

Unilateral superficial cervical plexus block as pre-emptive analgesia for hemithyroidectomy under general anaesthesia

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Abstract

Background and Aims: Superficial cervical plexus block is effective in reducing pain following thyroid surgeries. We studied the sensory block onset time and duration of analgesia time on adding dexmedetomidine to ropivacaine for unilateral superficial cervical plexus block (USCPB) in patients undergoing hemithyroidectomy. **Methods:** Forty American Society of Anaesthesiologists (ASA) Class I or II adult patients who were scheduled to undergo hemithyroidectomy were randomly allocated to the following groups to receive unilateral superficial cervical plexus block: 9ml 0.5% Ropivacaine (Group C) or 9ml 0.5% Ropivacaine + 1 mcg/kg Dexmedetomidine (Group D) as USCPBs before induction of general anaesthesia. The sensory block onset time, duration of analgesia, mean arterial pressure (MAP), heart rate (HR), and the incidences of side effects, such as hypotension and bradycardia were recorded. **Results:** The addition of dexmedetomidine to ropivacaine (Group D) shortened the sensory block onset time compared with the ropivacaine group (Group C) ($p < 0.05$). The duration of analgesia of cervical plexus block in Group D was significantly longer than that in Group C ($p < 0.001$). The Ramsay sedation score after extubation in Group D was significantly higher than that in Group C ($p < 0.05$). MAP level and HR level in Group D were significantly lower than that in Group C ($p < 0.05$). **Conclusion:** The addition of 1 mcg/kg dexmedetomidine to ropivacaine for cervical plexus block could shorten the sensory block onset time and extend the duration of analgesia, and increase the quality of analgesia than with 0.5% ropivacaine alone in patients undergoing hemithyroidectomy.


Keywords: Unilateral superficial cervical plexus block, Hemithyroidectomy, Ropivacaine, Dexmedetomidine, General anaesthesia, Pre-emptive analgesia.

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INTRODUCTION

Superficial cervical plexus block (SCPB) has been found to be very effective in procedures of neck such as thyroid surgeries, clavicular surgery, carotid endarterectomy and tracheostomy. The use of regional anaesthesia in combination with general anaesthesia reduce the requirements for opioid analgesics and provide prolonged postoperative analgesia. Thyroid surgery is considered to be moderately painful with a mean postoperative pain

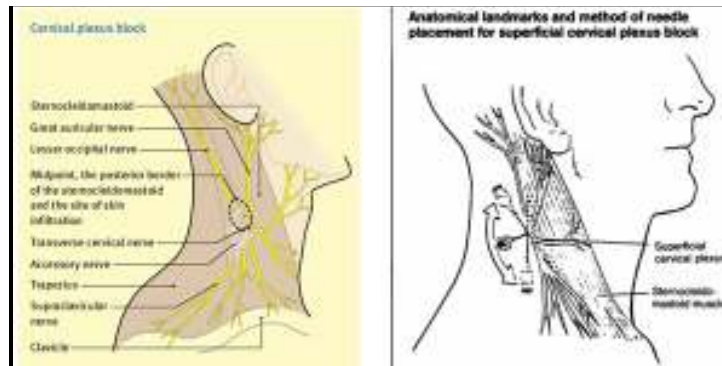
score of 6.9 on a visual analog scale (VAS) from 0 to 10. Recent reports suggest that patients experience a significant amount of pain, especially in the early postoperative hours despite modern and less invasive surgical techniques. In addition, thyroid surgery is reported to be associated with a high risk of postoperative nausea and vomiting (PONV). Analgesics inducing nausea or vomiting, such as opioids or nefopam, should be avoided. We hypothesized that the unilateral superficial cervical plexus block will reduce the intraoperative anaesthetic requirements and prolong the postoperative analgesia in patients undergoing hemithyroidectomy. The aim of this study is to evaluate the intra- and postoperative analgesic efficacy of unilateral superficial cervical plexus block for hemithyroidectomy surgeries.

MATERIAL AND METHOD

After ethical committee approval and informed written consent, 40 adult patients ASA physical status I-II scheduled for elective hemithyroidectomy under general

anaesthesia were eligible for the study. Exclusion criteria include history of allergy to the drugs used, local sepsis, bleeding disorder, diaphragmatic motion abnormality, inability to understand the study protocol or pain scale and patients under medications that would influence autonomic or cardiovascular responses to the surgery. Patients were premedicated with midazolam (0.03 mg/kg IV). Patients were allocated randomly into two equal groups, to receive either 9ml 0.5% Ropivacaine (Group C) or 9ml 0.5% Ropivacaine + 1 mcg/kg Dexmedetomidine (Group D) as USCBs before induction of general anaesthesia. Using a three-point injection technique, 9 ml of the prepared mixture was injected using a 23-gauge needle, inserted at the midpoint of the sternocleidomastoid muscle, corresponding to the C3 transverse apophysis, in three directions. Six mL of the solution was injected up and down at the posterior border of the sternocleidomastoid muscle to block the occipital, auricular, and supraclavicular branches of the superficial cervical plexus and 3 mL was injected horizontally above the muscle to block the transverse cervical nerve. After establishing standard monitoring, anaesthesia was induced with pentazocine 0.5 mg/kg IV and propofol 2mg/kg IV. Orotracheal intubation was facilitated by the administration of atracurium 0.5mg/kg IV. Anaesthesia

was maintained with isoflurane in a N2O/ O2 (50/50) mixture. Mean arterial blood pressure and heart rate were recorded after administration of block, every 5 min (the values during every hour of surgery were averaged to give a single mean value) and at 1 h after recovery. At the completion of surgery, isoflurane was discontinued and residual neuromuscular block was antagonized with neostigmine 0.05 mg/kg and glycopyrrolate 0.01mg/kg. Visual analogue score were recorded at (1 h, 2 h, 6 h, 12 h, 24 h) postoperatively. Diclofenac sodium 75 mg i.v was used as a rescue analgesic when VAS score was more than 5. Postoperative monitoring of side effects related to the cervical plexus block were recorded (such as phrenic nerve block, brachial plexus block, vagus nerve block, Horner’s syndrome). Any episodes of bradycardia (HR<40 beats/min), hypotension (MAP<65 mm Hg), nausea, vomiting, and excessive sedation were recorded during the first 24 h after surgery. The primary outcome was the duration of analgesia of cervical plexus block in both groups. The secondary outcomes include hemodynamic differences between the groups. The quality of analgesia and all adverse events related to surgery and the regional anaesthetic technique were also recorded.



Statistical Analysis

Results were expressed as mean ± SD, analyzed using tests of significance to identify the variables significantly to differences in different groups: Paired t– test, student t-test. We used Fisher's Exact Test to compare between numbers of patients in each group. Statistical significance was considered at the level of p<0.05.

RESULTS

Table 1: Demographic data associated with two groups

	Group D (n=20)	Group C (n=20)
Age (y)	33.30 ± 5.17	33.15 ± 6.71
Sex (F/M)	3/17	4/16
Weight (kg)	55.25 ± 3.95	53.65 ± 5.33
Duration of surgery (min)	109.13 ± 12.37	113.48 ± 11.84

Values are presented as mean standard error or number of patients. There were no significant differences between groups.

Table 2: Characteristics of superficial cervical plexus block

	Group D (n=20)	Group C (n=20)	p
Onset time of sensory block (min)	4.72 ± 1.15	6.64 ± 1.27	<0.05
Duration time of sensory block (min)	403.54 ± 16.18	249.52 ± 24.25	<0.001

Values are presented as mean standard error.

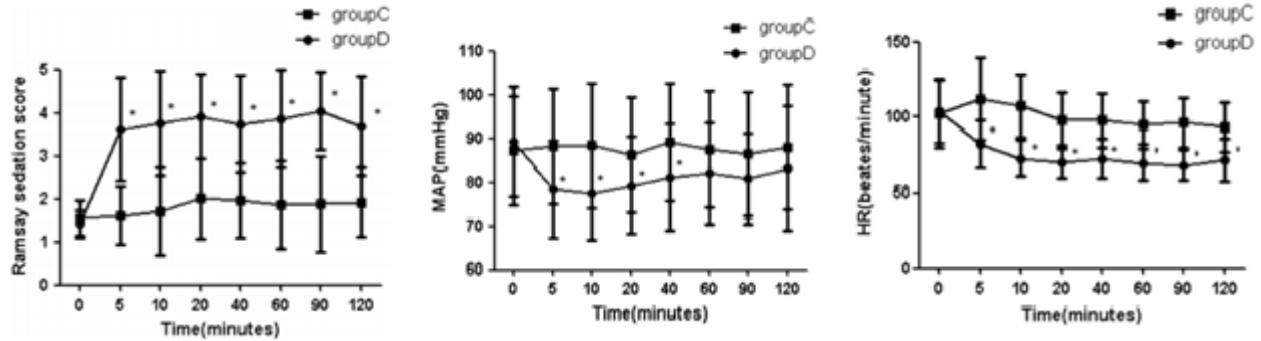


Figure 1: The Ramsay sedation score for the groups after extubation. Asterisks indicate time points of a statistically significant difference ($p < 0.05$) when the groups were compared.

Figure 2: Mean arterial pressure (MAP) for the groups after administration of block. Asterisks indicate time points of a statistically significant difference ($p < 0.05$) when the groups were compared.

Figure 3: Heart rate (HR) for the groups after administration of block. Asterisks indicate time points of a statistically significant difference ($p < 0.05$) when the groups were compared.

RESULTS

The Demographic data were similar in each group (Table 1). The Sensory block onset time in Group D is shorter than in Group C; the difference was statistically significant (Table 2, $p < 0.05$). The Sensory block duration time in Group D is longer than in Group C; the difference was statistically significant (Table 2, $p < 0.001$). The Ramsay sedation score at 5, 10, 20, 40, 60, 90, and 120 minutes after extubation in Group D was significantly higher than in Group C. (Fig. 1, $p < 0.05$). The Mean Arterial Pressure (MAP) level in Group D at 5, 10, 20, and 40 minutes was significantly lower than in Group C (Fig. 2, $p < 0.05$). The Heart Rate (HR) in Group D at 5, 10, 20, 40, 60, 90, and 120 minutes was significantly lower than in Group C. (Fig. 3, $p < 0.05$).

DISCUSSION

The main result of our study demonstrated that the addition of 1 mg/kg dexmedetomidine to ropivacaine for cervical plexus block can shorten sensory block onset time and extend the duration of analgesia. Administering the block prior to the incision as pre-emptive analgesia is more effective because an analgesic treatment initiated before, as opposed to after the surgical procedure, protects the central nervous system from the deleterious effects of noxious stimuli, and the patient from the resulting allodynia and increased pain⁹. Ropivacaine was chosen for its lesser cardiac toxicity compared with bupivacaine⁸. Aunac and colleagues have demonstrated the safety of ropivacaine 0.5% for combined superficial and deep cervical plexus block in thyroid surgery⁶. Meanwhile, study Group D had more hemodynamic stability than Group C. The Ramsay sedation score was higher in Group D, so we predict that the local area use of dexmedetomidine also can produce sedation. Although

dexmedetomidine is commonly used as an intravenous agent, interest in the addition of perineural dexmedetomidine to local anaesthetic is increasing. In a volunteer study, perineural dexmedetomidine with 0.75% ropivacaine prolonged block of ulnar nerve by 60%, while systemic administration of 20 μ g dexmedetomidine resulted in only 10% prolongation of the same, suggesting the peripheral effect of dexmedetomidine when added to the local anaesthetic⁸. Esmoğlu *et al*³ and Ammar and Mahmoud¹⁰ showed that the use of dexmedetomidine was safe as an addition in local anaesthetic for brachial plexus block. Another study¹¹ carried out on volunteers showed that dexmedetomidine could be used as an additive to local anaesthetics for various regional anaesthetic techniques, while simultaneously prolonging perineural nerve block. Dexmedetomidine is well known as a highly selective α_2 adrenoreceptor agonist shown to have both sedative and analgesic effects¹²⁻¹⁴. Its intravenous role has been well recognized as sedative, analgesic and anxiolytic without any respiratory depression^{15,16}. It is well known that in peripheral myelinated and nonmyelinated fibers, membrane hyperpolarization develops that can produce sensory effects and pain during or after stimulation and mainly results from the activation of the sodium-potassium pump after the transient influx of sodium ions¹⁷. Dalle *et al*¹⁸ found that clonidine (an α_2 adrenoreceptor agonist) enhances the sensory blockade by blocking the inhibiting hyperpolarization activated cation current to enhance the level of hyperpolarization and thus inhibits subsequent action potentials. Another effect may be the mechanism of α_2 adrenoreceptor agonist in peripheral nerve by reducing release of norepinephrine and causing α_2 receptor independent inhibitory effects on nerve fibre action potentials. The use of clonidine in peripheral nerve blocks has been reported to be safe and beneficial (it prolongs the duration of anaesthesia and

analgesia) by Singelyn *et al*^{19,20}. Dexmedetomidine is a selective α_2 adrenoceptor agonist similar to clonidine; its effect on the peripheral nerve may be the same as or more potent than that of clonidine. In studies on humans, Esmoglu *et al*,³ Kaygusuz *et al*,⁴ and Rancourt *et al*²¹ showed that dexmedetomidine was safe when used as an addition to local anaesthetic for brachial plexus block and posterior tibial nerve sensory blockade. Esmoglu *et al*³ and Kaygusuz *et al*⁴ also showed that when dexmedetomidine is used as an addition to local anaesthetic, it can provide faster onset and longer duration for brachial plexus block, but resulted in some side effects, such as hypotension and bradycardia. On the basis of our discussions, we assumed that the addition of dexmedetomidine to the local anaesthetic for cervical plexus block could provide the same effect as that of the study by Esmoglu *et al*³. In our study, the addition of dexmedetomidine (1 mcg/kg) to local anaesthetics extended the duration of anaesthesia and increased the quality of analgesia, and patients were sedated and arousable, thus avoiding the use of intravenous sedatives that may induce respiratory depression. The possible mechanism is that the dual role of dexmedetomidine of the central and peripheral, which include slight intravenous effect that cause by the tissue capillary absorption and its direct effect on the peripheral nerves. Dexmedetomidine has a double effect, playing an central sympatholytic role and activating the vagus nerve to lower plasmas catecholamine levels which can lower blood pressure BP and HR, providing stable hemodynamics. However, it also has a dosage related inhibition for BP and HR. Esmoglu *et al*²² showed some side effects such as hypotension and bradycardia in their test group (100 mcg of dexmedetomidine added to levobupivacaine for brachial plexus block), along with its effects such as sedation and anxiolysis. In our study, HR in Group D was lower than that of the Group C, but the incidence of bradycardia was lower than that recorded in the study by Esmoglu *et al*²² (2/20 vs. 7/30); the difference may be because the dose of dexmedetomidine (1 mcg/kg vs. 100 mcg) was lower in our study, and was tolerant and reversible without adverse effects to the patients. In conclusion, USCPB is an effective technique to reduce analgesic requirements during and after hemithyroidectomy surgeries. The addition of 1 mcg/kg dexmedetomidine to ropivacaine for cervical plexus block can decrease sensory block onset time and extend the duration of analgesia, and increase the quality of analgesia. Our data suggest that it is possible to manage pain after thyroid surgery with regional anaesthesia. Further studies are needed to determine the safe optimal dose of dexmedetomidine adding to local anaesthetics for cervical plexus block.

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