

Sex Differences in Remifentanyl Requirements for Preventing Cough during Anesthetic Emergence

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Purpose: Target-controlled infusion (TCI) of remifentanyl can suppress coughing during emergence from general anesthesia; nevertheless, previous studies under different clinical conditions recommend significantly different effective effect-site concentrations (effective Ce) of remifentanyl for 50% of patients (EC₅₀). The differences among these studies include type of surgery and patient sex. In recent years, study of sex differences in regards to anesthetic pharmacology has drawn greater interest. Accordingly, we attempted to determine the effective Ce of remifentanyl for preventing cough for each sex under the same clinical conditions. **Materials and Methods:** Twenty female and 25 male ASA physical status I-II grade patients between the ages of 20 and 46 years who were undergoing thyroidectomy were enrolled in this study. The effective Ce of remifentanyl for preventing cough was determined for each sex using the isotonic regression method with a bootstrapping approach, following Dixon's up-and-down method. **Results:** Isotonic regression with a bootstrapping approach revealed that the estimated EC₅₀ of remifentanyl for preventing coughing during emergence was significantly lower in females {1.30 ng/mL [83% confidence interval (CI), 1.20-1.47 ng/mL]} than in males [2.57 ng/mL (83% CI, 2.45-2.70 ng/mL)]. Mean EC₅₀ in females was also significantly lower than in males (1.23±0.21 ng/mL vs. 2.43±0.21 ng/mL, *p*<0.001). Mean arterial pressure, heart rate, and respiratory rate over time were not significantly different between the sexes. **Conclusion:** When using remifentanyl TCI for cough prevention during anesthetic emergence, patient sex should be a considered for appropriate dosing.

Key Words: Sex difference, effect-site concentration, general anesthesia, remifentanyl, target-controlled infusion, cough prevention

INTRODUCTION

Airway reflexes, including coughing, are commonly considered normal responses during emergence from general anesthesia.¹⁻³ However, coughing during emergence may lead to a number of potentially dangerous side effects, including laryngospasm, detrimental hemodynamic changes, and increased intraocular and intracranial pressure. Reportedly, post-thyroidectomy bleeding resulting in acute airway obstruction

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or re-operation occurs in 1-4% of patients,^{4,7} and severe cough may cause such bleeding.^{4,7} In addition to surgeon-related factors, anesthetic factors may also assist in prevention of post-thyroidectomy bleeding. A smooth extubation without significant coughing or retching to avoid raised venous or arterial pressures are important considerations in minimizing the risks of postoperative hemorrhage.⁷ During awakening from endotracheal anesthesia, the trachea may be stimulated by endotracheal tubes, by noxious effects of the anesthetic gas itself, or by uncleared secretions.³ The presence of an endotracheal tube during emergence from general anesthesia is often associated with severe cough.^{3,8} Therefore, prevention of cough during emergence from general anesthesia with the presence of a tracheal tube may be important in patients undergoing certain operations.

Intravenous opioids delivered at the end of surgery can facilitate smooth emergence by reducing coughing^{9,10} and attenuate deleterious hemodynamic changes.¹¹ Specifically, administration of remifentanyl via a target-controlled infusion (TCI) makes it possible to reach a defined target concentration without concern that remifentanyl levels will rapidly change or increase beyond the intended range.¹² Several previous studies, differing in type of surgery, combined anesthetics, and patient sex, have demonstrated reductions of airway reflexes during emergence with the use of remifentanyl TCI,^{13,14} however, the recommended effective effect-site concentrations (effective Ce) of remifentanyl for 50% of patients (EC₅₀) reported in these studies range from 1.46 ng/mL to 2.35 ng/mL.^{13,14}

Numerous animal and human studies suggest that sex may affect opioid analgesia: that is, males require higher doses of opioids to achieve similar levels of analgesia as females.¹⁵⁻¹⁷ This sex difference is known to occur mainly with mu- and kappa-receptor agonists. Sex differences in appropriate opioid dose may affect not only the analgesic system but also act on other inherent properties of the endogenous opioid receptor system, such as respiratory function.¹⁸ Mu- and kappa-opioid receptors in the brainstem contribute to regulation of the cough reflex.¹⁹ If there is the significant sex difference in remifentanyl concentration for airway suppression during anesthetic emergence, a dose regimen suitable for one sex could be excessive for the other, and such information would be clinically important not only in respect of analgesic effect but also in preventing complications.

Accordingly, we hypothesized that men may require higher concentrations of remifentanyl for cough suppression dur-

ing anesthetic emergence than women. The purpose of this study was to determine the EC₅₀ of remifentanyl in effect-site TCI for preventing cough in males and females under the same clinical conditions and evaluate whether there are sex differences in EC₅₀ of remifentanyl for emergence cough suppression.

MATERIALS AND METHODS

This study was approved by the Institutional Review Board of Severance Hospital, Yonsei University Health System (Ref: 4-2011-0356) and registered at <http://ClinicalTrials.gov> (Ref: NCT01614535). Written informed consent was obtained from all subjects. We enrolled 30 male and 23 female patients of ages 20 to 46 years and of ASA I-II status, all of whom underwent general anesthesia for elective thyroidectomy. Exclusion criteria included patients who exhibited signs of a difficult airway, upper respiratory infection in the previous two weeks, hypertension, or diabetes mellitus. Current smokers were excluded, and female patients who were pregnant, breast-feeding, or menopausal were also excluded. In addition, we withdrew patients who experienced delayed emergence and had not regained consciousness by 15 min after the main anesthetic agent had been stopped.

The patients were not premedicated. After arrival at the operating room, routine monitoring, including electrocardiogram, peripheral oxygen saturation, non-invasive arterial pressure, end-tidal carbon dioxide (EtCO₂), and nasopharyngeal temperature, were performed at 1-5 min intervals. Bispectral index (BIS) (BIS[®] monitor, Covidien Medical, Boulder, CO, USA) monitoring was also performed. Anesthesia was induced using propofol 1.5-2.5 mg/kg I.V. and remifentanyl TCI. For TCI of remifentanyl, a commercial TCI pump (Orchestra[®] Base Primea, Fresenius Vial, France) was used, and pump operation was based on the pharmacokinetic model of Minto, et al.²⁰ After non-response to verbal orders and loss of eyelid reflex, rocuronium 0.6 mg/kg I.V. was administered. Tracheal intubation was performed in all patients using a 6.5 mm (internal diameter) tracheal tube for females and a 7.5 mm tracheal tube for males. Cuff pressure was maintained at 20-25 cmH₂O using a hand pressure gauge (Hi-Lo[™] Hand Pressure Gauge, VBM Medizintechnik GmbH, Germany). Anesthesia was maintained with sevoflurane at 1.5-2.5 vol% in an air/oxygen mixture (FIO₂: 0.5, 2 L/min) to achieve a BIS value of 40-55, and a remi-

fentanyl Ce of 2-5 ng/mL was used to maintain patients' mean arterial pressure (MAP) and heart rate (HR) within 20% of baseline values. Mechanical ventilation was maintained with a tidal volume of 8 mL/kg, and ventilator frequency was adjusted to maintain EtCO₂ at 35-40 mm Hg. Nasopharyngeal temperature was maintained at 36-37°C.

Three anesthetists carried out this study. The first conducted anesthetic induction, maintenance, and control of remifentanyl concentration during the emergence period. The second, who did not know the remifentanyl concentration during emergence, led anesthetic emergence and extubation. The third anesthetist recorded all variables of interest to this study, including cough grading and was blinded to the Ce of remifentanyl. Ten min before the end of surgery, sevoflurane was adjusted to an approximate BIS level of 60, and the remifentanyl Ce was titrated to a pre-determined concentration (the initial Ce of remifentanyl being 2 ng/mL). This pre-determined Ce was maintained for at least 15 min throughout emergence. When the operation was completed, 30 mg of ketorolac I.V. and 4 mg of ondansetron I.V. were given, and 0.004 mg/kg of glycopyrrolate I.V. and 0.02 mg/kg of neostigmine I.V. were given to reverse neuromuscular block. Simultaneously, sevoflurane was discontinued, and fresh gas flow was increased to 10 L/min. Mechanical ventilation was converted to manual ventilation 3 min after sevoflurane discontinuation, and EtCO₂ was maintained at 45-55 mm Hg. During this phase, the patients were not disturbed, except for a verbal request to open their eyes. When the patients opened their eyes, deep breathing was encouraged. The tracheal tube was pulled when spontaneous respiration with an adequate tidal volume was confirmed. Immediately after extubation, remifentanyl infusion was stopped, and oxygen was supplemented via a facemask for at least 10 min.

Emergence cough was defined as the cough occurring from the time sevoflurane was turned off to 5 min after extubation. The level of cough was assessed and recorded by the following cough grading system: Grade 0, no cough; Grade 1, single cough with mild severity; Grade 2, cough persistence less than 5 s with moderate severity; or Grade 3, severe, persistent cough for more than 5 s. For estimation of the effective remifentanyl Ce, the up-and-down sequential allocation design was used²¹ (i.e., the predetermined Ce of remifentanyl was determined according to the cough response of the previous patient). If the patient did not cough throughout the peri-extubation period, the pre-determined concentration of remifentanyl for the subsequent patient was decreased by 0.4

ng/mL. Similarly, if the patient coughed at any time before, during, or after extubation, the pre-determined concentration was increased by 0.4 ng/mL. The upper concentration was limited to 3.0 ng/mL due to concerns for patient safety after extubation.^{8,14} The stopping rule, which required at least six pairs of failure/success, necessitated the recruitment of more than 20 patients.²² This up-and-down method was conducted independently for each sex.

The MAP and HR were recorded at the following time points: T1, before induction of anesthesia (baseline); T2, end of surgery; T3, before extubation; T4, just after extubation; T5, 5 min after extubation; T6, 10 min after extubation; T7, before transfer out from the postanesthetic care unit. If MAP <60 mm Hg or HR <50 beats per min (beats/min), we injected ephedrine 4 mg I.V. or atropine 0.5 mg I.V. immediately. In addition, respiratory rate (RR) was recorded at the T4-7 points. The time to eye opening (sevoflurane off to eye opening) and time to extubation (sevoflurane off to extubation) were also recorded. Bradypnea, which was defined as a RR <8 breaths per min (bpm), and SpO₂ below 95% despite oxygen supplementation were monitored during the entire emergence period.

Patient data are presented as mean±standard deviation, or median and range, or numbers of patients. According to previous studies in which the EC₅₀ was estimated by the Dixon's method,¹³ the stopping rule required at least six pairs of failure/success. The EC₅₀ of remifentanyl was defined as the mean value of the independent crossover pairs for each sex, and we compared the mean EC₅₀s using a t-test. To specify the precision of the target concentration, the isotonic regression method was also used for estimating EC₅₀ and EC₉₅ along with a confidence interval (CI).²³ From an observed response rate, which represents the ratio of the number of successful patients to the number of subjects at each concentration level, an adjusted response probability was calculated by pooled adjacent-violators algorithm (PAVA) in order to adhere to the assumption in dose-response determinations that drug effect increases with increased dosage. The CI was estimated by a boot strapping approach.^{22,24} If the value of EC₅₀ did not overlap at the level of 83% CI, the null hypothesis of equal concentration could be rejected as a type I error of 0.05.²⁵ Other normally distributed variables were analyzed using t-tests. Chi-square tests were used for intergroup comparisons of ordinal variables. Statistical analyses were performed using SAS (version 9.2, SAS Inc., Cary, NC, USA), and all *p*-values less than 0.05 were considered significant.

RESULTS

Twenty-three female patients and 30 male patients were enrolled in the study. Among the enrolled patients, three female patients were excluded due to surgical factors (nerve injury and delayed skin suture) and an incorrect pre-determined concentration of remifentanyl. Five male patients were excluded due to surgical factors (converted from simple thyroidectomy to modified radical neck dissection) and delayed emergence. The patients' characteristics are presented in Table 1. Age was comparable between the females and males; however, due to sex differences, patients' height, weight, body surface area, and body mass index were significantly lower in females. Also, the number of patients for each cough grade was comparable between the females and males.

The up-and-down results in consecutive patients are shown in Fig. 1. The EC₅₀ of remifentanyl needed to prevent cough

during emergence based on Dixon's method was significantly less in females (1.23±0.21 ng/mL) than in males (2.43±0.21 ng/mL) ($p<0.001$). The use of isotonic regression estimator, with CIs derived by a bootstrapping approach, revealed that the EC₅₀ of remifentanyl was significantly lower in females than in males [1.30 ng/mL (83% CI, 1.20-1.47 ng/mL) vs. 2.57 ng/mL (83% CI, 2.45-2.70 ng/mL)]. Similarly, the EC₉₅ was significantly lower in females than in males [1.86 ng/mL (95% CI, 1.56-1.96 ng/mL) vs. 2.96 ng/mL (95% CI, 2.77-2.98 ng/mL)] (Table 2). PAVA response rates of both sexes are presented in Fig. 2.

RR was 6.2, 9.1, 11.2, and 13.8 bpm at the T4-7 points, respectively, and RR at T5 and T6 points was significantly faster in females, though there was no difference of RR pattern between sexes over time (p -value=0.28) (Fig. 3). In females, bradypnea, defined as a RR <8 bpm, was observed just after extubation (T4 point) in two of seven patients with a Ce of 1.6 ng/mL. In males, bradypnea was observed in two of 10 patients with a Ce of 2.4 ng/mL, in one of nine

Table 1. Patient Characteristics

	Male group (n=25)	Female group (n=20)	<i>p</i> value
Age	38.36±4.40	37.25±6.81	0.533
Height	173.20±4.22*	161.12±5.61	<0.001
Weight	75.62±11.46*	56.30±5.84	<0.001
BMI	25.22±3.80*	21.76±2.70	0.001
BSA	1.90±0.148*	1.59±0.09	<0.001
ASA physical status classification (%)			0.622
1	22 (88)	18 (95)	
2	3 (12)	1 (5)	
Sevoflurane vol% at BIS60	1.08±0.21	1.06±0.16	0.764

BMI, body mass index; BSA, body surface area.

Values are presented as median±SD or number (percentage).

* $p<0.05$ between groups.

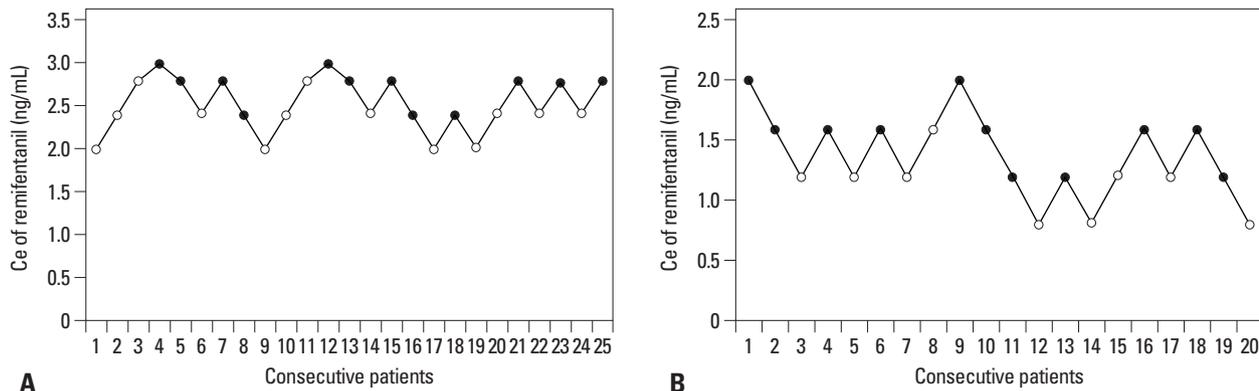


Fig. 1. Assessment of success or failure of smooth emergence over the predetermined concentration of remifentanyl based on consecutive patients by Dixon's up-and-down method. Mean EC₅₀ for smooth emergence was calculated from cross-over pairs from failure (open circle) to success (closed circle) in 25 male patients (A) and 20 female patients (B). EC₅₀, effective effect-site concentration (Ce) of remifentanyl for suppression of emergence cough in 50% of patients.

Table 2. EC₅₀ and EC₉₅ of Remifentanyl for Emergence without Cough

	M	F
Isotonic regression (EC ₅₀ , 83% CI)*	2.57 (2.45-2.70)	1.30 (1.20-1.47)
Isotonic regression (EC ₉₅ , 95% CI) [†]	2.96 (2.77-2.98)	1.86 (1.56-1.96)

PAVA, pooled-adjacent-violators algorithm; CI, confidence interval; EC₅₀, effective Ce of remifentanyl for suppression of emergence cough in 50% of patients; EC₉₅, effective Ce of remifentanyl for suppression of emergence cough in 95% of patients.

Values are presented as the mean (83%* or 95%[†] CI). EC₅₀ and EC₉₅ were calculated by isotonic regression method using the PAVA, and CIs were calculated using a bootstrapping approach.

Table 3. The Time to Eye Opening and the Time to Extubation in Males and Females

	Male group (n=25)	Female group (n=20)	p value
Time to eye opening (min)	10.01±4.43	6.50±1.64	<0.001
Time to extubation (min)	11.27±4.95	7.66±1.91	0.002

SD, standard deviation.

Values are presented as mean±SD. The time to eye opening was defined as the time from sevoflurane discontinuation to eye opening. The time to extubation was defined as the time from sevoflurane discontinuation to extubation.

patients with a Ce of 2.8 ng/mL, and in one of two patients with a Ce of 3.0 ng/mL. One male patient with a Ce of 2.8 ng/mL developed upper airway obstruction and experienced a brief desaturation episode. All of these patients who had adverse respiratory events returned to a normal respiratory pattern within 5 min by encouragement of deep breathing via facial mask without ventilatory support.

The time to eye opening was significantly shorter in females (6.50±1.64 min) than in males (10.01±4.43 min) ($p < 0.001$). The time to extubation was also significantly shorter in females (7.66±1.91 min) than in males (11.27±4.95 min) ($p < 0.001$) (Table 3).

MAP and HR over time did not differ significantly between the sexes [p -value=0.35 (MAP) and 0.082 (HR)] (Fig. 3). The MAP of females was significantly lower than that of males at the T2, T6 and T7 points. HR did not differ between sexes at any time point.

DISCUSSION

In the present study, we found that the remifentanyl requirement for cough suppression during anesthetic emergence in 50% patients was 1.30 ng/mL in females, which was about half of the concentration in males (2.57 ng/mL).

Several studies have investigated the effective Ce of remifentanyl for preventing airway reflexes during anesthetic emergence. Jun, et al.²⁶ demonstrated that 1.5 ng/mL of remifentanyl effectively suppressed cough during emergence from sevoflurane-remifentanyl anesthesia for thyroid surgery. However, Chen, et al.⁸ reported that 2-2.5 ng/mL was needed to achieve the antitussive effect of remifentanyl, following

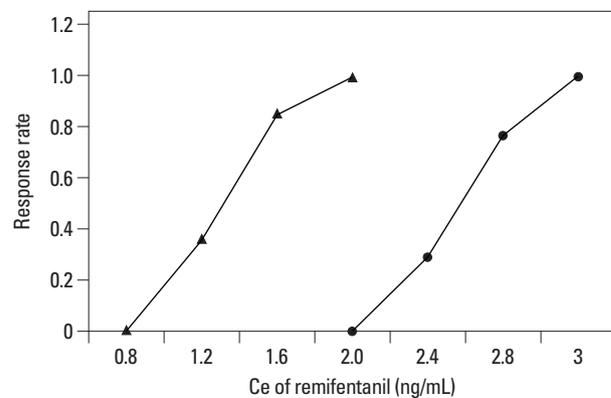


Fig. 2. Pooled-adjacent-violators algorithm response rate in female (▲) and male (●) groups. EC₅₀ of remifentanyl in females was 1.30 ng/mL (83% CI, 1.20-1.47 ng/mL) and 2.57 ng/mL (83% CI, 2.45-2.70 ng/mL) in males. EC₉₅ in females was 1.86 ng/mL (95% CI, 1.56-1.96 ng/mL) and 2.96 ng/mL (95% CI, 2.77-2.98 ng/mL) in males. Both EC₅₀ and EC₉₅ were significantly lower in the female group than in the male group. EC₅₀, effective Ce of remifentanyl for suppression of emergence cough in 50% of patients; EC₉₅, effective Ce of remifentanyl for suppression of emergence cough in 95% of patients; CI, confidence interval.

propofol anesthesia for nasal surgery. The most notable difference between the two studies was the type of surgery; in addition, the proportion of male patients in the former study was 14%, whereas it was 62% in the latter. It is not surprising that the opioid requirements differed by sex, but other confounding factors should be excluded. Therefore, we conducted the present study for sex comparison only. Based on our results, the concentrations required for cough suppression during anesthetic emergence under the same clinical conditions were significantly different between sexes. One interesting point is that the cough sensitivity of females was lower than that of males. Dicipinigaitis and Rauf²⁷ reported that healthy women have a more sensitive cough reflex than healthy men. Nevertheless, the antitussive effect was

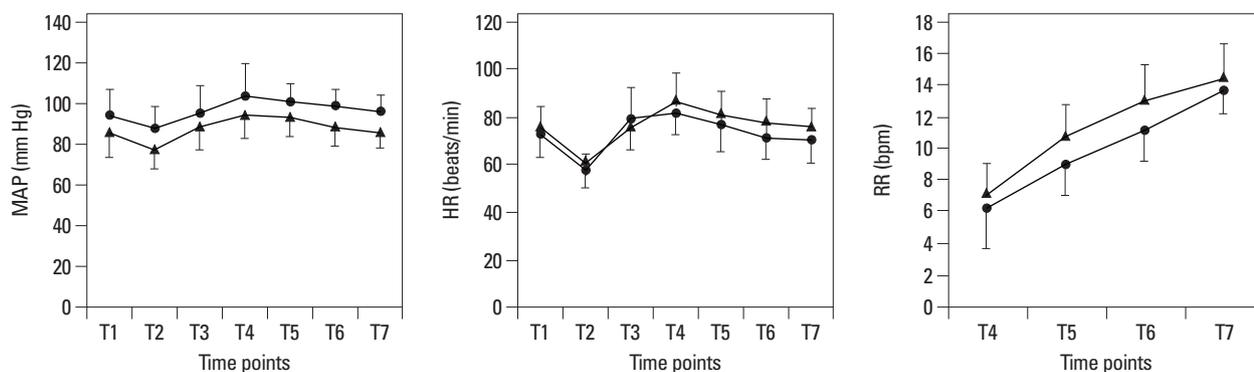


Fig. 3. MAP, HR, and RR were not significantly different in females (▲) and males (●) over time (p -values for MAP, HR, and RR=0.351, 0.082 and 0.277, respectively). The female MAP was significantly higher than that of males at the end of the surgery, 10 min after extubation, and before transfer out from the post-anesthetic care unit ($p=0.012$, $p<0.001$, and $p<0.001$, respectively). With regard to HR, there was no significant difference at any time point. RR increased continuously from each prior time point in both males and females: just after extubation to 5 min after extubation ($p<0.001$), 5 min after extubation to 10 min after extubation ($p<0.001$), and 10 min after extubation to before transfer out from the post-anesthetic care unit ($p<0.001$). MAP, mean arterial pressure; HR, heart rate; RR, respiratory rate; bpm, breaths per min.

achieved at a lower concentration of remifentanyl in females than in males. Male patients with a Ce of 2.0 ng/mL will definitely suffer from severe cough during emergence, while female patients with the same Ce will never cough.

In recent years, there has been growing interest in studying sex differences in anesthetic pharmacology. Compared to males, females are less sensitive to the anesthetic effects of propofol^{28,29} and ropivacaine,³⁰ but are more sensitive to vecuronium^{31,32} and rocuronium.³³ Of the anesthetic drugs, morphine is the most commonly studied with regard to sex differences, and several studies have shown that females are more sensitive to the analgesic effects of morphine than are males.^{34,35} Sex differences related to opioids are not restricted to analgesic properties, but are also present in other opioid-mediated responses, including the respiratory system. For instance, morphine has a greater effect on decreasing ventilatory response to carbon dioxide and hypoxic sensitivity in females than in males.¹⁸ Most of these studies did not measure morphine or morphine glucuronide concentration and pharmacokinetic differences could not be excluded; nevertheless, the major difference is thought to be pharmacodynamic.¹⁷

Remifentanyl TCI was derived from the results of pharmacokinetic and pharmacodynamic studies, which demonstrated that there are no sex differences related to pharmacokinetic or electroencephalogram (EEG) variables.²⁰ However, there is a lack of studies about the differential effect of remifentanyl based on sex,³⁶ and the clinical effects of opioids may differ from EEG variables. Experimental studies for mu-opioid receptor binding, measured by positron emission tomography (PET), illustrated that premenopausal females have significantly higher mu-receptor binding potential than males in the

cortical and sub-cortical areas.³⁷ Similarly, Zubieta, et al.³⁸ found that tonic experimental pain produced a greater decrease in mu-opioid receptor availability in several brain regions among men compared to women, apparently due to increased pain-induced binding of endogenous ligand to the receptor. Therefore, the authors suggested that the higher mu-opioid receptor availability among females might explain their increased analgesic responses to exogenous opioids. This pharmacodynamic difference can be expected in remifentanyl also, and our results did demonstrate a sex difference for cough suppression.

Several clinical studies have reported respiratory complications at concentrations over 2.5 ng/mL (Ce of remifentanyl),^{8,14} and our research did as well. Though the respiratory complications were not severe and resolved spontaneously or with minimal treatment, further studies regarding the safety of the suggested EC₉₅ are necessary for generalized application of high concentrations of remifentanyl.

There are some limitations to the present study. First, we did not measure plasma concentrations of remifentanyl. Rather, we calculated the predicted value using Minto's pharmacokinetic model, which is widely used in clinical settings with acceptable levels of bias and inaccuracy.³⁹ Second, the types of surgery included were limited. Thyroid surgery may involve more tracheal irritation than other surgeries, and therefore, caution is needed in extrapolating the findings of this study to other types of surgery.

In conclusion, when remifentanyl TCI is used for cough prevention during anesthetic emergence, patient sex may be an important factor for the determination of optimal remifentanyl concentrations. Overall, females require lower concentrations of remifentanyl for this purpose than do males.

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