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Management of hypertrophic pylorus stenosis with ultrasound guided single shot epidural anaesthesia – A retrospective analysis of 20 cases

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**Management of hypertrophic pylorus stenosis
with ultrasound guided single shot epidural
anaesthesia – A retrospective analysis of 20 cases**

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Epidural anaesthesia for hypertrophic pylorus stenosis

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None

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Summary

Aim: To retrospectively describe the performance of ultrasound guided thoracic epidural anaesthesia under sedation for anaesthesia management of open pyloromyotomy.

Background: Anaesthesia management for hypertrophic pylorus stenosis is usually performed under general anaesthesia with tracheal intubation. Only a few publications describe avoidance of tracheal intubation in infants by using spinal or caudal anaesthesia. The present retrospective analysis describes the performance of ultrasound guided thoracic epidural anaesthesia under sedation for anaesthetic management of open pyloromyotomy.

Methods: Twenty consecutive infants scheduled for pyloromyotomy according to the Weber-Ramstedt technique were retrospectively analysed. After sedation with nalbuphine and propofol, an ultrasound guided single shot thoracic epidural anaesthesia was performed with 0.75 ml kg⁻¹ ropivacaine 0.475%. Insufficient blockade was defined as increase of HR > 15% from initial value and / or any movements at skin incision. In those cases we were prepared for rapid sequence intubation according to the departmental standard.

Results: All pyloromyotomies could be performed under single shot thoracic epidural anaesthesia and sedation. One case of moderate oxygen desaturation was treated with intermittent ventilation via face mask.

Conclusions: Thoracic epidural anaesthesia under sedation for pyloromyotomy has been a useful technique in this retrospective series of infants suffering from hypertrophic pylorus stenosis. In 1/20 infants short term assisted ventilation via face mask was required. Undisturbed surgery was possible in all cases.

Keywords: Hypertrophic pylorus stenosis; thoracic epidural anaesthesia; ultrasound

Introduction

Hypertrophic pyloric stenosis (HPS) is a frequent disease in infants with an incidence of 0.9-5.1 per 1000 cases (1-4). The main symptoms of HPS are progressively worsening “projectile” vomiting, poor feeding and dehydration caused by a gastric outlet obstruction due to a hypertrophic pylorus. The average age and weight of infants with HPS is 5 weeks and 4 kg, respectively (5).

Anaesthesia management for HPS is usually performed under general anaesthesia with tracheal intubation. Tracheal intubation puts these infants at risk of regurgitation, with the potential of aspiration of gastric contents, and rapid sequence intubation is indicated. Beside the special character of anaesthesia induction in children with HPS, rapid sequence intubation in infants should be always considered as high risk procedure. Despite preoperative correction of acid-base balance and hypovolemia, prolonged mechanical ventilation might be required due to remaining metabolic alkalosis and a subsequent delayed equilibrium of the cerebrospinal fluid with the systemic circulation (6, 7).

Only a few publications describe avoidance of tracheal intubation and mechanical ventilation in infants undergoing surgery for treatment of HPS by using spinal (8, 9) or caudal anaesthesia (10, 11). The major drawback of high spinal anaesthesia is the unpredictable cranial subarachnoidal spread of local anaesthesia with subsequent respiratory failure. Caudal anaesthesia on the other hand might be insufficient for pyloromyotomy with a skin incision above the umbilicus.

Thoracic epidural anaesthesia might be an option for anaesthetic management of HPS. Until today, a lot of practitioners have concerns against thoracic epidural punctures in infants due to safety reasons. Recently our study group has developed a technique to directly observe the spread of local anaesthetic inside the epidural space in neonates and infants by ultrasound (12,

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3 13). Consequently, we used this technique for single shot thoracic epidural punctures in
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5 infants undergoing pyloromyotomy, and analysed the first 20 consecutive cases in a
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7 retrospective manner.
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Methods

We included 20 consecutive infants with HPS in this retrospective analysis. Parent’s informed consent included an exact description of the anaesthesia procedure (aspiration of gastric juice via a naso-gastric tube, sedation, ultrasound guided single shot epidural anaesthesia) and possible need for rapid sequence induction. After initial diagnosis of HPS by clinical status, ultrasound and blood gas analysis (BGA), 10 mL·kg⁻¹·h⁻¹ Elo-Paed balanced plus glucose 1% (Fresenius Kabi Inc., Graz, Austria) was administered via a peripheral venous access until a HCO₃ of ≤ 28 mmol l⁻¹ and a BE of < +2 was achieved.

Pre-epidural preparation

The epidural puncture site between the T10 and T11 vertebral levels was prepared with EMLA cream 30 min prior epidural puncture and infants were premedicated with midazolam 1 mg·kg⁻¹ via the rectal route. After transfer to the operation room, children were placed on a forced-air warming device (Bair Hugger warming blanket, Arizant Inc., Eden Prairie, MN, USA). Standard monitoring included ECG, SpO₂ and non-invasive blood pressure. Sedation was induced with nalbuphine 0.1 mg·kg⁻¹ and a loading dose of propofol 1.0-2.0 mg·kg⁻¹, administered over 30 seconds. If necessary, supplemental doses of propofol 0.5 mg·kg⁻¹ were administered until adequate sedation was achieved. Sedation was considered adequate, when the patient slept, arousable only with significant physical stimulation. This type of sedation was previously published by Machata et al. and Brenner et al. in 500 and 512 cases, respectively (14, 15).

Gentle aspiration of gastric juice via a naso-gastric tube was performed before initiation of the sedation procedure. The maintenance of spontaneous respiration was continuously verified by an end-tidal CO₂ line placed inside a face mask. Via this face mask oxygen / air (FiO₂ 50%) was administered. Infants received 10 mL·kg⁻¹·h⁻¹ Elo-Paed balanced plus glucose 1%.

Epidural puncture

The single shot epidural anaesthesia was performed under sterile conditions in left lateral position between the T10 and T11 vertebral spaces. The neuraxial structures were directly visualized with a sterile covered 38 mm 13-6 MHz linear ultrasound probe and a transportable ultrasound machine (M-Turbo, SonoSite Inc., Bothell, WA, USA) from paramedian. Once the dura mater and the epidural space were identified, the puncture was performed with a 20G, 50 mm Tuohy needle and an 8 ml loss-of-resistance (LOR) syringe (BBraun Inc., Melsungen, Germany) via a median approach (Figure 1) using ropivacaine 0.475%. A total volume of $0.75 \text{ ml} \cdot \text{kg}^{-1}$ ropivacaine 0.475% ($= 3.56 \text{ mg} \cdot \text{kg}^{-1}$) was administered under ultrasound observation of the spread of local anaesthetic (Figure 2). After performance of the epidural blockade the children were turned in supine position.

Surgical procedure

Fifteen minutes after performance of the block, skin incision was performed via a right lateral horizontal approach, according to the Weber-Ramstedt technique (16). After pyloromyotomy, saline was administered in the surgical wound, and a moderate volume of air was insufflated via the gastric tube to exclude accidental perforation of the pylorus.

Emergency management

All equipment for advanced airway management was prepared in cases of respiratory failure. Respiratory failure was defined as the development of paradox ventilation, disappearance of end-tidal CO_2 curve and / or decrease in $\text{SpO}_2 < 92\%$. The following sequential airway management was initiated to re-establish adequate oxygenation:

- careful ventilation via face mask with inspiratory pressure < 10 mmHg
- rapid sequence intubation according to the departmental standard (propofol 8.0 mg.kg⁻¹, rocuronium 0.6 mg.kg⁻¹).

Definitions for bradycardia and hypotension were a decrease in heart rate and MAP > 25% from initial values and treated with atropine 0.01 mg.kg⁻¹ and a fluid bolus of 10 ml.kg⁻¹, respectively.

Insufficient blockade was defined as increase of HR > 15% from initial value and / or any movements at skin incision. In those cases we were prepared for rapid sequence intubation according to the departmental standard.

Postoperative management

After transfer to the recovery room, pain status of the children was monitored via OPS score, in which objective behavioural variables (crying, facial expression, position of torso and legs, motor restlessness) are assessed. Each pain variable is scored on a three-point scale (0 = none, 1 = moderate, 2 = severe) to give a maximum cumulative score of 10. The scores were evaluated after admission in the recovery room and every 30 min during the first 2 postoperative hours. If the OPS score was ≥ 6 in two subsequent measurements, the child received acetaminophen 40 mg.kg⁻¹ rectally. Due to the retrospective nature of this study no OPS scores could be evaluated on the ward.

Postoperative nutrition was performed by ad libitum feeding (17). The epidural puncture site was examined 24 h postoperatively to detect local infection, according to the departmental standard.

Results

We analyzed the first 20 consecutive infants undergoing pyloromyotomy according to Weber-Ramstedt with single shot epidural anaesthesia. Pertinent patient data are illustrated in Table 1. The relevant blood gas values after admission in the hospital and before surgery are illustrated in Table 2.

Anaesthesia management via sedation (details are described in Table 2) and ultrasound guided single shot thoracic epidural blockade was successful in all infants. Thus, no rapid sequence intubation as described in the methods section and reversal of neuromuscular blockade was required.

As expected, median (range) decrease in heart rate after administration of 0.75 ml kg^{-1} ropivacaine 0.475% was 18% (5-30%) and remained stable on post-epidural lower level during the entire surgical procedure (Figure 3). Oxygen saturation remained stable between 97 and 100% in all cases throughout the entire anaesthesia and surgical procedure, except in one case where SpO_2 decreased to 92% 10 min after epidural anaesthesia. Treatment of this short episode of decrease in SpO_2 was performed by assisted positive pressure ventilation via face mask.

All OPS scores remained < 5 and therefore no child received additional systemic pain therapy in the recovery room. The examination of the epidural puncture site 24 hours postoperatively was uneventful in all cases.

Discussion

This consecutive case series describes a novel anaesthesia management for pyloromyotomy in infants suffering from HPS. Ultrasound guided single shot thoracic epidural anaesthesia under sedation and spontaneous respiration has been a useful technique in this retrospective series of infants for pyloromyotomy. In this particular retrospective study of 20 cases, airway manipulation and mechanical ventilation could be avoided.

Pyloromyotomy is usually performed under general anaesthesia, thus requiring tracheal intubation and rapid sequence induction of general anaesthesia (5). The use of regional anaesthesia techniques depends on the exact site of surgery. As most surgical techniques for pyloromyotomy require a supraumbilical skin incision, spinal and caudal blockade seem inadequate. Kachko et al. suggest spinal anaesthesia only for low abdominal procedures (8), whereas Somri et al. and Jetzek-Zader et al. describe high spinal blockade with bupivacaine 0.5% (0.8 mg·kg⁻¹ and 1.3 mg·kg⁻¹, respectively) as possible regional anaesthetic technique and as an alternative to general anaesthesia for pyloromyotomy (9, 18). Moyao-Garcia et al. suggest caudal blockade with bupivacaine 0.25% and a volume of 1.6 ml for pyloromyotomy and describe a success rate of 96% (11). In spite of all these encouraging reports, our clinical experience was based on the observation that caudal blockade is insufficient for Weber-Ramstedt repair of HPS due to a required cranial analgesic level between T4 and 6.

Preliminary and unpublished data show that only in the minority of cases a spread above T12 can be achieved via the caudal approach, even with a volume of local anaesthetic of 1.5 mg·kg⁻¹. On the other hand, spinal anaesthesia for pyloromyotomy may cause uncontrolled high blockade and subsequent respiratory insufficiency.

During the past 10 years our study group acquired a substantial experience in the area of central (12, 13, 19) regional anaesthetic techniques with ultrasound guidance. In the light of

the findings above and our own significant experience with epidural anaesthesia in infants, we considered ultrasound guided single shot thoracic epidural blockade with sedation as a possible alternative to other techniques. The analysis of our first consecutive cases showed that thoracic epidural blockade under ultrasound guidance is a useful anaesthesia method for Weber-Ramstedt pyloromyotomy. Once the epidural space is identified via a combination of LOR and direct visualization, the spread of local anaesthetic can be directly observed.

We administered from the first case ropivacaine 0.475% (1:1 mixture of ropivacaine 0.75 % and 0.2 %) with a volume of $0.75 \text{ ml} \cdot \text{kg}^{-1}$ and observed an adequate cranial spread of local anaesthetic with no alterations in ventilation. This volume and concentration may be considered as relatively large, but pharmacokinetics of epidural ropivacaine in infants is insufficiently described. Anyway, systemic resorption of ropivacaine from the epidural space seems to be slower as compared with bupivacaine, thus increasing the safety of epidural ropivacaine in infants (20). Due to the fact that no data existed regarding optimal volume and concentration of local anaesthetic for this particular indication, we have been prepared for alterations in volume and concentration of local anaesthetic. Fortunately the initial choice of volume and concentration of local anaesthetic was effective and seems to be safe regarding spontaneous ventilation. Anyway, further studies should investigate lower concentrations of epidural ropivacaine for treatment of HPS.

It is important to highlight that the described technique requires particular training and handskills. Anyway, paediatric anaesthesia nowadays is a highly specialized profession and therefore children, independent of the severity of their disease, should be treated only by dedicated paediatric anaesthetists and surgeons (21). In experienced hands, ultrasound guided thoracic epidural anaesthesia may be considered as safe technique. Careful miscellaneous management (sedation under maintenance of spontaneous respiration, suctioning of the stomach, etc.) is equally important for the safe and successful management of these cases.

The presented technique is only possible when open surgical procedures are performed. Nowadays laparoscopic procedures become more and more popular, which is also the case for pyloromyotomy. Two recent meta-analyses are available comparing open versus laparoscopic pyloromyotomy. Sola et al. identified 6 studies with 303 patients sufficient to be included in a meta-analysis and found slight advantages for laparoscopic procedures in terms of shorter time to full feeding, shorter postoperative length of stay and a reduced rate of total complications (22). Conversely, Hall et al. published a meta-analysis based on 8 studies and 595 patients and found fewer complications and a higher efficacy when open pyloromyotomy was performed (23). None of these publications consider the implications of anaesthesia management on morbidity. Anyway, from today's point of view laparoscopic pyloromyotomy shows no clear advantages compared with the open technique. Moreover, avoidance of tracheal intubation, subsequent ventilation and possible postoperative respiratory depression when thoracic epidural anaesthesia is performed may serve as another argument for open pyloromyotomy.

A major advantage of the reported technique is the fact that it can be used for all surgical open approaches. Despite not investigated in this retrospective analysis, paraumbilical procedures can be also treated with epidural single shot blockade. Another possible advantage is the operation time saving effect of pure regional anaesthesia based methods as compared with general anaesthesia. Kachko et al. reported about the time saving effects of spinal anaesthesia for HPS (24). Despite we did not exactly evaluate the anaesthesia control time in our retrospective analyses, it seems to be obvious that the avoidance of emergence from general anaesthesia is directly associated with faster procedural times.

The clear limitation of this report is that it is just a descriptive consecutive case series. However, from our point of view descriptive reports are useful and sufficient to describe anaesthetic techniques for particular surgical procedures. Another limitation is that we were

not able to evaluate OPS scores on the ward and therefore no statement regarding pain and behaviour after transfer from the recovery room can be provided.

In summary, thoracic epidural anaesthesia under sedation for pyloromyotomy **has been a useful technique in this retrospective series of infants** suffering from HPS. We did not observe any haemodynamic or respiratory complications in this consecutive series of 20 infants undergoing Weber-Ramstedt pyloromyotomy. Undisturbed surgery was possible in all cases.

For Peer Review

Tables

Table 1. Pertinent patient data. Values are median (*min-max*) except where indicated otherwise

<i>Patient data</i>	
Gender (m/f)	17/3
Age (months)	1.7 (1.5 - 4.0)
Weight (g)	3895 (1800 - 5000)
Duration of surgery (min)	25 (10 - 40)

Table 2. Relevant blood gas values, sedation details and volume of local anaesthetic for epidural single shot blockade. Values are mean (*min-max* or *SD*)

Relevant blood gas values	
pH at admission	7.52 (7.41 - 7.67)
HCO ₃ ⁻ (mmol·l ⁻¹) at admission	30.9 (21.1 - 44.2)
pH preoperative	7.40 (7.36 - 7.49)
HCO ₃ ⁻ (mmol·l ⁻¹) preoperative	25.3 (21.6 - 28.0)

Sedation details

Total propofol (mg) 13.3 (7.8)

Nalbuphine (mg) 0.6 (0.2)

Local anaesthetic volume 2.9 (1.4 - 3.8)

Figure legends

Figure 1

Position of the ultrasound probe relative to the Tuohy epidural needle

Figure 2

Ultrasound observation of the spread of local anaesthetic inside the epidural space. LA = local anaesthetic inside the epidural space; DM = dura mater; CM = conus medularis

Figure 3

Heart rate of all infants before and after single shot thoracic epidural blockade. Bold line = mean heart rate

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Position of the ultrasound probe relative to the Tuohy epidural needle
1236x824mm (72 x 72 DPI)

Review

Ultrasound observation of the spread of local anaesthetic inside the epidural space. LA = local anaesthetic inside the epidural space; DM = dura mater; CM = conus medularis
133x91mm (96 x 96 DPI)

review