The Effect of Bupivacaine Scalp Infiltration on the Hemodynamic Response to Craniotomy under General Anesthesia


We have observed that skin incision for elective craniotomy is frequently accompanied by tachycardia and arterial hypertension, despite an apparently adequate prior depth of general anesthesia (GA). The potential for morbidity is high in patients with intracranial pathology where hypertension may cause significant elevation of intracranial pressure¹ or precipitate intracranial hemorrhage, particularly in the presence of vascular tumors or cerebrovascular anomalies, such as aneurysms or malformations.² The problem may be overcome by increasing anesthetic depth or by administration of antihypertensive drugs. Because these treatments take time to initiate and risk hypotension, we sought to prevent or attenuate the hemodynamic response by infiltration of the scalp with local anesthetic prior to skin incision. As the practice had not previously been systematically evaluated, we compared the effect of scalp infiltration with 0.5% bupivacaine (without vasoconstrictor) that of normal saline in a double blind randomized fashion.

MATERIAL AND METHODS

Forty-two patients presenting for elective craniotomy for tumor within the cranial vault were studied, as approved by our local committee. Patients with unstable hypertension, receiving beta adrenergic blocking drugs, or with previous scalp incision were excluded from the study.

Forty-two pairs of 20-ml sterile glass ampules were prepared by the hospital pharmacy, 21 pairs containing normal saline and 21 pairs containing bupivacaine 0.5% (without vasoconstrictor). The ampules were identical in outward appearance, except for a code number which identified the contents. The availability of two vials for each code number allowed for breakage, the need for more than 20 ml for infiltration, and the need to repeat studies in the event of technical difficulties with recording equipment. The code was kept by the department secretary, and was not broken until all the data were collected and tabulated.

Each patient was premedicated with Temazepam, an oral benzodiazepine derivative, 20 mg, 1–1.5 h prior to arrival in the operating room. Prior to induction of GA, the radial artery was cannulated under local anesthesia. An electrocardiogram (ECG) monitor was attached. Arterial blood pressure was measured with a pressure transducer. ECG, heart rate (HR), mean arterial pressure (MAP), and arterial pressure wave form were recorded continuously from at least 3 min prior to induction of anesthesia until dural incision was complete. The times of induction of GA, endotracheal intubation, scalp infiltration, scalp incision, scalp flap reflection, craniotomy, and dural opening were noted on the recordings.

Induction of GA was accomplished with thiopental 5 mg/kg and phenoperidine, a potent narcotic analgesic, 0.04 mg/kg iv. Succinylcholine 1.2 mg/kg was administered iv to facilitate endotracheal intubation. Anesthesia was maintained with nitrous oxide 67%, oxygen 33%, and d-tubocurarine 0.4 mg/kg. Ventilation was controlled to maintain end tidal \( P_{CO2} \) at 30–35 mmHg.

Infiltration of the scalp was performed by the surgeon after skin preparation, prior to draping the patient with the intention of allowing a minimum of 5 min to elapse before incision. Both the incision line and the proposed line of scalp flap reflection were infiltrated. Sustained severe hypertension (>20% rise in MAP above the pre-induction levels) was treated with additional thiopental, phenoperidine, or hydradrazine, iv according to the discretion of the anesthetist. Such treatments were noted on the recordings.

Eight events were identified for the purpose of comparison: 1) pre-induction of GA, 2) post-endotracheal intubation, 3) pre-scalp infiltration, 4) post-scalp infiltration, 5) post-scalp incision, 6) scalp flap reflection, 7) craniotomy, and 8) dural incision. Mean HR and MAP data for each of these events was extracted from the polygraph tracings, using a digital plotter and micro-
Fig. 1. Mean arterial blood pressure (mmHg) from induction of general anesthesia to dural opening in the bupivacaine and saline (control) groups. Bars indicate ± one standard deviation. The statistical significance of the differences between each group at each event are displayed along the bottom (NS = not significant). Events: 1 = pre-induction of general anesthesia; 2 = post-endotracheal intubation; 3 = pre-scalp infiltration; 4 = post-scalp infiltration; 5 = post-scalp incision; 6 = post-scalp flap reflection; 7 = craniotomy; 8 = dural opening.

computer program to average the data over the period of extraction. This period was between 1 and 3 min (2.2 ± 0.8 min, mean ± SD) to ensure that it truly characterized the levels at the time. To verify that the data extraction was accurate, it was also extracted manually on a separate occasion.

After data extraction and final tabulation, the code was broken and the two groups (saline [control] and bupivacaine) identified. Statistical comparison between the two groups at each event was made using an unpaired t test.

Fig. 2. Mean heart rate (min⁻¹) from induction of general anesthesia to dural opening in the bupivacaine and saline (control) groups. Bars indicate ± one standard deviation. The statistical significance of the differences between each group at each event are displayed along the bottom (NS = not significant). Events: 1 = pre-induction of general anesthesia; 2 = post-endotracheal intubation; 3 = pre-scalp infiltration; 4 = post-scalp infiltration; 5 = post-scalp incision; 6 = post-scalp flap reflection; 7 = craniotomy; 8 = dural opening.

RESULTS

The two groups were well matched for age, weight, sex, and type of craniotomy. The saline group had ten men and 11 women with mean age 55.5 ± 12.8 yr. The bupivacaine group had nine men and 12 women with mean age 55.4 ± 15.3 yr.

The mean volumes of saline and bupivacaine injected were 14.5 ± 2.5 ml and 16.4 ± 5 ml, respectively. The mean time between infiltration of the scalp and skin incision was 5.5 ± 1.9 min in the saline group and 6.3 ± 3.2 min in the bupivacaine group.

No significant difference in MAP or HR occurred between the groups from induction of anesthesia until scalp incision commenced (Figs. 1, 2). During scalp incision, a marked difference developed between the saline group, in whom hypertension and tachycardia occurred, and the bupivacaine group, where there was little change in these variables. The difference in MAP and HR persisted through scalp reflection. The difference in MAP remained apparent during craniotomy. It was not until dural opening that both the tachycardia and hypertension in the saline group had resolved and the data of the two groups converged. Eleven patients in the saline group required additional thiopental, phenoperidine, and, in two cases, hydralazine because of sustained severe hypertension. No patients in the bupivacaine group required this treatment. No clinically apparent adverse reactions attributable to systemic absorption of bupivacaine were noted.

DISCUSSION

The notable cardiovascular stability seen in the bupivacaine group confirmed our impression that local anesthetic scalp infiltration prior to craniotomy minimizes the hemodynamic responses to skin incision. Prevention of tachycardia and hypertension circumvents the need for increase in anesthetic depth or antihypertensive treatments which take time to initiate and may transiently produce hypotension.

The benefits of local anesthetic infiltration persisted beyond scalp incision to include the craniotomy itself. Furthermore, by the time of dural opening, the hypertension and tachycardia observed earlier in the saline group had resolved. These findings are attributable to the mode of innervation of the scalp and cranium and the effect of local anesthesia. Subcutaneous local anesthetic scalp infiltration blocks peripheral nerves supplying the skin, subcutaneous tissue, muscle, and periosteum of the outer table of the skull. The peristomeum of the inner table of the skull (dura mater) is innervated by nerves accompanying the meningeal arteries which are not blocked by this technique. However, the dura is relatively insensitive to painful stimuli, except in prox-
Clinical Reports

Intraoperative Cardiac Dysrhythmias in a Patient With Bulimic Anorexia Nervosa

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Anorexia nervosa is a syndrome characterized by extreme weight loss, distorted body image, and a fear of becoming obese. Bulimia, a distinct syndrome, is characterized by binge-eating episodes followed by self-induced vomiting, fasting, and the use of diuretics and/or laxatives.

Preoccupation with food is common to both syndromes. The potential for marked weight fluctuations exists with bulimia, and severe weight loss often occurs with anorexia nervosa. In addition, bulimic symptoms may be part of the anorexia nervosa syndrome. Anorexia and bulimia are estimated to effect 5–10% of adolescent girls and young women. We report a case of cardiac dysrhythmias during general anesthesia in a patient with bulimic anorexia nervosa.

CASE REPORT

A 16-yr-old girl presented in the ambulatory surgery unit for dental extractions. The patient had a 1-yr history of anorexia nervosa with

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Key words: Complications: dysrhythmias. Ions: potassium. Metabolism: anorexia nervosa; bulimia.