Interfascial Plane Blocks Reduce Postoperative Pain and Morphine Consumption in Thoracic Outlet Decompression

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Interfascial Plane Blocks Reduce Postoperative Pain and Morphine Consumption in Thoracic Outlet Decompression

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eindhoven, the Netherlands

Background: Postoperative analgesia in patients undergoing transaxillary thoracic outlet decompression (TATOD) is challenging because of the invasive surgery, the complex innervation of the axillary region, and the preoperative use of opioids by many patients. Commonly, postoperative pain is managed with additional opioids that introduce well-known side effects. To investigate the analgesic efficacy of 2 novel regional anesthesia techniques, we performed a retrospective study comparing the combined pectoral block type 1 and erector spinae block (PECS 1 + ESB) and the pectoral block type 2 (PECS 2) and systemic intravenous opioids regimen (no block) in patients undergoing TATOD.

Materials and methods: We performed 10 PECS 1 + ESB and 10 PECS 2 blocks in patients undergoing TATOD. Twenty patients were randomly selected as controls. The primary endpoint was pain. Secondary endpoints were opioid use, nausea, and vomiting.

Results: Postoperative maximal numeric rating scale scores on recovery were significantly lower in patients receiving either a PECS 1 + ESB or a PECS 2 block compared with controls without block (no block: median 6.00, interquartile range [IQR] 3.00; PECS 1 + ESB: median 4.50, IQR 4.00; PECS 2: median 4.00, IQR 5.00; P = 0.031). Postoperative intravenous morphine consumption was 43% lower in the PECS 1 + ESB group and 56% lower in the PECS 2 group compared with the group with no block (oral morphine equivalents: no block: mean 16.05 ± SD 6.79 mg; PECS 1 + ESB mean 9.05 ± SD 6.24 mg; PECS 2: mean 7.00 ± SD 6.16; P = 0.03 and P = 0.003, respectively). There was no statistical difference in both nausea and vomitus (no block 45% nausea and 30% vomitus, PECS 1 + ESB 40% nausea and 20% vomitus, PECS 2 10% nausea and 0% vomitus, P = 0.17 and P = 0.14, respectively).

Conclusions: There was a significant reduction in postoperative pain and opioid consumption for patients treated with either the PECS 1 + ESB block or PECS 2.

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INTRODUCTION

Thoracic outlet syndrome (TOS) is a group of potentially disabling conditions thought to be caused by the compression of neurovascular structures going to the upper extremity.1–4 Three distinct types of TOS exist: arterial, venous, and neurogenic TOS, by compression of the artery, vein, or plexus, respectively.1 In certain cases—when patients do not improve with physiotherapy—surgery can be performed to release the compressed neurovascular structures at the thoracic outlet.1,5,6 Several approaches (transaxillary, supraclavicular, infraclavicular, and transthoracic) are used for surgical decompression.7 In our center, we perform a transaxillary thoracic outlet decompression (TATOD) with resection of the first rib, partial scalenectomy, and lysis of either the plexus, vein, or artery. In every patient with pain over the pectoral minor insertion, a tenotomy of the most proximal 4–5 cm of the pectoral minor muscle is performed as well during the TATOD procedure.

Postoperative analgesia in TATOD is difficult because of the extensive nature of the surgery, the complex innervation of the thoracic wall and axillary region, and the limited possibilities of oral pain relief in patients already using oral pain relief on a daily base.8,9 Commonly, pain is managed with additional opioids, which introduces well-known side effects like nausea, vomitus, constipation, drowsiness, and respiratory depression. This ultimately leads to worse patient experience and longer hospital stay.10–12

Regional anesthesia techniques are a good supplement to reduce postoperative pain and reduce the need for opioids. These techniques are part of a multimodal analgesia approach.13 Techniques such as thoracic paravertebral, thoracic epidural, and interscalene brachial plexus blocks are routinely used in thoracic and axillary surgery.8,12,14–18 However, these techniques are not easily performed and have their own complications. The thoracic interfascial plane blocks are easy alternatives to provide regional anesthesia with a very low risk of complications.7,19,20 The pectoral nerve blocks (PECS) type I and type II and the erector spinae block (ESB) are thoracic interfascial plane blocks providing analgesia of the thoracic wall and axillary region.8,12,14–18

We hypothesized that the combined pectoral block type I and erector spinae block (PECS 1 + ESB) and PECS 2 might provide additional analgesia in patients undergoing TATOD, blocking intercostal nerves and sensory brachial plexus nerves. The motor branches of the brachial plexus compressed at the thoracic outlet are not blocked with these techniques. This is an advantage as these techniques do not interfere with motor function of the arm and thus, do not obscure early postoperative complications of the brachial plexus nerves after TATOD surgery.

After introduction of both the PECS 1 + ESB and PECS 2 techniques in our clinical practice, we retrospectively evaluated pain and opioid consumption in patients diagnosed with TOS treated with TATOD. Our secondary outcome parameters were postoperative morphine consumption and postoperative nausea and vomitus (PONV).

MATERIALS AND METHODS

This retrospective case-control study was approved by the Medical Research Ethics Committees United (W18.227).

Case and Control Selection

In a 6-month period (May 2018–October 2018), we performed 20 thoracic wall blocks in patients undergoing TATOD: 10 patients with PECS 1+blocks combined with an ESB and 10 patients with a PECS 2 block. These blocks were part of clinical expertise and performed by the same anesthesiologist (B.V.). All patients undergoing TATOD were asked by this anesthesiologist if they would like a supplementary analgesia through a block. If a case was identified, a control patient treated with TATOD without the thoracic interfascial plane block in the same time period was included as a control. In total, 20 controls were selected. All patients were treated by the same surgeon, who agreed with the implementation of the blocks. All participating patients consented to the anonymous use of their data for study purposes. This question to consent is part of the routine preoperative anesthesia questionnaire, and patients are free to accept or not. There is ample time (>1 week) between the preoperative consultation and the date of surgery for patients to re-evaluate their consent.

Analgesia Technique

The axillary region is innervated by the thoracic spinal nerves and branches of the brachial plexus. The PECS 1 is an interfascial plane block in which local anesthetic is deposited between the pectoralis major and minor muscles, blocking the lateral and medial pectoral nerves, provided by the brachial plexus and the intercostobrachial nerve.9,19,21,22 This block provides pain relief in the area of the pectoral major muscle and is used in breast surgery, breast...
expander surgery, catheter surgery, and pacemaker insertion.23

The PECS 2 is a modification of the PECS 1 where an additional deposit of local anesthetic is injected between the pectoralis minor and serratus anterior muscles. This blocks the lateral branches of at least the T2–T4 spinal nerves and the long thoracic nerve, as well as the intercostobrachial nerve and the lateral and medial pectoral nerves from the PECS 1 deposit. The PECS 2 block provides pain relief for the pectoral and axillary region and is suitable for more extensive surgery: mastectomy and axillary clearance.23–26

The ESB is administered beneath the erector spinae muscle group, blocking dorsal and ventral rami of the spinal nerves causing sensory blockade over the dorsolateral and anterolateral thorax, provided by the spinal nerves20,27–31

Erector Spinae Block. The patient is placed in a lateral decubitus with the side of interest upward. A curve array probe or a high-frequency linear probe, depending on BMI of the patient, is placed in a longitudinal position 2–3 cm lateral to the vertebral column. The erector spinae muscles are identified in relation to the ipsilateral fifth thoracic vertebra (T5) transverse process. A needle is inserted with an in-plane technique in a cephalad to caudal direction until bone contact with the top of the transverse process is reached. After slight retraction of the needle, 20 mL of 0.5% ropivacaine hydrochloride is injected.

Pectoral Nerves Block Type 1. The patient is in supine position. A high-frequency linear probe is placed horizontally at the level of the third rib and vertically below the lateral third of the clavicle. Then the probe is rotated 45° counterclockwise. The corresponding ultrasound image shows the pectoralis major and minor muscles and the pectoral branch of the thoraco-acromial artery in the interfascial plane between both muscles. The needle is introduced in plane from medial to lateral and 20 cc of 0.5% ropivacaine is injected in this interfascial plane. In addition to a PECS 1 block, in a PECS 2 block, an instillation of 20 mL of 0.5% ropivacaine hydrochloride is injected underneath the pectoralis minor muscle. If necessary, the local anesthetic is diluted with saline to achieve the right amount of volume without exceeding the maximal dose of 3 mg/kg ropivacaine.

Preoperative Management
As part of the standard protocol, all patients were given an intravenous (IV) access and receive acetaminophen as part of the preoperative management.

Intraoperative Management
On arrival in the OR, standard American Society of Anesthesiologists (ASA) monitoring was applied, and IV access was obtained. After completion of the sign-in procedure, general anesthesia (GA) was induced. After induction of GA, the PECS 1 + ESB or the PECS 2 block was performed as described. Surgery was commenced 10 min after finalizing the block.

Postoperative Management
At the postanesthesia care unit (PACU) and ward, postsurgical pain management was performed as per the hospital’s postoperative pain protocol. Patients are clinically assessed in a routine fashion for postoperative complications. Nausea, vomiting, and numeric rating scale (NRS) scores are documented by the PACU nurses. If a patient has an NRS score more than 3, IV boluses of morphine (1 mg/mL) are titrated until pain relief with NRS < 4 was achieved. The maximum amount of morphine is decided by the attending anesthesiologist and does not exceed 20 mg. Ketamine is given to reduce postoperative pain, if morphine alone is inadequate (NRS > 3, after morphine titration). All patients received standard PONV prophylaxis consisting of granisetron 1 mg and dexamethasone 4 mg during surgery. Patients were discharged from the PACU if Aldrete’s scores ≥ 8, NRS scores < 3, and PONV was absent or treated. At the surgical ward, a medical assessment is documented routinely at 4 different time intervals. All medication administration is documented in the patient’s clinical file. Postoperative pain at the surgical ward is managed with paracetamol, nonsteroid anti-inflammatory agents, and oral opioids.

Data Collection
All data were collected from the electronic patients file. The attending anesthesiologist collected data with regard to the anesthesia and surgical procedure. This was performed to evaluate the results of these blocks, based on clinical expertise. Nurses and surgical residents documented the patient’s status on the PACU and the surgical ward as part of standard practice. Morphine and sufentanil dosages were converted to morphine equivalent dose (MED). Oral opioids, only administered at the surgical ward, are not included in the calculation of the MED. We collected data on 4 different time points: preoperative, at the PACU, during the evening round at the ward, and the following morning. Because a local block only acts for a short period of
time (+- 4 hr), the data at admission and during morning rounds postoperatively will serve as baseline data.

**Statistical Analysis**

The primary endpoint was the NRS score. Secondary endpoints were opioid use, nausea, and vomiting. Continuous variables were presented as mean and standard deviation (SD) or median and interquartile range (IQR), depending on normality. Categorical demographic variables were reported as number. Percentages were used to report on PONV. Differences in normal distributed continuous variables across the 3 groups were tested using a one-way analysis of variance. Differences between groups were estimated using the Bonferroni correction. Differences in not-normal distributed continuous variables were compared with a Kruskal-Wallis test across the 3 groups and with the Mann-Whitney U-test comparing 2 groups. The Bonferroni correction was used ($P$-value < 0.016) to correct for multiple comparison. Differences in categorical variables between the 3 groups were tested using the Fisher’s exact test. A $P$ value < 0.05 was considered statistically significant when comparing 3 groups. Statistical analyses were performed using SPSS, version 25 (SPSS Inc, Chicago IL).

**RESULTS**

Patient demographics, ASA classification, type of TOS, and duration of surgery were comparable in all groups (Table I). The length of stay was comparable between all groups. In total, 3 VTOS patients and 17 NTOS patients were treated in the ‘no block’ group. 1 VTOS patient and 9 NTOS patients were treated in the PECS 1 + ESB group, and 10 NTOS patients were treated in the PECS 2 group. There were no patients with ATOS. The NRS scores on admission of the patient were similar in all groups (no block: median 1.50, IQR 4.00; PECS 1 + ESB: median 3.50, IQR 4.00; PECS 2: median 3.00, IQR 4.00; $P$ = 0.40) (Table II and Fig. 1). Comparing the 3 groups, postoperative maximal NRS scores on the recovery were significantly lower in patients with no block (no block: median 6.00, IQR 3.00; PECS 1 + ESB: median 4.50, IQR 4.00; PECS 2: median 4.00, IQR 5.00; $P$ = 0.031). There was no statistical difference between NRS scores at the evening ($P$ = 0.060) and the next morning ($P$ = 0.13).

When comparing groups, a trend for lower pain scores was seen on the recovery in patients receiving a thoracic wall block than in no block patients, however statistically nonsignificant ($P$ = 0.019 for PECS 1 + ESB versus no block and $P$ = 0.069 for PECS 2 versus no block). If we compare the pain scores of PECS 1 + ESB to the PECS 2 group, no statistical differences were found (Table II).

Postoperative IV morphine consumption was lowered by 43% in the PECS 1 + ESB and by 56% in the PECS 2 compared with no block (oral morphine equivalents: no block: mean 16.05 ± SD 6.79 mg; PECS 1 + ESB mean 9.05 ± SD 6.24 mg; PECS 2: mean 7.00 ± SD 6.16; $P$ = 0.029 and $P$ = 0.003, respectively). Intraoperative pain relief with sufentanil was comparable in all groups (no block: mean 68.40 MED ± SD 6.71 mg; PECS 1 + ESB mean 57.60 MED ± SD 6.24 mg; PECS 2: mean 56.40 ± SD 6.16, $P$ = 0.27) (Table III). In the PECS 1 + ESB group, 4 patients (40%) experienced nausea and 2 patients (20%) vomited. In the PECS 2 group, 1 patient (10%) experienced nausea and no patients (0%) suffered from vomiting (Table IV). The no-block group had 9 patients with nausea (45%) and 6 patients (30%) who vomited after surgery. For both nausea ($P$ = 0.17) and vomiting ($P$ = 0.14), no statistically significant difference was found. No patients were treated with IV morphine in the ward.

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**Table I.** Demographic data for patients without thoracic wall block, a PECS 1 + ESB block and a PECS 2 block

<table>
<thead>
<tr>
<th>Variable</th>
<th>No block (n = 20)</th>
<th>PECS 1 + ESB (n = 10)</th>
<th>PECS 2 (n = 10)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, (y)</td>
<td>38.00 (17.00)</td>
<td>40.50 (30.00)</td>
<td>45.00 (27.00)</td>
<td>0.41</td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>3/17</td>
<td>3/7</td>
<td>2/8</td>
<td>0.63</td>
</tr>
<tr>
<td>ASA classification (1/2)</td>
<td>6/14</td>
<td>7/3</td>
<td>3/7</td>
<td>0.082</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>23.35 (7.28)</td>
<td>24.60 (5.43)</td>
<td>24.70 (7.97)</td>
<td>0.54</td>
</tr>
<tr>
<td>Length of surgery, (h)</td>
<td>1:12 (0:15)</td>
<td>1:10 (0:26)</td>
<td>1:09 (0:10)</td>
<td>0.54</td>
</tr>
<tr>
<td>NTOS/VTOS/ATOS</td>
<td>17/3/0</td>
<td>9/1/0</td>
<td>10/0/0</td>
<td></td>
</tr>
</tbody>
</table>
DISCUSSION

This case-control study compared the PECS 1 + ESB block and PECS 2 block with the standard of care and demonstrated less postoperative pain on the recovery with a reduced need for morphine administration. Morphine consumption is associated with well-documented side effects such as PONV, drowsiness, constipation, ileus, respiratory depression, and urinary retention. In this study, we only assessed nausea and vomiting as a side effect of morphine.

As baseline demographic and preoperative NRS scales are similar in all groups, we believed we managed to sample representative groups of patients.

To our knowledge, this is the first report of the use of thoracic interfascial plane blocks in TOD surgery. If we compare our results with the use of thoracic interfascial plane blocks used in breast, axillary, or thoracic surgery, our findings are consistent. A recent meta-analysis about PECS blocks in breast and axillary surgery shows a decreased use of morphine within the first 24 hr after surgery. The use of this plane block has also been described in thoracic surgery, cardiac surgery, and chronic thoracic pain cases. PECS blocks are easily performed and are able to relieve postoperative pain more effectively than traditional analgesics with a reduced need for opioids. There are only limited data comparing ESB with other techniques, as this technique was only conceived in 2016. A review of 242 cases reports a reduction in postoperative pain and opioid use in breast, thoracic, and
cardiac surgery cases.\textsuperscript{27,32,34} ESB was less effective in breast surgery than PECS blocks.\textsuperscript{27}

In most (invasive) thoracic surgery, the paravertebral block (PVB) has taken over the place of the thoracic epidural analgesia (TEA) as the gold standard.\textsuperscript{8,18} However, the PVB is a complex technique with potentially serious side effects and requires a substantial amount of operating time to perform. This may explain why many anesthesiologists are not fond of using the PVB in daily routine.\textsuperscript{33}

In our center, we first decided to perform a PECS 2 block for TATOD. Initial cases were promising; however, posterior thorax pain at the level of the scapula was still present. Therefore, we decided to perform a combination of the PECS 1 and ESB blocks to target the ventral and the lateral posterior side of the thorax, respectively. Nonetheless, patients reported similar pain complaints. Most likely, this pain at the scapula is due to the positioning and the traction onto the nerves during surgery (patients are placed onto their sides with the arm in anteverision and abducted up to 120°). As both blocks gave similar promising results, except for the shoulder pain, we decided to implement the PECS 2 block. As the block is performed after induction and the patient can stay supine to perform a PECS 2 block, it is easiest to implement in daily practice.

There is a significant decrease in NRS scores at the recovery comparing between the fascial plane block groups and the control group. There were no statistical differences comparing between both fascial plane groups. This is probably caused by the small numbers in the groups. Furthermore, we see a trend for lower NRS scores at the recovery and at the ward for both fascial plane blocks compared to the control group. Also, the PECS 1 + ESB block shows a trend to last longer (as NRS scales in the evening are lower) and the PECS 2 block shows a trend to lower PONV levels. Both blocks show a decrease in morphine used postoperatively.

Our results show that the effect of a block is biggest at the PACU/recovery. The NRS scores 24 hr after surgery are similar in all groups. This is consistent with the fact that the effect of ropivacaine hydrochloride lasts for 2–8 hr. Because there was no morphine administration at the surgical ward, there is no need to extend the effect of the locoregional blocks (through an indwelling catheter or adding steroids to the injectate). There was no difference in the length of stay. All patients were able to leave the hospital 1 day after surgery. If we compare with other centers, this is quite exceptional: the mean length of stay varies from 2 to 5 days.\textsuperscript{35,36} This is most likely explained by the efforts of the anesthesiology team to offer patients multimodal analgesia—of which local blocks are a part of. There are multiple publications that connect high levels of opioid prescription to a longer length of stay because of opioid-induced side effects.\textsuperscript{10,35,36} In a publication by the group of Karl Illig, they were able to reduce the length of stay after TATOD from 4 to 2.6 days by introducing multimodal analgesia (with nonsteroid anti-inflammatory agents and benzodiazepines mixed with lower dosage of opioids).\textsuperscript{36}

The decision not to block the nerves of the brachial plexus itself can be discussed. Although

<table>
<thead>
<tr>
<th>Variable</th>
<th>No block (n = 20)</th>
<th>PECS 1 + ESB (n = 10)</th>
<th>PECS 2 (n = 10)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sufentanil intraoperative (MED)</td>
<td>68.40 23.06</td>
<td>57.60 20.23</td>
<td>56.40 20.45</td>
<td>0.63</td>
</tr>
<tr>
<td>Morphine recovery (MED)</td>
<td>16.05 6.79</td>
<td>9.05 6.24</td>
<td>7.00 6.16</td>
<td>1.000</td>
</tr>
</tbody>
</table>

Table IV. PONV for patients without thoracic wall block, a PECS 1 + ESB block and a PECS 2 block

<table>
<thead>
<tr>
<th>Variable</th>
<th>No block (n = 20)</th>
<th>PECS 1 + ESB (n = 10)</th>
<th>PECS 2 (n = 10)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nausea, %</td>
<td>45%</td>
<td>40%</td>
<td>10%</td>
<td>0.17</td>
</tr>
<tr>
<td>Vomitus, %</td>
<td>30%</td>
<td>20%</td>
<td>0%</td>
<td>0.14</td>
</tr>
</tbody>
</table>
the transaxillary approach for thoracic outlet decompression is still open surgery, the size of the incision and extent of surgical dissection is quite minimal. Blocking the nerves that are freed from the overlying tissue (although not blocking the rami of the thoracic spinal nerves) might provide patients with excellent pain relief. However, we are weary to block the motor nerves in the brachial plexus, leaving the patient with ‘a dead arm’ for up to 24 hr, obscuring possible complications and possibly increasing the length of stay. Therefore, it was a conscious decision of the anesthesiologist not to block any motor nerves of the brachial plexus.

There are several limitations to our study. This is a retrospective evaluation of a clinical implementation and therefore lacks all prospective, randomized, or placebo-controlled elements. Data collection was complete but was limited to the data that are routinely assessed at the PACU and ward. Therefore, only pain scores, opioid consumption, and the presence of nausea and vomiting were assessed in this research. Other side effects of morphine (drowsiness, urinary retention, constipation, and respiratory depression) are not routinely documented and could therefore not been included. We did not include oral opioid therapy in the calculation of total morphine consumption (MED). This might have affected outcomes. The number of patients in this study was small.

In this retrospective case-control study, we found a significant reduction in postoperative pain and opioid consumption for patients treated with either the PECS 1 + ESB block or PECS 2 block compared with no block in patients undergoing thoracic outlet decompression for TOS. There was a trend for less nausea and vomiting.

REFERENCES