

Continuous spinal anaesthesia using a standard epidural set for extracorporeal shockwave lithotripsy

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Purpose: Continuous spinal anaesthesia (CSA) offers considerable advantages over "single shot" spinal or epidural anaesthesia since it allows titration of anaesthesia using small doses of local anaesthetics (LA). We evaluated the feasibility of CSA using a standard epidural set for extracorporeal shockwave lithotripsy (ESWL).

Methods: Charts of 100 consecutive CSAs for ESWL were retrospectively reviewed. Lumbar CSA was performed using a 20G epidural catheter through an 18G Tuohy needle. The CSA was preplanned, or followed inadvertent dural puncture. Small LA boluses were injected to achieve the desired sensory level of anaesthesia. Demographic data, anaesthetic duration, LA doses, the most cephalad sensory level to pinprick, arterial blood pressure, heart rate, use of systemic sympathomimetics and complications were recorded.

Results: Mean age was 66.2 ± 9.9 (SD). The ASA status was III-IV in 54.1% and 5.5% of the preplanned and inadvertent patients, respectively. In 85 anaesthetics, hyperbaric bupivacaine 0.1% (9.7 ± 7.5 mg) was used as the sole anaesthetic. Sensory level was T₄-T₈. Maximal decrease in systolic and diastolic blood pressures and heart rate was $19.0 \pm 9.8\%$, $13.4 \pm 13.3\%$, and 7.2 ± 11.7 respectively. Intravenous sympathomimetics were used in nine of 82 (11.0%) preplanned, and in six of 18 (33.3%) inadvertent anaesthetics. Post dural puncture headache appeared following two of 82 (2.5%) preplanned, and four of 18 (22.2%) inadvertent anaesthetics. No postanaesthetic neurological deficit was detected.

Conclusion: Continuous spinal anaesthesia, using a standard epidural set and hyperbaric bupivacaine is feasible for ESWL in high risk patients. Inadvertent dural puncture does not preclude CSA under these circumstances.

Objectif : Parce qu'elle permet de titrer l'anesthésie avec des doses minimales d'anesthésique local (AL), la rachianesthésie continue (RAC) offre des avantages considérables sur la rachianesthésie et l'épidurale à injection unique. Nous avons recherché s'il était possible d'administrer la RAC avec un plateau épidural standard pour la lithotripsie extracorporelle par ondes de choc (LEOC).

Méthodes : L'étude a consisté à revoir rétrospectivement les dossiers de 100 interventions de LEOC consécutives. La rachianesthésie lombaire était réalisée avec un cathéter épidural 20G introduit à travers une Tuohy 18G. La RAC faisait partie de la planification originale ou suivait une ponction accidentelle de la dure-mère. Le niveau sensoriel d'anesthésie désiré était atteint par l'injection de petits bolus d'AL. Les données démographiques, la durée de l'anesthésie, les doses d'AL, le niveau céphalique maximal déterminé par piqûre, la pression artérielle, la fréquence cardiaque, l'utilisation de sympathicomimétiques et les complications ont été enregistrées.

Résultats : L'âge moyen était $66,2 \pm 9,9$ (ÉT). Le pourcentage d'état physique ASA III-IV représentait 54,1% des patients prévus et de 5,5% chez les patients imprévus. Quatre-vingt-cinq patients n'ont reçu que de la bupivacaine hyperbare à 0,1% ($9,7 \pm 7,5$ mg). Le niveau sensoriel atteint se situait entre T₄ et T₈. Les chutes maximales des pressions systolique et diastolique et de la fréquence cardiaque étaient de $19 \pm 9,8\%$, $13,4 \pm 13,3\%$ et $7,2 \pm 11,7$ bpm. Il a fallu administrer des sympathicomimétiques intraveineux à 9 des 82 (11,0%) des patients prévus et à six des 18 (33,3%) des imprévus. Une céphalée postrachianesthésie s'est manifestée après 2 des 82 (2,5%) RAC prévues et quatre de 18 (22,2%) imprévues. Il n'y a pas eu de déficit neurologique.

Conclusion : Il est possible d'administrer une rachianesthésie continue avec un plateau épidural standard et de la bupivacaine hyperbare pour la LEOC chez des patients à haut risque. Une ponction accidentelle de la dure-mère n'exclut pas la RAC dans ces circonstances.

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CONTINUOUS spinal anaesthesia (CSA) offers considerable advantages over "single shot" spinal or epidural anaesthesia since it allows administration of well-controlled anaesthesia using small doses of local anaesthetics (LA).¹ Its use has been reported for various procedures such as Caesarean section,¹ orthopaedic,¹⁻³ trauma³ or peripheral vascular⁴ procedures in the lower limb, lower abdominal and pelvic surgery,^{1,5} urological procedures,¹ and for gynaecological, rectal and perineal surgery.¹ It has been used in old^{2,6} and high-risk^{1,2,4-6} patients. However, CSA has not gained wide popularity because of the fear of post anaesthetic cauda equina syndrome,^{3,6-9} and because of the difficulty in placing microcatheters into the subarachnoid space.^{3,7}

The patient population referred for extracorporeal shockwave lithotripsy (ESWL) usually includes elderly patients, many of whom suffer from coexisting systemic illness. Moreover, patient positioning and immersion in the water bath can lead to haemodynamic changes.¹⁰

In this study, we analysed retrospectively 100 consecutive CSAs for ESWL using a standard epidural set, performed in 92 patients.

Methods

Subjects: Of the 1146 ESWLs performed in our hospital, all with anaesthesia, over five years, 100 (8.7%) were performed with CSA.

Procedures: The CSAs were performed using a standard epidural set (Perifix, B. Braun Melsungen AG, Germany). The patients were positioned in the seated or the lateral decubitus position, and an 18G Tuohy needle was inserted into the subarachnoid space in the L₂-L₃, L₃-L₄, or L₄-L₅ intervertebral space, through which a standard 20G epidural catheter was introduced. The catheter was threaded 3-4 cm into the subarachnoid space. Paraesthesia were uncommon and, when elicited, the catheter was slightly withdrawn. Location of the needle and the catheter in the subarachnoid space was verified by obtaining a free flow of cerebrospinal fluid (CSF). When epidural anaesthesia was originally planned, but the dura was inadvertently penetrated, the catheter was inserted through the needle, and continuous spinal, rather than epidural anaesthesia was conducted. Local anaesthetics (e.g., 1-3 mg boluses of bupivacaine 0.1%) were injected into the subarachnoid space, until the desired sensory level of anaesthesia relative to the location of the stone, was reached. The selection of anaesthetic agent was according to the preference of the consultant anaesthetist. Routine monitoring included pulse oximetry, ECG and non-invasive (or invasive) mea-

surement of arterial blood pressure (BP). Usually, non-invasive BP was measured every 1-2.5 min until approximately 10 min after anaesthetic induction, and later, every 2.5-5 min until completion of the procedure. Sympathomimetic agents were administered *iv* when systolic BP decreased below 100 mmHg, or when the anaesthetist has considered that the value was too low (particularly in hypertensive patients). The patients were then transferred to the hydraulic chair of the lithotripter (HM3 Dornier, Munchen, Germany) and immersed into the warm water bath (about 36°C) where the shockwaves were delivered. After the procedure, the spinal catheter was withdrawn and the patients were transferred to the post anaesthesia care unit. For all patients, the following variables were recorded: age, weight, ASA physical status, duration of procedure, anaesthetic drugs and doses, the most cephalad sensory level to pinprick, the use of additional systemic sedatives and analgesics, baseline and minimal BP and heart rate (HR), the use of sympathomimetic drugs, volume of *iv* fluid, and peri-anaesthetic complications. The last included technical failure to thread the catheter, evidence of inadequate anaesthesia (either insufficient sensory level or quality of the block), "high" spinal anaesthesia, hypotension, bradycardia, post dural puncture headache (PDPH) and adverse neurological sequelae including persistent pain, motor or sensory deficits, haematoma or infection.

Results

Ninety-two patients underwent 100 anaesthetics for lithotripsy: 86 had one anaesthetic, five had two and one had four anaesthetics.

Of the 100 CSAs, 82 were preplanned (CSA-P), and 18 (in 18 patients), followed inadvertent dural puncture following attempted epidural blockade (CSA-I). Fifty-nine (64.1%) of patients were male. The mean age of the patients was 66.2 ± 9.9 (SD) yr and weight was 78.5 ± 13.9 kg. Forty-one patients (44.6%) were graded as ASA physical status III or IV. Of these, 40 were in the CSA-P group (54.1% of the 74 patients in the subgroup), as opposed to only one patient (5.5%) in the CSA-I group. Twenty-one patients had one concurrent disease, 21 had two, 18 had three, 13 had four, six had five, one had six concurrent diseases, and one patient had ten concurrent diseases (Table I). In 85 anaesthetics, hyperbaric, diluted bupivacaine was used as the sole intrathecal anaesthetic drug. It was prepared as a 0.1% solution by diluting plain 0.5% bupivacaine (Astra, Sodertalje, Sweden) in 10% glucose (prepared by our institutional pharmacy). The mean total administered dose of bupivacaine was 9.7 ± 7.5 (SD) mg. Lidocaine 2% was added intrathecally in two patients and used as the sole agent in one additional patient. Intrathecal fentanyl in doses of

TABLE I Pre-existing medical conditions in 92 patients

Condition	n (%)	Condition	n (%)
ischaemic heart disease	25 (27.2)	COPD,* bronchial asthma	9 (9.8)
valvular heart disease	3 (3.3)	pulmonary hypertension	1 (1.1)
dysrhythmia/conduction defect	12 (13.0)	diabetes mellitus	19 (20.7)
cardiomyopathy/cardiomegaly	7 (7.6)	metabolic/other hormonal†	7 (7.6)
congestive heart failure	7 (7.6)	obesity/morbid obesity	29 (31.5)
S/A coronary artery bypass	3 (3.3)	chronic renal failure	13 (14.1)
systemic hypertension	43 (46.7)	CNS disorders‡	8 (8.7)
smoking	14 (15.2)	others§	7 (7.6)

* COPD: Chronic obstructive pulmonary disease

† hyperlipidaemia, gout, hypothyroidism

‡ impaired level of consciousness, S/A stroke, myelodysplasia

§ peptic ulcer disease, anaemia, asymptomatic carotid arterial narrowing, polyarthralgia

10–20 µg was used in 12 patients anaesthetized with bupivacaine. The observed sensory level was T₄-T₈, and all patients achieved satisfactory anaesthesia.

Intravenous sedatives and analgesics (i.e., 1–5 mg midazolam, 1 mg droperidol, 25–100 µg fentanyl and 300–600 µg alfentanil), were used in 19, one, five, and three patients, respectively. Anaesthetic duration, fluid administration, the use of sympathomimetic drugs, the appearance of PDPH and the need for epidural blood patch are presented in Table II. Haemodynamic variables are presented in Table III.

Sympathomimetic drugs were needed for the treatment of hypotension in nine patients (11%) in the preplanned group and in six patients (33%) in the inadvertent group. Ephedrine was used in 12 patients (six in each group), at a dose of 5–25 mg in 5 mg increments. Five patients received 5 mg, one received 7.5 mg, three received 10 mg, and three received 15 mg, 20 mg

and 25 mg each. Phenylephrine was administered to two patients (CSA-P), at a dose of 100 and 150 µg, in 50 µg increments. Both drugs (5 mg and 50 µg) were used in one patient (CSA-P). Post dural puncture headache appeared after six CSAs, but in only two (2.5%) of the preplanned group. Four of these were male, and two female. The ages of these patients were 76, 70, 67, 65, 43 and 40 yr. One patient received an epidural “blood patch,” while another was treated with an epidural injection of morphine.¹¹ In both patients, treatment resulted in resolution of pain. In the other four patients, satisfactory pain relief was achieved by intravenous fluid loading and systemic analgesics. No postanaesthetic neurological deficit was detected in any of our patients.

Discussion

This study describes the use of continuous spinal anaesthesia with a standard 20G epidural catheter as an effective and safe anaesthetic approach for ESWL, and specifically in high risk patients.

Continuous spinal anaesthesia combines the advantages of both epidural and spinal anaesthesia: failure rate is very low as placement of the Tuohy needle in the subarachnoid space is easily ascertained by the escape of CSF.^{1,2,6,12} Onset of anaesthesia is easy to control and can be either gradual,^{1,12} or rapid,¹ depending on the sequence of injection of the LA. The administered dose of LA is small (9.7 ± 7.5 mg bupivacaine in our study), reducing the possibility of systemic toxicity if the catheter is accidentally inserted intravascularly.^{1,2} The sensory level of anaesthesia can be adjusted by careful titration,^{1,5,6} thus reducing the risk of haemodynamic instability.^{5,6} As the level of the block is established after patient positioning, haemodynamic^{1,2,5,12} and respiratory⁵ stability may be better preserved. Duration of the block is unlimited due to the possibility of repeated injections *via* the spinal catheter.^{1,12} With the use of low

TABLE II Clinical detail

Duration (min)*	92.1 ± 31.9
Fluids (ml)*	1330 ± 578
Sympathomimetics (n)	15 (15%)
PDPH (n)†	6 (6%)
Epidural blood patch (n)	1 (1%)

* Mean ± SD

† PDPH = Post dural puncture headache

TABLE III Haemodynamic variables

Baseline systolic BP (mmHg)	147.8 ± 22.9
Maximal systolic BP decrease (%)	19.0 ± 9.8
Baseline diastolic BP (mmHg)	84.5 ± 12.4
Maximal diastolic BP decrease (%)	13.4 ± 13.3
Baseline HR (beats per minute)	80.3 ± 13.7
Maximal HR decrease (%)	7.2 ± 11.7

BP: blood pressure, mean ± SD

HR: heart rate, mean ± SD

doses of LA, the recovery time is usually short.^{1,5,12} The deep motor and sensory blockade achieved provides excellent analgesia and tends to eliminate movement which can disturb calculi fragmentation.

All patients achieved satisfactory anaesthesia. This may be related to the injection of the LA into the subarachnoid space. However, since the use of microcatheters may be associated with technical failure¹⁻³ and with anaesthetic maldistribution^{3,7-9} our success may be related to the use of 20G catheters.

Hypotension is one of the disadvantages of "single shot" spinal anaesthesia^{13,14} which can cause considerable morbidity and mortality.¹³ It may be additive to the haemodynamic changes associated with ESWL, related to patients positioning and to the immersion in the water bath. The head-up positioning leads to peripheral pooling of blood and thereby to diminished venous return to the heart. This, however, may be offset by a possible increase in venous return due to immersion in water.¹⁰ In addition, the warm water in the bath may lead to vasodilatation, and therefore decreased blood pressure.

Prophylactic *iv* infusion of crystalloids, or ephedrine, failed to prevent severe hypotension after "single shot" spinal anaesthesia in 55% and 22% of patients, respectively.¹³ In contrast, although volume loading and sympathomimetics were not used prophylactically in our study, and despite the T₄-T₈ sensory level of anaesthesia, the average BP decrease was only 13.4-19.0% (12.9-14.0% in the CSA-P group). Moreover, administration of sympathomimetics for BP control was needed in only 15% of our anaesthetics, compared with 65% of the patients receiving "single shot" spinal anaesthesia for lower limb surgery in another study.²

We used 1-3 mg boluses of bupivacaine, comparable to the dosage range suggested by Sutter *et al.*² Gradual titration of LA according to clinical response allows the circulation to accommodate to the vasodilatation induced by the CSA, thus minimizing the decrease in BP. As long as the block is kept below the T₄ dermatome, cardioaccelerator sympathetic nerves can still react to hypotension. This enables the clinician to use CSA in patients that cannot tolerate wide fluctuations in BP, HR and/or rapid and massive volume loading. Several of the high risk patients in our study received < 500 ml crystalloid during the procedure, while still maintaining haemodynamic stability. However, fluid administration was not restricted, since high urinary output was encouraged to flush calculi fragments from the urinary system. Despite the use of the wide-bore 18G Tuohy needle and standard epidural catheter, there was a low incidence PDPH, and no neurological sequelae were encountered.

The use of microcatheters (i.e., 28-32G) for CSA, was introduced in an attempt to reduce PDPH,^{1,3,7} but it was found to be technically difficult to thread the microcatheter into the subarachnoid space despite good CSF flow through the needle.^{1,3,7} Using epidural sets, however, makes subarachnoid catheter insertion technically simple. The use of microcatheters was later abandoned following several reports of postanaesthetic cauda equina syndrome.^{3,6-9,15}

The incidence of PDPH, which decreases with patient age, but increases with needle diameter,¹⁶ was low (2.5%) in our CSA-P group despite the use of the 18G needle. This is in accordance with the 1% incidence found by others.^{4,17} The explanation for the reduced incidence of PDPH after CSA is not known. Perhaps an early inflammatory response develops in the dura surrounding the catheter at the puncture site, so that when the catheter is removed, fibrinous exudate and/or oedema seal the hole in the dura, thus preventing leakage of CSF.^{4,17} Adverse neurological sequelae after CSA may be related to the small diameter of microcatheters,^{3,8,9,15} or to a high LA dose^{8,18} or concentration.^{9,18} However, such a complication occurred even after CSA was performed with macrocatheters.¹⁹ Hence, it is possible that the lack of adverse neurological sequelae in our patients may be related to the relatively large diameter of the catheter, or to the low LA dose and concentration used. It is possible that such a complication could have been detected in a larger study population.

In summary, the results of this study suggest that CSA, utilizing standard 20G epidural catheters, can be performed effectively and safely in patients undergoing ESWL. Moreover, if the epidural needle is inadvertently introduced into the subarachnoid space during attempted epidural blockade, the catheter may then be intentionally inserted, and CSA carried out. In elderly patients for ESWL, however, and especially in high risk individuals, preplanned CSA is particularly desirable, since slow titration of LA to achieve the required sensory level seems to result in relative haemodynamic stability despite minimal use of pre-emptive volume-loading or cardiotoxic agents. Despite the use of a relatively large diameter needle, the incidence of post dural puncture headache is low.

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