for hemophilia and not vWD, provides the following corrective components: 2.5 IU vWF:RCo, 1 IU of FVIII, and a near-normal count of high molecular weight multimers.1,4,14,17,18 Threshold values of vWF:RCo and FVIII greater than 50 IU/dl at the time of delivery and for 72–96 h after an uncomplicated vaginal delivery and for 96–120 h after a cesarean section have been proposed.18

Although the treatment and goals for vWF:RCo and FVIII are relatively straightforward, the response to thrombocytopenia remains controversial. In part, this is because of the variation in thrombocytopenia observed in patients with type 2B vWD, which can be both intermittent and variable in expression, from mild to severe. In addition, the etiology of the thrombocytopenia is often unclear. Platelet aggregation has been confirmed by a number of investigations to occur spontaneously and with the use of desmopressin in patients with type 2 vWD.15,19 However, recent work suggests that desmopressin with this subtype may cause a “false” thrombocytopenia, resulting from short-term agglutination, but not aggregation.20 Moreover, the significance of the thrombocytopenia is unclear; some have suggested that the thrombocytopenia per se observed in type 2B vWD is not a major risk factor for bleeding, representing sequestration and margination of platelets, with later availability, rather than consumption.3

In our case, FVIII was given to increase the vWF:RCo and FVIII concentrations. The decision to transfuse platelets was prompted by the desire to have a platelet count greater than an arbitrary threshold of 50 giga/l for the spinal anesthesia and cesarean section. Although these platelets may have provided additional substrate for further aggregation and consumption of both vWF and platelets,4 we hypothesized that the degree of thrombocytopenia would be further accentuated by the surgical stress. Moreover, therapy with platelet transfusions has been found useful in some patients with type 2B vWD.15 Whether FVIII given simultaneously with platelet transfusions is more efficacious in treating type 2B vWD induced thrombocytopenia is currently unknown. What our case demonstrates, however, is that the quantitative response to FVIII and platelet transfusions is slow.

Ultimately, our decision to proceed with general anesthesia was based on a risk-benefit analysis, weighing the risk of neuraxial bleeding with a regional anesthetic versus the more common failed airway scenario observed in the parturient.21–25 Although difficult airways can not always be identified a priori,24,25 especially during pregnancy,22,26 we elected the general anesthetic based on our patient having a Mallampati class 2 airway, a wide mouth opening, a 5 cm thyromental distance, and a reasonable body mass index. If a more challenging airway had been observed, a regional anesthetic would have been more strongly considered, however, not before adequate vWF:RCo, FVIII, and platelet concentrations were obtained. Had our parturient presented in labor with a vertex presentation, intravenous patient-controlled analgesia with opioids would have been offered; were the desire for additional analgesia requested and hematologic goals achieved, a regional labor analgesic would have been considered. Despite these care algorithms, which were consistent with the desire of the patient to remain awake during her cesarean delivery, the falling platelet concentration despite hematologic intervention resulted in a general anesthetic being used.

This report emphasizes the severity of thrombocytopenia that can result in parturients with type 2B vWD. Early consultation with all involved healthcare providers, including obstetricians, hematologists, and anesthesiologists, and appropriate considerations should be made. Experience with this subtype of vWD suggests a correlation between vWF:RCo and FVIII concentrations and normal hemostasis exists,1 and therapy should be directed to these endpoints. The implications and treatment of thrombocytopenia deserve further investigation. Consideration of neuraxial analgesic or anesthetic techniques in these patients should weigh the risks and benefits of various approaches and the observation that certain therapies for type 2B vWD may be ineffective or slow.
**Spinal Epidural Hematoma after Spinal Anesthesia in a Patient Treated with Clopidogrel and Enoxaparin**

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IN recent years newly developed antiplatelet drugs such as the thienopyridine derivatives ticlopidine and clopidogrel are being increasingly used in patients with cardiovascular disease. Knowledge of their pharmacokinetics is crucial for appropriate perioperative care. Therefore, updated guidelines for the concomitant use of these drugs and locoregional anesthesia have been issued by the American Society of Regional Anesthesia and Pain Medicine and by European societies of anesthesiologists concerning a risk-benefit ratio as well as by European societies of anesthesia.

### References


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disease grade II (Canadian Cardiovascular Society angina classification) with atrial fibrillation, compensated renal insufficiency (creatinine clearance 34 ml/min), hypertension, and insulin-dependent diabetes. Her oral medication included molsidomine (Corvaton™; Aventis Pharma Deutschland GmbH, Bad Soden, Germany), piretanide/ramipril (Arelix ACE™; Aventis Pharma Deutschland GmbH), spironolactone/furosemide (Furorese™; Hexal AG, Holzkirchen, Germany), metoprolol (Beloc zok™; AstraZeneca GmbH, Wedel, Germany), and xipamid (Aquaphor™; Lilly Deutschland GmbH, Bad Homburg, Germany) for treatment of hypertension, ischemic heart disease, and renal insufficiency. Diabetes was treated with subcutaneous insulin. In addition, she was treated daily with 75 mg clopidogrel (Iscover™; Bristol-Myers Squibb GmbH, München, Germany) for ischemic heart disease which was discontinued 7 days before surgery.

Preoperatively the activated partial thromboplastin time was 39 s (normal range, 30–40 s), the prothrombin time was 85% (normal range, 70–120%) and the international normalized ratio was 1.02 (therapeutic level, 2–4.5), platelet count was 161 × 10^9/l, at the lower limit of the normal range (150 × 10^9/l – 400 × 10^9/l).

Spinal anesthesia was performed by an experienced staff anesthesiologist using a 22-gauge Sprotte needle. However, the first attempt failed at L3–4 because of repeated bone contact as well as a sanguine puncture presumed to be outside the epidural space. At the L4–5 level subarachnoid puncture was performed uneventfully in the first attempt and 3 ml of plain bupivacaine 0.5% was injected. Within 10 min maximum level of sensory block reached the T8 dermatome. Surgery was performed uneventfully and the regression of the block was complete within 4 h. The patient was mobilized on the evening of the day of surgery. For prophylaxis of thromboembolism the patient received two doses of low molecular weight heparin (enoxaparin) 40 mg 8 and 36 h after lumbar puncture. During the second postoperative night, 4 h after the second enoxaparin administration, the patient complained of voiding difficulty, and a urinary catheter was applied. At that time neither sensory nor motor deficiency was observed. On the morning of the second postoperative day the patient complained of numbness and weakness in both lower limbs. No back pain was reported. Emergency magnetic resonance imaging revealed spinal epidural hematoma extending from T12 to L3, with a maximum diameter between T12 and L1 (figs. 1–3). The patient was transferred to the university hospital and presented with lower paraplegia and a Th12 sensory loss. Emergency decompressive laminectomy was performed 20 h after the first signs of bladder dysfunction were observed. Before the second surgical procedure coagulation profile was documented as activated partial thromboplastin time 44, prothrombin time 81, international normalized ratio 1.22, and a platelet count of 216 × 10^9/l. The operative site showed a partially liquid and partially organized hematoma, which was difficult to resect as organized parts showed strong adhesion to the conus and seemed to obstruct epidural veins. No lesions related to puncture or vessel malformations were seen.

Postoperatively, the patient could partially be mobilized and only slowly recovered from her neurologic deficiencies. Three weeks after laminectomy sensory deficiency up to T12 and motor block Bromage 2 were documented.

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Fig. 1. T2 sagittal magnetic resonance image showing the large extension of the hematoma (arrows), as well as its heterogeneity.

Fig. 2. T2 axial magnetic resonance image showing the hematoma (L2).

Fig. 3. T1 axial magnetic resonance image showing the hematoma (L2).
Discussion

Beneficial effects have been demonstrated for new antiplatelet drugs in cardiovascular patients, especially for thienopyridine derivatives in preventing stroke. As an increasing number of patients is treated with these drugs, anesthetists must be familiar with their mechanism of action. In Europe two orally applicable adenosine diphosphate receptor antagonists are available that differ in clinical pharmacology. Both drugs inhibit platelet aggregation by noncompetitive irreversible interaction with the adenosine diphosphate (P2Y12) receptor on the outer platelet surface. Blockade leads to an inhibition of adenosine diphosphate mediated platelet activation via an increase in intracellular cyclic adenine monophosphate. In addition, release of calcium, fibrinogen, and serotonin that usually boost platelet activation is averted. Concurrently, alteration of GP IIb/IIIa conformation is inhibited.

Clopidogrel is six times more potent than ticlopidine and is therefore effective within 3–7 (ticlopidine, 8–10) days after its application. Normalization of platelet function is restored accordingly to the half-life time of platelets within 7 days (ticlopidine, 7–14 days). An animal model high-dose aprotinin partially reversed the effects of clopidogrel. Platelet transfusion is the only effective treatment in cases of severe bleeding in humans.

For clopidogrel a therapy-free interval of 7 days is recommended before performance of neuraxial blocks. In our patient 7 days, as recommended, were maintained and no sustaining clopidogrel impact was suspected.

Nevertheless, an epidural hematoma developed after spinal anesthesia. No case of epidural hematoma occurrence after central neuraxial block performance has been previously reported in patients shortly after a 7 day clopidogrel therapy-free interval. The first sign of epidural bleeding was bladder dysfunction, which was not considered unusual by the nursing staff at that time after spinal anesthesia in a patient of that age. Onset of sensory and motor deficiencies were delayed by approximately 12 h. Once discovered, magnetic resonance imaging was performed immediately and the patient was transferred to our hospital for laminectomy.

We consider four points to be of possible importance in this case:

• First, sanguine puncture occurred, which has been shown to be a general risk factor in development of spinal hematoma. Even though in the first attempt it was presumed that the epidural space was not reached, puncture of an epidural vein can not be excluded.

• Second, as the platelet count was at the lower limit of normal range, platelet regeneration may have been reduced, hence prolonging the effect of the adenosine diphosphate antagonist.

• Third, the exact response to subcutaneous low molecular weight heparin therapy in our patient remains speculative, as subcutaneous blood flow may vary considerably interindividually in the elderly and may result in therapeutic anticoagulation. A further risk factor might have been the moderate impairment of renal function. A linear correlation between anti-factor Xa concentrations and renal function has recently been demonstrated in patients receiving low molecular weight heparin, and the authors concluded that a dose adjustment is necessary in patients with a creatinine clearance of less than 30 ml/min. In an older retrospective study even clearance rates lower than 40 ml/min were associated with higher anti-factor Xa concentrations. In our patient the clearance was slightly greater than 30 ml/min.

• Finally, a 7–10 day therapy-free interval before operative procedures is recommended according to some manufacturer guidelines. The additive effect of anticoagulatory drugs acting at different sites of the coagulatory cascade should not be underestimated.

In summary, several factors may have equally contributed to the occurrence of an epidural hematoma in our patient. For thienopyridine derivatives clinical possibilities of testing thrombocyte function are limited. This case further underscores the necessity of close neurologic monitoring after spinal anesthesia, especially in applying locoregional anesthetic techniques shortly after the recommended therapy-free interval of clopidogrel.

References

Transient Profound Neurologic Deficit Associated with Thoracic Epidural Analgesia in an Elderly Patient

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PERIOPERATIVE epidural analgesia has been reported to improve patient outcome after thoracic surgery. Unfortunately, this procedure is not devoid of inherent potential risk. Specifically, numerous case reports describing neurologic deficits resulting from direct needle trauma or epidural mass effect (e.g., hematoma or abscess) have been published.1–3 We report a unique case of transient, yet profound, neurologic deficit that resulted from an unusual epidural mass.

Case Report

A 91 yr-old, 152-cm, 52-kg female was scheduled to undergo elective thoracotomy for repair of a symptomatic type III paraesophageal hiatal hernia using the Belsey Mark intravaneous procedure. Her past medical history was remarkable for well-controlled hypertension, primary cancer of the breast, colon, and bladder (all treated without recurrence), chronic anemia, glaucoma, and profound kyphosis (Cobb angle 72°) secondary to osteoporosis. Medications included ranitidine, raloxifene, latanoprost, carbidopa, levodopa, carisoprodol, vitamin B12, iron supplements, latanoprost, and timolol maleate.

Two hours before surgery she received 5000 U unfractionated heparin subcutaneously for deep venous thrombosis prophylaxis. On arrival in the operating room, sterile placement of a T6–7 epidural catheter was attempted unsuccessfully. The patient was repositioned, and catheter placement attempted at the T7–8 interspace resulted in dural puncture. A third attempt at the T8–9 level proved successful, as there was a discrete identification of the epidural space (i.e., loss-of-resistance) and the catheter was advanced 4 cm without difficulty. Catheter aspiration was negative for blood, and no discernible neurologic or hemodynamic changes resulted from a 3-ml test dose of lidocaine 1.5% with 1:200,000 epinephrine. Subsequently, the patient was placed supine, general anesthesia was induced, and the epidural catheter was loaded with 4 ml bupivacaine 0.25% plus 0.4 mg hydromorphone. Shortly thereafter, an epidural infusion of 0.075% bupivacaine with hydromorphone 5 μg/ml was initiated (and maintained postoperatively) at 6 ml/h.

Postoperatively, she reported good analgesia from the time of arrival in the postanesthesia care unit through the morning hours of the second postoperative day. During the afternoon of postoperative day 2, she developed profound bilateral lower extremity weakness. She denied back pain and was noted to be afibrile. Physical examination confirmed complete bilateral lower extremity paralysis with areflexia and absence of bilateral great toe proprioception. Light touch and temperature sensations were also diminished. Of note, she had received a total volume of approximately 220 ml of epidural infusate since the time of catheter placement. After confirming her coagulation status was normal, the epidural catheter was promptly removed, and emergent neurosurgical consultation and magnetic resonance imaging were obtained. Magnetic resonance imaging revealed an epidural mass in the dorsal left lateral aspect of the spinal canal extending from T6 to T9 with anterolateral spinal cord displacement and compression (fig. 1). Interestingly, the mass was initially thought to be an epidural hematoma. However, upon closer scrutiny of the magnetic resonance scan with our neuroradiologist (Dr. Rydberg), these images revealed features that were more characteristic of a nonheme-containing fluid. Specifically, the T2-weighted image intensity of the mass was consistent with a high water-content containing fluid (e.g., cerebrospinal fluid, local anesthetic, or opioid) with only a scant amount of blood (fig. 1).

The duration from discontinuation of the epidural infusion and completion of magnetic resonance imaging was approximately 90 min. During this time, the patient gradually regained lower extremity motor function. Within 2 h of discontinuing the epidural infusion, the neurologic deficits completely resolved, thereby circumventing the need for emergent surgical decompression. The remainder of her hospital stay was uneventful and she was discharged to an acute rehabilitation facility on postoperative day 7 with no adverse sequelae.

Discussion

We report a unique case of transient profound neurologic deficit in an elderly patient. To our knowledge, this is the first reported case of analgesic infusate and epidural cerebrospinal fluid causing clinically significant spinal cord compression.

Complications of epidural anesthesia include but are not limited to back pain, postdural puncture headache, infection, intravascular injection, inadvertent subdural or intrathecal injection resulting in high neuraxial blockade, arachnoiditis, epidural hematoma, anterior spinal artery thrombosis, and transient or persistent neurologic...
injury. With regard to neurologic injury, spinal cord damage may result from direct needle trauma, epidural hematoma formation, epidural abscess, arachnoiditis, or compromised vascular supply (e.g., injury or spasm of the spinal arteries). We believe the etiology of our observation was multifactorial. Specifically, dural puncture likely caused extravasation of cerebrospinal fluid into the epidural space. More importantly, the epidural infusate continuously added further volume to this expanding mass, which was particularly problematic because of the advanced age of the patient and severe kyphosis.

In the setting of normal neuraxial anatomy, epidural fluid is readily redistributed longitudinally along the craniocaudal axis, laterally towards the paravertebral region, or anteriorly. In elderly patients or patients with abnormal anatomy (i.e., severe kyphosis), dispersion of the fluid longitudinally and laterally may be limited, thereby resulting in anterior dispersion (i.e., the pathway offering minimal tissue resistance). In our patient, this resulted in mass effect and spinal cord compression. Motor deficit without sensory deficit likely resulted from disruption of anterior horn cells as the spinal cord was compressed against the thoracic vertebral bodies.

When faced with new onset neurologic deficits in patients receiving epidural analgesia, a neuraxial mass must be expeditiously excluded from the differential diagnosis. In addition to making a prompt diagnosis, it has been suggested that surgical decompression should be performed within 8 h of neurologic deficit onset for complete recovery to occur.

In summary, we report a unique case of transient profound neurologic deficit resulting from accumulation of epidural analgesic infusate and epidural cerebral spinal fluid. In our case, we observed early recognition of the epidural mass and discontinuation of the epidural infusion circumvented surgical intervention yet resulted in complete resolution of the profound deficit.

References


Fig. 1. T2 weighted sagittal (a) and axial (b) magnetic resonance images depicting a large epidural mass (arrows) in the dorsal left lateral aspect of the spinal canal extending from T6 to T9 with anterolateral spinal cord displacement and compression (chevron). The epidural mass was predominantly clear fluid (i.e., cerebrospinal fluid and analgesic infusate; large arrows) with only a scant amount of blood (small arrow).
POSTHERPETIC neuralgia (PHN) is one of the most difficult problems encountered by physicians. Sympathetic nerve blockade has been used with variable success as a component of therapy for PHN in cranial, cervical, thoracic, and lumbar distributions. Sacral dermatomal involvement of PHN in the sacral distribution was made, and the patient was treated with Neurontin and tramadol with limited improvement in symptoms. She was referred to our clinic for further evaluation. Initial physical examination revealed that she sat on one hip on a padded cushion as a result of her discomfort. Examination of the genital and rectal areas revealed no visible abnormalities. She was noted to have allodynia of the external genitalia and in the perirectal area. There were no areas of anesthesia noted. She described her pain as sharp and burning and noted painful bowel movements as well as dyspareunia. She initially received a caudal epidural steroid injection, which gave some mild relief of her symptoms near the coccyx but no relief of her vaginal and rectal pain. On her follow-up visit, she was treated with blockade of the ganglion impar using 12 ml of 0.25% bupivacaine and 40 mg of triamcinolone using a paramedial approach and 22-gauge 3.5-inch spinal needle (Becton-Dickinson, Franklin Lakes, NJ) prepared with 2000 units of neostigmine. † Consultant, Pacific Care Institute.

Discussion

Varicella zoster virus, a member of the herpes virus family, is a significant cause of morbidity. The incidence of Varicella (chickenpox) is over 4 million cases per year whereas the incidence of zoster is 300,000 cases per year. Primary infection with Varicella zoster virus results in Varicella (chickenpox); this is followed by a latent phase where the virus remains sequestered in dorsal root ganglia or the trigeminal ganglion. The virus usually remains dormant; however, it may be reactivated, leading to herpes zoster (shingles). Reactivation may be triggered by infection, immunosuppression, trauma, irradiation, malignancy, and advancing age. Herpes zoster is characterized as severe pain in one or more dermatomes usually followed by a vesicular skin eruption 7 to 10 days later. Persistence of painful symptoms for 4–6 weeks after the termination of the acute phase of infection is referred to as PHN. Prevalence of PHN is 9–45% of all cases of herpes zoster. Early institution of sympathetic blockade has been advocated to decrease the incidence and severity of PHN. This therapy remains controversial because a direct correlation...
between sympathetic blockade and prevention of PHN has not been proven by randomized prospective trials.

The ganglion impar is a solitary retroperitoneal structure. It is the fused terminal end of the paravertebral sympathetic chain and is found at the level of the sacrococcygeal junction (fig. 2). Neural blockade of the ganglion impar has been advocated as a means of managing intractable pelvic and perineal pain in cancer patients. A review of the literature reveals very limited information on the usefulness of blockade of the ganglion impar for pain other than that of neoplastic origin.

The original technique for blockade of the ganglion impar described by Plancarte and Valazquez involves a midline approach through the anococcygeal ligament with advancement of the needle tip cephalad to the anterior surface of the sacrococcygeal ligament. This technique involves placing one or two bends near the tip of the needle to more easily approximate the location of the ganglion. However, a gently curved needle has also been advocated to negotiate around the natural curve of the coccyx. Insertion of a finger in the rectum has not been proven by randomized prospective trials.

The pain caused by postherpetic neuralgia may involve both peripheral and central mechanisms. This understanding likely explains why postherpetic neuralgia is so difficult to treat effectively and also stresses the importance of multiple modality therapy.

The role of corticosteroids in pain relief for this patient is uncertain. Corticosteroid injections with nerve blocks have had unpredictable and limited success in treatment of PHN. Perhaps if further treatment is required for this patient, blockade of the ganglion impar with local anesthetic and no corticosteroid would offer some clarification of the role of corticosteroid in this instance.

Sacral dermatomal distribution of zoster and PHN is unusual but can lead to severe and disabling pain. Blockade of the ganglion impar may play a significant role in treatment of this disabling pain; however, the dramatic results in this patient should be interpreted cautiously. Further investigation is needed to determine efficacy of this sympathetic block in the treatment of the pain of zoster and PHN in the sacral dermatomes. Because of the low incidence of complications with blockade of the ganglion impar, it may have a role in early aggressive therapy of zoster in the sacral dermatomes.

Fig. 2. Needle placement for the paramedial approach to the ganglion impar.

References


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