

Concentrations and Costs of a Thirty Second Priming Technique with Sevoflurane Using the Circle Circuit

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Abstract

Background : Priming the circuit for an application of inhalation agent depends on the time taken, the fresh gas flow (FGF) rate, the concentration setting of the agent on the vaporizer, and the priming technique. Based on previous studies, the priming time for a desirable concentration of sevoflurane varied between 30 seconds and 5 minutes. Remarkably, although the cost of priming with sevoflurane was lowest at 30 seconds, concentrations in the circuit were not clearly stated. The aim of this study was to test the success (and cost) of a 30-second priming technique to achieve a 4.5 per cent sevoflurane concentration in the circle circuit.

Method : Analyses were done on 20 samplings. Each time the adjustable pressure-limiting (APL) valve on the assembly was closed, the 2-liter reservoir bag emptied and the patient end occluded. For 30 seconds, the circuit was filled with oxygen ($8 \text{ L} \cdot \text{min}^{-1}$) and 8 per cent sevoflurane. The pressure in the circuit was kept constant at 10 mbar by partially releasing the excess gas at the patient end. After 30 seconds, the sevoflurane concentration at the patient end of the circuit was analyzed and the highest concentration recorded. The cost of the sevoflurane used was then calculated.

Results : The sevoflurane concentration in all of the samples was more than 4.5 per cent (mean 6.40%; SD 0.30%). The average amount of liquid sevoflurane used was 1.8 ml at a cost of 42.30 baht (0.9 US\$).

Conclusion : A 30-second priming technique was sufficient to achieve the required concentration of sevoflurane in the circle circuit thus reducing costs for gaseous induction.

Key word : Sevoflurane, Anesthetic Circuit, Cost, Priming Technique

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Gaseous induction of anesthesia in adult patients is an alternative to intravenous induction. Sevoflurane is a volatile anesthetic agent with low blood-gas solubility and lack of a pungent odor. Since its introduction there has been widespread interest in administering techniques. Examples of recently described induction techniques include single vital capacity inhalation induction^(1,2), three sequential vital capacity breaths⁽³⁾, and tidal breathing technique^(4,5).

Gaseous induction is a two-stage procedure: priming - an application of a high concentration of sevoflurane into the circuit - followed by inhalation of the gas mixture by the patient. The concentration of sevoflurane in the circuit depends on the time used, the FGF rate, the concentration settings of sevoflurane on the vaporizer and the priming techniques used. Previous studies confirmed that a sevoflurane concentration of at least 4.5 per cent in the circuit was sufficient for gaseous induction^(4,6) and increasing the concentration over 6 per cent did not markedly shorten induction⁽⁶⁾. The shortest priming time reported for sevoflurane was 30 seconds (range, between 30 to 45 seconds^(3,7-9) and 3 to 5 minutes^(1,10,11)). Remarkably, although the priming time was a major cost factor, its concentration in the circuit was not clearly stated. The authors' aim was to test the concentration (and cost) of sevoflurane in a circle circuit using a 30-second priming technique.

METHOD

Study design

Experimental, descriptive study.

Sample size

The authors postulated that a 4.5 per cent concentration of sevoflurane would be achieved after 30 seconds of priming. Based on pilot studies, the prediction of success according to the hypothesis approached 95 per cent. At the 0.05 level of significance and an acceptable error of 10 per cent, 20 samplings would be needed to confirm the hypothesis.

Priming technique

A standard circle anesthetic circuit with CO_2 absorber was assembled by closing the adjustable pressure-limiting (APL) valve, emptying the 2-liter reservoir bag connected with corrugated tube and occluding the patient end (an elbow connector) with

the thumb. A flowmeter, vaporizer and gas monitor were calibrated. For 30 seconds, the circuit was then filled with oxygen ($8 \text{ L}\cdot\text{min}^{-1}$) and 8 per cent sevoflurane (Drägerwerk AG vaporizer) resulting in a fully inflated reservoir. In order to keep the pressure in the circuit constantly at 10 mbar (approximately 10 cmH_2O) by using the AP monitor of the anesthesia machine (Sulla 808V: Dräger), the excess gas was partially released at the patient end of the circuit. After the 30 seconds, the circuit was re-blocked and the concentration of sevoflurane at the patient end was analyzed by an infrared gas analyzer (Agilent gas monitor: Omnicare; a real time monitor with gas sampling at 120 ml/min) and the highest concentration recorded. A concentration of more than 4.5 per cent was considered successful. Results were presented as the mean \pm standard deviation and the percentage of success. The liquid sevoflurane priming and cost were calculated using Dion's formula: $\text{PFTMC}/2412d$ where P = % gas concentration, F = fresh gas flow (L/min), T = time (min), M = molecular weight (200 g), C = cost/ml (24 baht/ml), and d = density (1.505 g/ml). The calculation was based on a FGF of $8 \text{ L}\cdot\text{min}^{-1}$ and 8 per cent sevoflurane for 30 seconds^(3,12).

RESULTS

Twenty samplings were analyzed and 100 per cent had a sevoflurane concentration greater than 4.5 per cent (range, between 5.90 and 6.95%; mean 6.40%; SD 0.30%). The average amount of liquid sevoflurane used was 1.8 ml at a cost of 42.30 baht (0.9US\$).

DISCUSSION

The mean (6.40%) sevoflurane concentration in the present study was high enough for gaseous induction. Muzy *et al*⁽³⁾ primed the circuit for 30 seconds with FGF $6 \text{ L}\cdot\text{min}^{-1}$ and reported the inspired sevoflurane concentration, the first reading on the monitor after placing the face mask on the patient, was about 8 per cent. Since their study measured the concentration at 30-second intervals, the first reading on the monitor after the 30-second priming meant approximately one minute had elapsed since priming began. Consequently, the sevoflurane concentration in the circuit at the end of the 30-second priming should be less than 8 per cent. Philip⁽⁹⁾ used a 45-second priming technique and claimed that the sevo-

flurane concentration in the inspired-limb was more than 6 per cent, a concentration similar to the present study though the priming time was 15 seconds longer. In fact, the authors' priming technique differed from the two previous studies, in that:

1. The circuit was sealed by closing the APL valve and occluding the patient end. This technique trapped a high concentration of sevoflurane gas in the circuit; resulting in the reservoir bag being fully inflated. In previous studies⁽³⁾, the open APL valve allowed gas to escape. When short-priming, the APL valve should be closed to save gas; only opening it before placing the mask on the patient's face.

2. The excess gas was only partially released at the patient end, allowing the trapped gas with a high concentration of sevoflurane in the reservoir bag to move into the inspired limb of the circuit. In the fill-empty technique⁽⁹⁾, gas was lost from the reservoir bag during the empty phase so that at least 15 seconds was needed to refill the bag. Since the patient end of the circuit was occluded during priming with the fill-empty technique, the concentration of sevoflurane in the inspired-limb increased slowly.

3. In a previous study using the circle circuit, a 2-liter reservoir bag was large enough for gaseous induction⁽¹³⁾. With a short priming time, both the bag size and the FGF rate may affect the concentration

of sevoflurane in the circuit. The authors used a FGF rate of $8 \text{ L} \cdot \text{min}^{-1}$ and 8 per cent sevoflurane (compare Philip)⁽⁹⁾. Therefore, the priming time required to adequately fill the circle circuit with a total volume of about 4 liters (include anesthetic circuit, CO_2 absorber, 2-liter reservoir bag with corrugated tube) was 30 seconds.

The amount of liquid sevoflurane for priming used in the present study was 1.8 ml, at a cost of 42.30 baht (0.9 US\$). The cost of sevoflurane would be lower if the FGF rate were reduced to $6 \text{ L} \cdot \text{min}^{-1}$. Baker⁽⁸⁾ used an FGF rate of $6 \text{ L} \cdot \text{min}^{-1}$ for 45 seconds, which cost £1.05 of sevoflurane. Since the induction time reported in many studies^(14,15) counted time from 'starting prime' to a loss of consciousness, the main cost of priming the circuit hardly differed.

In conclusion, a 30-second priming technique was demonstrably sufficient for achieving the required induction concentration of sevoflurane and it reduced the cost of gaseous induction. The clinical parameters used should be further explored.

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REFERENCES

1. Ti LK, Pua HL, Lee TL. Single vital capacity inhalational anaesthetic induction in adults- isoflurane vs sevoflurane. *Can J Anaesth* 1998; 45: 949-53.
2. Hall JE, Stewart JIM, Harmer M. Single-breath inhalation induction of sevoflurane anaesthesia with and without nitrous oxide: A feasibility study in adults and comparison with an intravenous bolus of propofol. *Anaesthesia* 1997; 52: 410-5.
3. Muzy M, Robinson BJ, Ebert TJ, O'Brien TJ. Induction of anaesthesia and tracheal intubation with sevoflurane in adults. *Anesthesiology* 1996; 85: 536-43.
4. Yurino M, Kimura H. Induction of anaesthesia with sevoflurane, nitrous oxide, and oxygen: A comparison of spontaneous ventilation and vital capacity rapid inhalation induction (VCRII) techniques. *Anesth Analg* 1993; 76: 598-601.
5. Yurino M, Kimura H. A comparison of vital capacity breath and tidal breathing techniques for induction of anaesthesia with high sevoflurane concentrations in nitrous oxide and oxygen. *Anaesthesia* 1995; 50: 308-11.
6. Yurino M, Kimura H. Efficient inspired concentration of sevoflurane for vital capacity rapid inhalation induction (VCRII) technique. *J Clin Anesth* 1995; 7: 228-31.
7. Hall JE, Ebert TJ, Harmer M. Induction characteristics with 3 per cent and 8 per cent sevoflurane in adults: An evaluation of the second stage of anaesthesia and its haemodynamic consequences. *Anaesthesia* 2000; 55: 545-50.
8. Baker CE, Smith I. Sevoflurane: A comparison between vital capacity and tidal breathing techniques for the induction of anaesthesia and laryngeal mask airway placement. *Anaesthesia* 1999; 54: 841-4.
9. Philip BK, Lombard LL, Roaf ER, Drager LR, Calalang I, Philip JH. Comparison of vital capacity induction with sevoflurane to intravenous induction with propofol for adult ambulatory anesthesia. *Anesth Analg* 1999; 89: 623-7.
10. Goto T, Nakata Y, Uezono S, Niimi Y, Uchiyama M, Morita S. Insertion of the cuffed oropharyngeal airway (COPA) with propofol or sevoflurane in adults. *J Clin Anesth* 1999; 11: 280-4.
11. Yamaguchi S, Egawa H, Okuda K, Mishio M, Okuda Y, Kitajima T. High concentration sevoflurane induction of anaesthesia accelerates onset of vecuronium neuromuscular blockade. *Can J Anaesth* 2001; 48: 34-7.
12. Dion P. The cost of anaesthetic vapours. *Can J Anaesth* 1992; 39: 633.
13. Joo HS, Perks WJ, Belo SE. Sevoflurane with remifentanil allows rapid tracheal intubation without neuromuscular blocking agents. *Can J Anaesth* 2001; 48: 646-50.
14. Fleischmann E, Akca O, Wallner T, et al. Onset time, recovery duration, and drug cost with four different methods of inducing general anaesthesia. *Anesth Analg* 1999; 88: 930-5.
15. Smith I, Terhoeve PA, Hennart D, et al. A multi-centre comparison of the costs of anaesthesia with sevoflurane or propofol. *Br J Anaesth* 1999; 83: 564-70.

ความเข้มข้นและต้นทุนของยาสลบเชิงฟลูเรนในการปั๊พนวจจรวงยาสลบชนิดรอบตัวยเวลา 30 วินาที

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บทนำ : การ prime วงจรวงยาสลบให้ได้ระดับความเข้มข้นของยาดมสลบที่ต้องการขึ้นอยู่กับ เวลาที่ใช้ในการ prime, fresh gas flow rate (FGF), ความเข้มข้นของยาดมสลบที่เปิดจาก vaporizer และเทคนิคของการ prime จากการศึกษาที่ผ่านมาพบว่า การ prime วงจรวงด้วย sevoflurane เพื่อให้ได้ระดับยาดามที่ต้องการ ใช้วิลัยแต่ก็ต้องใช้เวลา 30 วินาที ถึงแม้ว่าการ prime ด้วยเวลา 30 วินาทีจะเสียค่าใช้จ่ายน้อยที่สุด แต่ยังไม่มีรายงานการวัดความเข้มข้นของ sevoflurane ในวงจรวงยาสลบที่ชัดเจน

วัตถุประสงค์ : เพื่อหาอัตราความสำเร็จในการ prime วงจรวงยาสลบด้วย FGF 8 ลิตรต่อนาที เปิด sevoflurane 8% เป็นเวลา 30 วินาที ให้ได้ความเข้มข้นของ sevoflurane อย่างน้อย 4.5% ว่าจะมีอัตราความสำเร็จเท่าใด และหาค่าใช้จ่ายของการ prime ในส่วนของยา sevoflurane ว่าเป็นเท่าใด

วิธีการศึกษา : ศึกษาจากการวิเคราะห์ตัวอย่างก้าวจำนวน 20 ตัวอย่าง โดยใช้วิธีการ prime ตั้งต่อไปนี้ ปิด adjustable pressure-limiting valve ของเครื่องวงยาสลบ เป็นไอล์ก้าวใน reservoir bag ขนาด 2 ลิตรออกให้หมด อุดปลาย-วงจรวงนิด circle ด้านที่จะต่อเข้ากับผู้ป่วยให้สนิท แล้วเปิดออกอีกชิ้น 8 ลิตรต่อนาที เปิด sevoflurane ความเข้มข้น 8% นาน 30 วินาที รักษาความดันในวงจรวงยาสลบที่ 10 mbar โดยเปิดให้ก้าวส่วนเกินบางส่วนให้ล้อออกทางปลายวงจรวงด้านที่จะต่อเข้ากับผู้ป่วย บันทึกค่าความเข้มข้นสูงสุดที่ได้ตั้งแต่ 30 วินาที รวมทั้งค่าวนบprimair และราคาของ sevoflurane ที่ใช้ในการ prime ด้วย

ผลการศึกษา : ความเข้มข้นของ sevoflurane ของทุกตัวอย่าง วัดได้สูงกว่า 4.5% (mean 6.40%; SD 0.30%) ค่าวนบprimair เฉลี่ยของ sevoflurane ที่ใช้ได้ 1.8 มล คิดเป็นราคา 42.30 บาท

สรุป : การ prime วงจรวงยาสลบด้วยด้วยเทคนิคดังกล่าวข้างต้น เป็นเวลา 30 วินาที สามารถทำให้ความเข้มข้นในวงจรวงยาสลบเพียงพอที่จะใช้น้ำสลบได้ และช่วยลดค่าใช้จ่ายในการน้ำสลบได้ด้วย

คำสำคัญ : เชิงฟลูเรน, วงจรวงยาสลบ, ค่าใช้จ่าย, เทคนิคในการปั๊พนวจจรวงยาสลบ

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