

Planning for operating room efficiency and faster anesthesia wake-up time in open major upper abdominal surgery

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Abstract

Reducing anesthesia-controlled time (ACT) may improve operation room (OR) efficiency result from different anesthetic techniques. However, the information about the difference in ACT between desflurane (DES) anesthesia and propofol-based total intravenous anesthesia (TIVA) techniques for open major upper abdominal surgery under general anesthesia (GA) is not available in the literature.

This retrospective study uses our hospital database to analyze the ACT of open major upper abdominal surgery without liver resection after either desflurane/fentanyl-based anesthesia or TIVA via target-controlled infusion with fentanyl/propofol from January 2010 to December 2011. The various time intervals including waiting for anesthesia time, anesthesia time, surgical time, extubation time, exit from OR after extubation, total OR time, and postanesthetic care unit (PACU) stay time and percentage of prolonged extubation (≥ 15 minutes) were compared between these 2 anesthetic techniques.

We included data from 343 patients, with 159 patients receiving TIVA and 184 patients receiving DES. The only significant difference is extubation time, TIVA was faster than the DES group (8.5 ± 3.8 vs 9.4 ± 3.7 minutes; $P=0.04$). The factors contributed to prolonged extubation were age, gender, body mass index, DES anesthesia, and anesthesia time.

In our hospital, propofol-based TIVA by target-controlled infusion provides faster emergence compared with DES anesthesia; however, it did not improve OR efficiency in open major abdominal surgery. Older, male gender, higher body mass index, DES anesthesia, and lengthy anesthesia time were factors that contribute to extubation time.

Abbreviations: ACT = anesthesia-controlled time, BIS = bispectral index, BMI = body mass index, Ce = effect-site concentration, DES = desflurane, GA = general anesthesia, NDMR = nondepolarizing muscle relaxant, OR = operation room, PACU = postanesthetic care unit, TCI = target-controlled infusion, TIVA = total intravenous anesthesia.

Keywords: anesthesia-controlled time, desflurane, open major upper abdominal surgery, propofol

1. Introduction

Anesthesia-controlled time (ACT) and turnover time are 2 of the most important factors that regulate operation room (OR)

efficiency.^[1] Extubation time is of special interests to surgeons and anesthesiologists because it could be affected by different anesthetic agents or techniques.^[2–4] Prolonged extubation is an important factor that would decrease OR efficiency. Prolonged extubation time would cause slowing of work flow, having OR members staying idly waiting for extubation, and the surgeon have to wait longer for next operation. Surgeons always want patient quick to awaken.^[5] Accordingly, choosing appropriate anesthetic agents or techniques to avoid prolonged extubation is essential for anesthesiologists in order to improve the efficiency of OR. Dexter and Epstein^[6] recommended that recording extubation time and monitoring the incidence of prolonged extubation is very important especially at facilities that have at least 8 hours of cases and turnovers.

The ACT between total intravenous anesthesia (TIVA) with propofol and desflurane (DES) anesthesia was investigated, nevertheless, the results are controversial.^[4,7–15] The majority of these studies comparing the effects of different anesthesia regimens on OR efficiency have tended to focus on ambulatory or short-time surgery. As our best knowledge, we found no comparisons in different anesthetic techniques for the improvement of ACT in open major upper abdominal surgery under general anesthesia (GA). Moreover, different propofol delivery techniques such as target-controlled infusion (TCI) and syringe pump infusion were used in these studies and may lead to different results. The aim of our present study was to determine whether the use of TIVA with TCI system is more effective than DES anesthesia in reducing ACT in patients undergoing open major upper abdominal surgery.

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2. Methods

This retrospective study was approved by the Ethics Committee (TSGHIRB No: 100-05-168) of Tri-Service General Hospital, Taipei, Taiwan (Chairperson, Professor Pauling Chu) on August 29th, 2011. IRB allows waiving the requirement for obtaining informed consent, and patient records were anonymized and deidentified prior to analysis. The information was retrieved from the medical records and the electronic database of Tri-Service General Hospital (TSGH; Taipei, Taiwan, Republic of China). We enrolled 343 patients (American Society of Anesthesiology class I–III) who received elective open major upper abdominal surgery under TIVA with TCI or DES anesthesia from January 2010 to December 2011. Exclusion criteria include: body mass index (BMI) $>35 \text{ kg/m}^2$, liver resection, emergent surgeries, patient's age younger than 18 years, combined TIVA with inhalation anesthesia or epidural anesthesia, other inhalation anesthesia besides DES, patients were sent to the intensive care unit, or incomplete data. Other parameters included demographic data and American Society of Anesthesiology physical status.

There was no premedication before induction of anesthesia. Regular monitoring, such as noninvasive blood pressure, arterial line, electrocardiography (lead II), pulse oximetry, and end-tidal carbon dioxide (EtCO₂) pressure, was applied in each patient. Anesthesia was induced with fentanyl, propofol, and rocuronium in all patients. The patients were then intubated and maintained with propofol or DES and the analgesic fentanyl. In our common practice, we take patients to the postanesthetic care unit (PACU) after extubation and did not extubate in PACU.

In the TIVA group, anesthesia was induced using intravenous (i.v.) fentanyl (2 $\mu\text{g/kg}$) and 2% lidocaine (1.5 mg/kg). Continuous infusion of propofol (fresfol 1%) was delivered subsequently using Schneider kinetic model of TCI (Fresenius Orchestra Primea; Fresenius Kabi AG, Bad Homburg, Germany) with the effect-site concentration (Ce) of 4.0 $\mu\text{g/mL}$. Rocuronium (0.6 mg/kg) was administered when patients lost consciousness, followed by tracheal intubation. Anesthesia was maintained using TCI with propofol Ce 3 to 4 $\mu\text{g/mL}$ and an oxygen flow of 0.3 L/min. Repetitive bolus injections of cisatracurium and fentanyl were prescribed as required throughout the procedure.^[12,16]

In the DES group, the patients were induced with i.v. fentanyl (2 $\mu\text{g/kg}$), 2% lidocaine (1.5 mg/kg), and propofol (1.5–2 mg/kg). When patients lost consciousness, 0.6 mg/kg of rocuronium was administered, followed by endotracheal intubation. Anesthesia was maintained using 8% to 12% DES (inhaled concentration) in an oxygen flow of 300 mL/min under a closed system without nitrous oxide. Repetitive bolus injections of cisatracurium and fentanyl were prescribed as required throughout the procedure.^[12,16]

Maintenance of the Ce for the TCI with propofol and DES concentration was adjusted at the range of 0.2 $\mu\text{g/mL}$ and 0.5%, respectively, according to the hemodynamics. If 2 increments or decrements were unsuccessful, the range of Ce for TCI propofol and DES was increased to 0.5 $\mu\text{g/mL}$ or 2%, respectively. The EtCO₂ pressure was maintained at 35 to 45 mm Hg by adjusting the ventilation rate and maximum airway pressure. Once neuromuscular function returns, cisatracurium (2 mg, i.v.) was administered as required.^[12,16]

Ce of propofol or DES concentration was tapered to 2.0 $\mu\text{g/mL}$ or 5% respectively at the beginning of skin closure. At the last 5 stitches of surgery, propofol or DES was discontinued, but the oxygen flow was kept 300 mL/min. At the end of the skin closure, the lungs were ventilated with 100% oxygen at a fresh gas flow of

6 L/min. Reversal of neuromuscular function was achieved by administering neostigmine (0.03–0.04 mg/kg) with glycopyrrrolate (0.006–0.008 mg/kg) once spontaneous breathing returned to prevent residual paralysis. When the patient regained consciousness by name with spontaneous and smooth respiration, the endotracheal tube was removed and the patient was sent to the PACU for further care. An extubation time (from the end of skin closure until extubation) equal or longer than 15 minutes is considered prolonged extubation.^[17]

Data are presented as the mean and standard deviation, number of patients, or percentage. Demographic and perioperative variables were compared using Student *t* tests. Categorical variables were compared using chi-square test. Multivariable logistic regression analyses were performed to assess the association between variables contributed to prolonged extubation. The level of statistical significance was determined as $P < 0.05$.

3. Results

After excluded from the electrical record, another 56 patients were excluded from the analysis. Of those excluded, 20 patients received combined inhalation anesthesia with propofol and 15 patients received sevoflurane anesthesia, and 21 patients had BMI $>35 \text{ kg/m}^2$ (Fig. 1).

Our study included 343 patients, of which 184 received DES and 159 received TIVA anesthesia. Summary of surgical procedures was shown in Table 1. There was no significant difference in patient demographics (Table 2). The amount of opioids and nondepolarizing muscle relaxants (NDMRs) were significant higher in TIVA group than in DES group while reversal agents showed no significant difference between groups (Table 3). The emergence was faster for TIVA group than DES group (8.5 ± 3.8 vs 9.4 ± 3.7 minutes; $P = 0.04$). The waiting for anesthesia time, surgical time, anesthesia time, exit from OR after extubation, total OR time, PACU time, and the incidence of prolonged extubation were no difference between groups (Table 4).

The result of multivariable logistic regressions comparing prolonged extubation time between several variants in all patients is shown in Table 5. Age, gender, BMI, group, and anesthesia time were factors that contribute to extubation time. The results showed that patients with older age, male, higher BMI, DES anesthesia, and lengthy anesthesia time have slower emergence.

4. Discussion

The major findings in this retrospective study show that propofol-based TIVA by TCI reduced the extubation time relative to DES anesthesia. Although statistically significant differences were

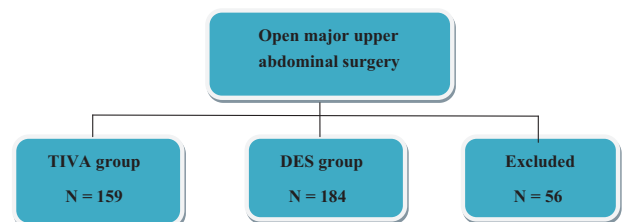


Figure 1. The flow diagram. DES=desflurane anesthesia, TIVA=total intravenous anesthesia.

Table 1**Summary of surgical procedures.**

Surgical procedure	DES (n)	TIVA (n)
Gastrectomy	95	86
Repair ventral hernia	21	25
Biliary track surgery	29	17
Exp Lap with gastrojejunostomy	35	25
Exp Lap with splenectomy	4	6

Data shown as number. DES=desflurane, Exp Lap=exploratory laparotomy, TIVA=total intravenous anesthesia.

found, 0.9 minutes reduction in extubation time in TIVA group suggested less clinical or economical effect on the ACT component of OR efficiency. In addition, we found that the factors of prolonged extubation are age, gender, BMI, DES group, and anesthesia time in patients undergoing open major upper abdominal surgery.

The 1st finding was consistent with several previous studies showing that GA using TCI system with propofol could achieve faster extubation than using DES in different surgeries.^[9-12,18] In our previous large case number retrospective studies, we showed that propofol-based TIVA by TCI reduced the extubation time were 1.8 and 5.4 minutes relative to DES in patients undergoing ophthalmic surgery^[12] and lengthy lumbar spine surgery.^[13] Because the awakening time can be predicted by TCI system.^[19] However, 4 studies compared DES anesthesia with propofol-based TIVA and failed to show any significant clinical difference in extubation in laparoscopic cholecystectomy,^[20] otological surgery,^[21] ear, nose, and throat surgery.^[22] These were different from our retrospective studies and other previous studies.^[9,11,12,18] The reason might be due to the DES maintenance flow rate of oxygen was different: 1 to 4 L/min versus 300 mL/min in our study. Using close circuit anesthesia in the DES group would also prolong the neuromuscular blockade and delay the extubation time.^[23] In another study, Dolk et al^[24] had reported that there were shorter extubation time for DES anesthesia compared with propofol delivered by TCI in knee surgery. The difference may cause by using nitrous oxide as an adjuvant to anesthetics, which reduce the requirement of DES during the maintenance period and facilitate early emergence.

Epstein et al^[25] concluded that prolonged extubation time should result in increased variable costs. Another study conducted by the same group demonstrated that the mean time from end of surgery to exit OR is at least 12.6 minutes longer in cases with prolonged extubation and that the percentage of cases for which the extubation was prolonged among anesthesia for intraperitoneal procedures in upper abdomen was 15.1% ± 0.6%.^[6] In our present study, the percentage of prolonged

Table 2**Patient's characteristics.**

	Group DES (n=184)	Group TIVA (n=159)	P
ASA II/III	151/33	126/33	0.51
Gender (M/F)	68/116	63/96	0.61
Age, year/o	68.0 ± 7.3	67.9 ± 7.8	0.91
Height, cm	168.4 ± 5.1	168.0 ± 4.7	0.47
Weight, kg	69.4 ± 7.1	69.7 ± 5.7	0.68
BMI	24.5 ± 2.2	24.7 ± 1.7	0.29

Data shown as mean ± SD or number. ASA=American Society of Anesthesiology, BMI=body mass index, DES=desflurane, SD=standard deviation, TIVA=total intravenous anesthesia.

Table 3**The amount of opioid, NDMRs, and reversal agents during surgical periods between DES and TIVA group.**

	Group DES (n=184)	Group TIVA (n=159)	P
Fentanyl, µg	168.5 ± 24.2	269.8 ± 24.5	<0.001
Cisatracurium, mg	13.2 ± 5.9	20.4 ± 8.0	<0.001
Neostigmine, mg	2.28 ± 0.45	2.27 ± 0.45	0.802
Glycopyrrolate, mg	0.46 ± 0.09	0.45 ± 0.09	0.802

Data shown as mean ± SD or number. DES=desflurane, NDMR=nondepolarizing muscle relaxant, SD=standard deviation, TIVA=total intravenous anesthesia.

extubation in DES group is 9.8%, while the percentage of prolonged extubation in TIVA group was 9.4% (Table 4). There was no significant difference in the incidence of prolonged extubation between TIVA and DES groups, which might be due to the similar BMI, gender, surgical time, and anesthesia time.

There were studies that investigated the confounding risk factors of prolonged extubation which included prone position, prolonged surgical time, significant blood loss, and larger volume of crystalloid and colloid infusion.^[6,26,27] Our previous study reported that DES anesthesia, lengthy anesthesia time, higher BMI, and shorter surgical time contribute to slower emergence in gynecologic laparoscopic surgery.^[14] In addition, Chan et al^[19] demonstrated that the confounding factors that predicted awaken under TCI with propofol are age, gender, and times of surgery and anesthesia (total consumption dose of propofol and fentanyl) in assortments of surgeries. In this study, old age, male gender, higher BMI, and lengthy anesthesia time resulting in prolonged extubation, which was consistent with our previous studies.^[14,19] Nevertheless, we showed that surgical time did not contribute to prolonged extubation, it might be due to the prolonged duration of neuromuscular relaxants resulting from the close circuit anesthesia.^[23]

Previous studies also implied that longer-than-average anesthesia times strongly influence the academic anesthesiology departments by increasing the staffing costs and decreasing hourly productivity.^[28,29] There is evidence that propofol may accumulate and redistributed from the fatty tissue and muscle to the plasma, which leads to delay recovery by using syringe pump with continuous infusion in adult.^[30] However, TCI could maintain the steady concentration of propofol instead of flow rate and predict awake time. Therefore, the effect of accumulation and redistribution of propofol on extubation should be less than syringe pump with continuous infusion of propofol. The inhaled DES is redistributed in the fatty tissue and muscle and

Table 4**OR time measurement between DES and TIVA group.**

	Group DES (n=184)	Group TIVA (n=159)	P
Waiting for anesthesia time, minute	7.3 ± 3.2	7.2 ± 2.8	0.75
Surgical time, minute	180.9 ± 85.4	182.3 ± 77.9	0.88
Anesthesia time, minute	211.4 ± 87.5	213.1 ± 79.9	0.86
Extubation time, minute	9.4 ± 3.7	8.5 ± 3.8	0.04
Exit from OR after extubation, minute	5.5 ± 3.1	5.7 ± 3.2	0.60
Total OR time, minute	224.2 ± 87.5	225.9 ± 79.6	0.85
PACU time, minute	57.4 ± 14.4	58.3 ± 15.1	0.59
Prolonged extubation (≥15 minutes)	18 (9.8)	15 (9.4)	0.91

Data shown as mean ± SD or number (percentage). DES=desflurane, OR=operation room, PACU=postanesthetic care unit, SD=standard deviation, TIVA=total intravenous anesthesia.

Table 5
Multivariable linear regression analyses of variables associated with extubation time in all patients (n=343).

	β	95% CI	P
Age	0.17	0.13–0.21	<0.001
Gender	1.14	0.63–1.66	<0.001
BMI	0.20	0.08–0.31	0.001
Group	–0.88	–1.30–0.46	<0.001
Surgical time	–0.01	–0.03–0.01	0.37
Anesthesia time	0.03	0.01–0.05	0.003

β , difference between each variant using emergence time as dependent variable, group=DES=0, TIVA=1. P values <0.05 were considered significant. BMI=body mass index, CI=confidence interval, DES=desflurane, TIVA=total intravenous anesthesia.

delayed emergence once the lengthy anesthesia time.^[18] Therefore, monitoring anesthetic depth such as bispectral index (BIS) to keep the hypnotic level within the recommended range improves anesthetic delivery and postoperative recovery from relatively deep anesthesia.^[31] In addition, we suggested prescribed BIS for patients were elder, higher BMI, and lengthy anesthesia time.

The amount of opioid and NDMRs in DES group was significantly lower than in TIVA group during surgical periods. It is reasonable because volatile anesthetics may increase the potency of NDMRs^[32] and demonstrate synergy effects with opioids.^[33] In addition, not until spontaneous breathing returned were the reversal agents administered. Therefore, we believed the final neuromuscular blockade status and amount of reversal agents given were matched between groups.

Our previous studies showing that GA using TCI system with propofol could achieve faster extubation than using DES anesthesia in different surgeries.^[9–14,18] Different anesthetic manipulations before emergence in various types of surgical procedures might explain the differences in findings. For example, in breast^[11] and gynecologic surgery,^[14] propofol was adjusted to a Ce of 2.0 $\mu\text{g/mL}$ and the vapor of DES was changed to 5.0% in the beginning of wound closure. After gauze coverage, propofol and DES were discontinued and lungs were ventilated with 100% oxygen at a gas flow of 6L/min. In ophthalmic surgery,^[12] DES or propofol was discontinued after the surgery, and the lungs were ventilated with 100% oxygen at a fresh gas flow of 6L/min. In spine surgery, we discontinued DES or propofol at the end of the operation or at the last 3 stitches of surgery. After turning the patients to supine position, the lungs were ventilated with 100% oxygen at a fresh gas flow of 6L/min.^[10,13] In addition, we used closed-circuit anesthesia in the DES patients, which would prolong neuromuscular blockade and contribute to delay emergence.^[23]

There are many limitations in the study. The first was our study is a retrospective study. Considering comparability and standardization of study groups, a retrospective study may contribute to bias. Although the choice of anesthetic management was not randomly allocated but rather by the availability of the TCI devices, the results showed no difference in the characteristics of the patients between 2 groups. The study, performed under clinical conditions and provided large sample size, reflects more precisely the clinical relevant benefit. Second, we excluded liver resection due to liver dysfunction resulting in major impacts on the pharmacokinetics and pharmacodynamics of anesthetics and the recovery from TIVA and inhalation anesthesia is delayed in hepatectomy patients.^[34,35] Third, we did not compare the effect of body temperature on extubation time, because hypothermia

may delay awakening.^[36] However, in our cases, we used the patient warming system including fluid warming kit and convective air warming system to keep their core temperature $\geq 35^\circ\text{C}$ perioperatively. Fourth, we excluded patients with BMI $> 35\text{ kg/m}^2$, because obesity may lead to prolonged extubation^[36] and it is the limitation in Schnider model of TCI machine. Fifth, we did not include patients receiving Whipple operation and blood loss $> 1500\text{ mL}$ because larger volume of fluid infusion may be the risk factor for delayed extubation.^[26] Finally, we did not use BIS in our common practice. But the depth of anesthesia was monitored by the experienced anesthesiologist, and our percentage of prolonged extubation was 9.6% less than overall 15.4% reported by a previous study.^[6]

Although anesthesia has the capacity to reduce operating room efficiency, a well-planned anesthesia technique like propofol-based TIVA or DES does not impede efficiency even after lengthy invasive surgery and even in an academic teaching hospital setting. Therefore, other factors (adequate preoperative patient workup, hospital transport, preparation in the preanesthesia unit, surgical time, patient comorbidities, etc.) need to be considered.

References

- Junger A, Klasen J, Hartmann B, et al. Shorter discharge time after regional or intravenous anaesthesia in combination with laryngeal mask airway compared with balanced anaesthesia with endotracheal intubation. *Eur J Anaesthesiol* 2002;19:119–24.
- Apfelbaum JL, Grasela TH, Hug CCJr, et al. The initial clinical experience of 1819 physicians in maintaining anesthesia with propofol: characteristics associated with prolonged time to awakening. *Anesth Analg* 1993;77:S10–4.
- Dexter F, Bayman EO, Epstein RH. Statistical modeling of average and variability of time to extubation for meta-analysis comparing desflurane to sevoflurane. *Anesth Analg* 2010;110:570–80.
- Wachtel RE, Dexter F, Epstein RH, et al. Meta-analysis of desflurane and propofol average times and variability in times to extubation and following commands. *Can J Anaesth* 2011;58:714–24.
- Vitez TS, Macario A. Setting performance standards for an anesthesia department. *J Clin Anesth* 1998;10:166–75.
- Dexter F, Epstein RH. Increased mean time from end of surgery to operating room exit in a historical cohort of cases with prolonged time to extubation. *Anesth Analg* 2013;117:1453–9.
- Juvin P, Servin F, Giraud O, et al. Emergence of elderly patients from prolonged desflurane, isoflurane, or propofol anesthesia. *Anesth Analg* 1997;85:647–51.
- Gupta A, Stierer T, Zuckerman R, et al. Comparison of recovery profile after ambulatory anesthesia with propofol, isoflurane, sevoflurane and desflurane: a systematic review. *Anesth Analg* 2004;98:632–41. table of contents.
- Hong HC, Kuo CP, Ho CC, et al. Cost analysis of three anesthetic regimens under auditory evoked potentials monitoring in gynecologic laparoscopic surgery. *Acta Anaesthesiol Taiwan* 2007;45:205–10.
- Chan SM, Horng HH, Huang ST, et al. Drug cost analysis of three anesthetic regimens in prolonged lumbar spinal surgery. *J Med Sci* 2009;29:75–80.
- Chen JL, Chen YF, Chen YW, et al. Do anesthetic techniques affect operating room efficiency? Comparison of target-controlled infusion of propofol and desflurane anesthesia in breast cancer surgery. *J Med Sci* 2013;33:205–10.
- Wu ZF, Jian GS, Lee MS, et al. An analysis of anesthesia-controlled operating room time after propofol-based total intravenous anesthesia compared with desflurane anesthesia in ophthalmic surgery: a retrospective study. *Anesth Analg* 2014;119:1393–406.
- Lu CH, Wu ZF, Lin BF, et al. Faster extubation time with more stable hemodynamics during extubation and shorter total surgical suite time after propofol-based total intravenous anesthesia compared with desflurane anesthesia in long-term lumbar spine surgery. *J Neurosurg Spine* 2016;24:268–74.
- Lai HC, Chan SM, Lin BF, et al. Analysis of anesthesia-controlled operating room time after propofol-based total intravenous anesthesia

- compared with desflurane anesthesia in gynecologic laparoscopic surgery: a retrospective study. *J Med Sci* 2015;35:157–61.
- [15] Liu FL, Cherng YG, Chen SY, et al. Postoperative recovery after anesthesia in morbidly obese patients: a systematic review and meta-analysis of randomized controlled trials. *Can J Anaesth* 2015;62:907–17.
- [16] Lin BF, Ju DT, Cherng CH, et al. Comparison between intraoperative fentanyl and tramadol to improve quality of emergence. *J Neurosurg Anesthesiol* 2012;24:127–32.
- [17] Agoliati A, Dexter F, Lok J, et al. Meta-analysis of average and variability of time to extubation comparing isoflurane with desflurane or isoflurane with sevoflurane. *Anesth Analg* 2010;110:1433–9.
- [18] Lu CH, Yeh CC, Huang YS, et al. Hemodynamic and biochemical changes in liver transplantation: a retrospective comparison of desflurane and total intravenous anesthesia by target-controlled infusion under auditory evoked potential guide. *Acta Anaesthesiol Taiwan* 2014;52:6–12.
- [19] Chan SM, Lee MS, Lu CH, et al. Confounding factors to predict the awakening effect-site concentration of propofol in target-controlled infusion based on propofol and fentanyl anesthesia. *PLoS One* 2015;10:e0124343.
- [20] Grundmann U, Silomon M, Bach F, et al. Recovery profile and side effects of remifentanyl-based anaesthesia with desflurane or propofol for laparoscopic cholecystectomy. *Acta Anaesthesiol Scand* 2001;45:320–6.
- [21] Fombeur PO, Tilleul PR, Beaussier MJ, et al. Cost-effectiveness of propofol anesthesia using target-controlled infusion compared with a standard regimen using desflurane. *Am J Health Syst Pharm* 2002;59:1344–50.
- [22] Mahli A, Coskun D, Karaca GI, et al. Target-controlled infusion of remifentanyl with propofol or desflurane under bispectral index guidance: quality of anesthesia and recovery profile. *J Res Med Sci* 2011;16:611–20.
- [23] Yeh CC, Kong SS, Chang FL, et al. Closed-circuit anesthesia prolongs the neuromuscular blockade of rocuronium. *Acta Anaesthesiol Sin* 2003;41:55–60.
- [24] Dolk A, Cannerfelt R, Anderson RE, et al. Inhalation anaesthesia is cost-effective for ambulatory surgery: a clinical comparison with propofol during elective knee arthroscopy. *Eur J Anaesthesiol* 2002;19:88–92.
- [25] Epstein RH, Dexter F, Brull SJ. Cohort study of cases with prolonged tracheal extubation times to examine the relationship with duration of workday. *Can J Anaesth* 2013;60:1070–6.
- [26] Li F, Gorji R, Tallarico R, et al. Risk factors for delayed extubation in thoracic and lumbar spine surgery: a retrospective analysis of 135 patients. *J Anesth* 2014;28:161–6.
- [27] Chan WH, Lee MS, Lin C, et al. Comparison of anesthesia-controlled operating room time between propofol-based total intravenous anesthesia and desflurane anesthesia in open colorectal surgery: a retrospective study. *PLoS One* 2016;11:e0165407.
- [28] Abouleish AE, Prough DS, Whitten CW, et al. Increasing the value of time reduces the lost economic opportunity of caring for surgeries of longer-than average times. *Anesth Analg* 2004;98:1737–42.
- [29] Abouleish AE, Prough DS, Zornow MH, et al. The impact of longer-than-average anesthesia times on the billing of academic anesthesiology departments. *Anesth Analg* 2001;93:1537–43.
- [30] Levitt DG, Schnider TW. Human physiologically based pharmacokinetic model for propofol. *BMC Anesthesiol* 2005;5:4.
- [31] Punjasawadwong Y, Phongchiewboon A, Bunchungmongkol N. Bispectral index for improving anaesthetic delivery and postoperative recovery. *Cochrane Database Syst Rev* 2014;6:CD003843.
- [32] Paul M, Fokt RM, Kindler CH, et al. Characterization of the interactions between volatile anesthetics and neuromuscular blockers at the muscle nicotinic acetylcholine receptor. *Anesth Analg* 2002;95:362–7.
- [33] Hendrickx JF, Eger EI, Sonner JM, et al. Is synergy the rule? A review of anesthetic interactions producing hypnosis and immobility. *Anesth Analg* 2008;107:494–506.
- [34] Murata F, Iwade M, Hidano G, et al. Recovery from propofol anesthesia is delayed in hepatectomy patients due to altered pharmacodynamics. *Masui* 2006;55:150–7.
- [35] Murata F, Iwade M, Nagata O, et al. Recovery from sevoflurane anesthesia delayed in hepatectomy patients due to influence of operation. *Masui* 2007;56:650–6.
- [36] Kleine S, Hofmeister E, Egan K. Multivariable analysis of anesthetic factors associated with time to extubation in dogs. *Res Vet Sci* 2014;97:592–6.