

# 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 lengthy lumbar spine surgery

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**OBJECTIVE** Anesthesia techniques can contribute to the reduction of anesthesia-controlled time and may therefore improve operating room efficiency. However, little is known about the difference in anesthesia-controlled time between propofol-based total intravenous anesthesia (TIVA) and desflurane (DES) anesthesia techniques for prolonged lumbar spine surgery under general anesthesia.

**METHODS** A retrospective analysis was conducted using hospital databases to compare the anesthesia-controlled time of lengthy (surgical time > 180 minutes) lumbar spine surgery in patients receiving either TIVA via target-controlled infusion (TCI) with propofol/fentanyl or DES/fentanyl-based anesthesia, between January 2009 and December 2011. A variety of time intervals (surgical time, anesthesia time, extubation time, time in the operating room, postanesthesia care unit [PACU] length of stay, and total surgical suite time) comprising perioperative hemodynamic variables were compared between the 2 anesthesia techniques.

**RESULTS** Data from 581 patients were included in the analysis; 307 patients received TIVA and 274 received DES anesthesia. The extubation time was faster ( $12.4 \pm 5.3 \text{ vs } 7.0 \pm 4.5 \text{ minutes}$ , p < 0.001), and the time in operating room and total surgical suite time was shorter in the TIVA group than in the DES group ( $326.5 \pm 57.2 \text{ vs } 338.4 \pm 69.4 \text{ minutes}$ , p = 0.025; and 402.6 ± 60.2 vs 414.4 ± 71.7 minutes, p = 0.033, respectively). However, there was no statistically significant difference in PACU length of stay between the groups. Heart rate and mean arterial blood pressure were more stable during extubation in the TIVA group than in the DES group.

**CONCLUSIONS** Utilization of TIVA reduced the mean time to extubation and total surgical suite time by 5.4 minutes and 11.8 minutes, respectively, and produced more stable hemodynamics during extubation compared with the use of DES anesthesia in lengthy lumbar spine surgery.

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**KEY WORDS** TIVA; propofol; desflurane; lengthy spine surgery; extubation; lumbar

PERATING room efficiency is regulated by several factors; the 2 most important factors are anesthesiacontrolled time and turnover time.<sup>13</sup> The interval between the end of surgery and extubation is of particular interest to surgeons and anesthesia care providers because it is affected by anesthetic agents.<sup>1,4,19,20</sup> Anesthesiologists associate prolonged tracheal extubation time with poor re-

covery from anesthesia.<sup>1</sup> When surgeons score anesthesiologists' attributes on a scale from 0 ("not important") to 4 ("a factor that would make me switch groups/hospitals"), the average score for "patient quick to awaken" is 3.9.<sup>19</sup>

The majority of studies comparing the effects of different anesthesia regimens on operating room efficiency have tended to focus on relatively short and painless surgery

**ABBREVIATIONS** DES = desflurane; LOS = length of stay; MBP = mean blood pressure; PACU = postanesthesia care unit; TCI = target-controlled infusion; TIVA = total intravenous anesthesia.

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(e.g., day-case surgery). Recently, a meta-analysis comparing operating room recovery times for desflurane (DES) and propofol reported that, relative to propofol, DES proportionally reduced the mean time to extubation and time to follow commands (21% and 23%, respectively).<sup>20</sup> However, only 2 studies<sup>9,14</sup> of prolonged anesthesia or surgery duration were included in this meta-analysis. In our previous prospective study, patients undergoing lengthy procedures were found to recover more quickly after total intravenous anesthesia (TIVA) via a target-controlled infusion (TCI) system, than with volatile anesthesia.<sup>2</sup> The present study aimed to determine whether the use of TIVA with TCI is more effective than DES anesthesia in reducing anesthesia-controlled operating room time in patients undergoing lengthy lumbar spine surgery.

### **Methods**

This study was approved by the ethics committee of Tri-Service General Hospital, Taipei, Taiwan. Medical records and electronic databases were collected from this hospital and were reviewed for all patients undergoing elective lengthy lumbar spine surgery (surgical time > 180 minutes), including posterolateral fusion and pedicle screw fixation (excluding patients with a deformity diagnosis), between January 2010 and December 2011. A total of 581 patients who received TIVA or DES anesthesia were included in this study. Exclusion criteria were patients younger than 18 years, emergency surgeries, combined propofol and DES anesthesia, non-DES inhalation anesthesia, failure to extubate, no postanesthesia care unit (PACU) stay, and incomplete data. Other parameters extracted from the electronic databases and medical records included demographic data, American Society of Anesthesiology physical status class, intraoperative fentanyl dosage, and perioperative hemodynamic parameters.

For the purposes of this study, the following times (in minutes) were calculated: 1) surgical time (incision to surgical completion and application of dressings), 2) anesthesia time (initiation of anesthesia to extubation), 3) extubation time (completion of surgery and dressing application to extubation), 4) time in the operating room (arrival in the operating room to departure from the operating room), 5) length of stay (LOS) in the PACU (time from arrival in the PACU to discharge from the PACU to the general ward), and 6) total surgical suite time (arrival in the operating room to discharge from the PACU to the general ward).

### **Patient Groups**

There was no administration of medication prior to induction of anesthesia; however, regular monitoring, such as electrocardiography (lead II) and measurements of pulse oximetry, noninvasive blood pressure, respiratory rate, and end-tidal carbon dioxide pressure, were performed. The selection of anesthesia was based on the attending anesthesiologist's preference, and anesthesia management was performed by the same anesthesiologist according to clinical practice demands. In all patients, anesthesia was induced with propofol and fentanyl. The patients were then intubated and maintained with the anesthetics DES or propofol and the analgesic fentanyl.

### Anesthesia

TIVA was induced with fentanyl (2  $\mu$ g/kg) and lidocaine (2%, 1.5 mg/kg). Following this, continuous infusion of propofol (Fresenius 1%) was initiated using a TCI system programmed with the Schneider model (Fresenius Orchestra Primea, Fresenius Kabi AG) at an effective target concentration of 4  $\mu$ g/ml. Rocuronium (0.6 mg kg<sup>-1</sup>) was administered when patients lost consciousness, followed by tracheal intubation. Anesthesia was maintained using TCI with propofol effective concentration 3–4  $\mu$ g/ ml and an oxygen flow of 0.3 L/min. Repetitive bolus injections of rocuronium and fentanyl were prescribed as required throughout the procedure.

In the DES group, anesthesia was induced with fentanyl (2  $\mu$ g/kg), lidocaine (2%, 1.5 mg/kg), and propofol (2 mg/kg). After loss of consciousness, rocuronium (0.6 mg/kg) was administered, and tracheal intubation was performed. Anesthesia was maintained with 8%–12% DES in an oxygen flow of 300 ml/min under a closed system without nitrous oxide, and repetitive bolus injections of rocuronium and fentanyl were prescribed as required throughout the procedure.

Maintenance of the effective concentration for the TCI propofol and DES concentrations was adjusted at a range of 0.2 µg/ml and 0.5%, respectively, according to the hemodynamics. If 2 increments or decrements were unsuccessful in stabilizing the hemodynamics, the effective concentration ranges for the TCI propofol and DES concentrations were increased to 0.5 µg/ml and 2%, respectively. The ventilation rate and maximum airway pressure were adjusted to maintain end-tidal carbon dioxide pressure at 35–45 mm Hg. Either cisatracurium (2 mg intravenously) or rocuronium (10 mg intravenously) was administered as required by the return of neuromuscular function.

At the end of the operation, DES or propofol was discontinued, and the lungs were ventilated with 100% oxygen at a fresh gas flow of 6 L/min. When the patient regained consciousness, with spontaneous and smooth respiration, the endotracheal tube was removed.

#### Statistical Analysis

Categorical and continuous variables were presented as proportions and means  $\pm$  SD, respectively. The differences in study variables between DES and TIVA were tested using the Student t-test or chi-square test where appropriate. In the multivariable analysis, patient characteristics (sex, age, height, and weight) were adjusted using linear regression analysis to obtain estimates of differences in all of the study variables. Since some study variables were the linear combination of each other, further analyses of individual variables were separated into 2 parts: 1) total surgical suite time, levels of fusion, and fentanyl dosage; and 2) anesthesia time, extubation time, PACU LOS, levels of fusion, and fentanyl dosage. These analyses were also adjusted according to the physiological state. The differences in heart rate and mean blood pressure (MBP) between the groups were also estimated. This study assumed that there were no interaction effects between any variables on any dependent variables. Statistical analyses were performed with R 3.0.1 software, and the level of statistical significance was determined as p < 0.05.

# Results

A total of 270 patients were excluded from the analysis, of whom 85 received combined inhalation anesthesia with propofol, 126 received sevoflurane anesthesia, and 59 had incomplete data (Fig. 1). Of the 581 patients included in this study, 307 received TIVA and 274 received DES anesthesia. There was no statistically significant difference in patient demographics between the groups, and the frequency of more than 2 levels of fusion was similar between them (Table 1).

Surgical and anesthesia time, and LOS in the PACU were not statistically significantly different between groups (Table 1). The extubation time was faster (12.4  $\pm$ 5.3 vs 7.0  $\pm$  4.5 minutes, p < 0.001), and the time in the operating room and surgical suites was shorter (326.5  $\pm$ 57.2 vs 338.4  $\pm$  69.4 minutes, p = 0.025; and 402.6  $\pm$  60.2 vs. 414.4  $\pm$  71.7 minutes, p = 0.033, respectively) in the TIVA group than in the DES group (Table 1). The dose of intraoperative fentanyl was significantly larger in the TIVA group than in the DES group  $(268.2 \pm 61.4 \text{ vs } 228.5)$  $\pm$  63.1 µg, p < 0.001; Table 1). Comparisons of surgical status by multiple variable analyses also demonstrated similar results between the groups (Table 2). Model 3 (Table 2) presents these results adjusted by other operating room times. Out of all the operating room times analyzed, the largest significant difference between the DES and TIVA groups was extubation time. Therefore, the differences in time in operating room and total surgical suite time between groups were mainly due to extubation time.

The heart rate and MBP during extubation in the TIVA group were lower than those in the DES group (91.7  $\pm$  20.0 vs 102.8  $\pm$  18.5 bpm, p < 0.001 and 108  $\pm$  17.3 vs 115.9  $\pm$  13.9 mm Hg, p < 0.001, respectively; Table 1). Changes in heart rate and MBP in the TIVA group were also lower than those in the DES group (17.1  $\pm$  20.1 vs 29.5  $\pm$  18.5 bpm, p < 0.001 and 5.7  $\pm$  19.0 vs 14.4  $\pm$  13.8 mm Hg, p < 0.001, respectively; Table 1). Multivariable analyses

revealed similar effects of physiological status between groups (Table 3).

To ensure unbiased results, the confounding effect of patient characteristics (sex, age, height, and weight), various time intervals, levels of fusion, and fentanyl dosage were investigated. The results revealed a similar frequency of more than 2 levels of fusion and no statistically significant difference in patient demographics between the DES and TIVA groups (Table 1). This indicated that the selection of anesthesia techniques was unaffected by levels of fusion. Furthermore, there were no intergroup interactive effects between any variables on any dependent variables for each time, and clinical characteristics were similar compared with the unadjusted results (Tables 2 and 3). These analyses adjusted for the physiological state and therefore guaranteed robust and unbiased results.

# Discussion

In this retrospective study, propofol-based TIVA by TCI reduced the mean time to extubation (by 5.4 minutes) and total surgical suite time (by 11.8 minutes) and produced more stable hemodynamics during extubation, relative to DES anesthesia in patients undergoing lengthy lumbar spine surgery. These findings contradict the limited existing information obtained from previous randomized trials. However, they importantly reveal that extubation time and total surgical suite time significantly differ according to duration of surgery and anesthetic drug selection.

Reductions in operating room time reduce direct labor costs either when the operating room has overused time or when there is more than 8 hours of appropriate staffing planned for the operating room and the staffing can be reduced to 8 hours.<sup>6,17,18</sup> Facilities that consistently use operating rooms for more than 8 hours daily will benefit from the largest direct cost reductions. Among these operating rooms, each 1-minute reduction in operating room time

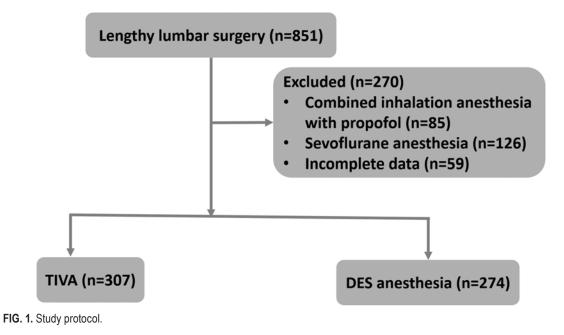


TABLE 1. Patient characteristics and various time intervals in both group	tics and various time intervals in both grou	variou	and	cteristics	chara	. Patient	TABLE 1	
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Variable	DES (n = 274)	TIVA (n = 307)	p Value†
Female sex	45.3%	46.3%	0.875
Mean age, yrs	59.9 ± 10.0	63.8 ± 15.4	0.757
Mean height, cm	164.5 ± 9.0	164.4 ± 8.9	0.919
Mean weight, kg	70.2 ± 12.6	70.3 ± 12.5	0.903
Mean arrival in OR to start of anesthesia, mins	9.0 ± 3.9	7.9 ± 3.9	0.001
Mean surgical time, mins	253.1 ± 64.1	254.5 ± 53.0	0.775
Mean anesthesia time, mins	306.6 ± 68.4	302.3 ± 56.3	0.417
Mean extubation time, mins	12.4 ± 5.3	7.0 ± 4.5	<0.001
Mean extubation to departure from OR, min	10.5 ± 5.4	$9.3 \pm 4.9$	0.004
Mean time in OR, mins	338.4 ± 69.4	326.5 ± 57.2	0.025
Mean PACU LOS, mins	76.0 ± 15.8	76.0 ± 16.8	0.953
Mean total surgical suite time, mins	414.4 ± 71.7	402.6 ± 60.2	0.033
Mean fentanyl, µg	228.5 ± 63.1	268.2 ± 61.4	<0.001
Mean baseline HR, bpm	73.3 ± 14.3	74.6 ± 15.3	0.299
Mean baseline MBP, mm Hg	101.4 ± 11.9	102.4 ± 10.8	0.337
Mean max HR during extubation, bpm	102.8 ± 18.5	91.7 ± 20.0	<0.001
Mean max MBP during extubation, mm Hg	115.9 ± 13.9	108 ± 17.3	<0.001
Mean ΔHR, bpm	29.5 ± 18.5	17.1 ± 20.1	<0.001
Mean ΔMBP, mm Hg	14.4 ± 13.8	5.7 ± 19.0	<0.001
PLF & PSF, >2 levels	32.5%	34.9%	0.546

HR = heart rate; MBP = mean blood pressure; ΔHR = maximal heart rate during extubation – baseline heart rate; ΔMBP = maximal MBP during extubation – baseline MBP; OR = operating room; PLF = posterolateral fusion; PSF = pedicle screw fixation.

\* The baseline patient data indicate the measurements before the start of anesthesia. Mean data are shown as mean ± SD.

† p values < 0.05 were considered significant.

results in an overall 1.1- to 1.2-minute reduction in regularly scheduled labor costs.<sup>6,7</sup> Consequently, small reductions in operating room time, achieved through the reduction of extubation and total surgical suite time, as reported in this study (in which the operating room workday in this study exceeded 8 hours), can be reasonably considered to be of economic benefit. Additionally, an intangible value of time saved may be achieved from more predictable recovery (e.g., fewer frustrated surgeons complaining of delays in beginning the next case).

Differences in operating room recovery times between anesthetic agents have been extensively studied because they can limit operating room throughput; this is based on data showing that nonanesthesia operating room personnel must wait for the patient to be extubated during emergence from anesthesia in the majority of cases (> 66%).<sup>8,16</sup> Previously, Masursky et al.<sup>17</sup> demonstrated that longer times to extubation are associated with an increased risk of at least one person waiting or being idle in the operating room (slowing the workflow). In addition, cases with prolonged tracheal extubation times have longer times from operating room exit to the start of the surgeon's next case in the operating room.<sup>4</sup> Therefore, selection of an anesthetic technique associated with faster extubation is associated with rapid operating room workflow and reductions in the time from operating room exit to the start of the surgeon's next case.<sup>4,17</sup> An anesthesia technique with shorter extubation times would decrease waiting time for the operating room staff and reduce the time from the end of surgery to operating room exit.<sup>5</sup>

Extubation times differ significantly among anesthetic drugs.4,15,20 Propofol has become popular for general anesthesia, especially in the ambulatory setting. It is frequently used in combination with remifentanil since both drugs have been reported to enable rapid emergence from anesthesia and early return to normal activities.<sup>10,15</sup> Until recently, remifentanil was unavailable in Taiwan. However, the duration of emergence from anesthesia has been shown to be significantly shorter with TIVA than with inhalation anesthesia in breast cancer and gynecological laparoscopic surgeries.<sup>3,12</sup> In addition, in a previous prospective study, a propofol and fentanyl combination was found to be cost-saving and resulted in faster emergence and extubation in lengthy spinal surgery when compared with DES and sevoflurane anesthesia.2 Furthermore, the use of TIVA has been shown to reduce the mean time to extubation by at least 9% and duration in PACU by more than 1% compared with the use of DES anesthesia in ophthalmic surgery.22

In this retrospective study of patients undergoing lengthy lumbar spine surgery, propofol-based TIVA by TCI was also found to reduce the mean time to extubation and total surgical suite time relative to DES. Our findings contradict those of a recent meta-analysis<sup>20</sup> comparing the operating room recovery time of DES with that of propofol. The discrepancy in these findings is most likely due to differences

		Model 1			Model 2			Model 3			Model 4			Model 5	
Variable	β	SE	p Value	В	SE	p Value	g	SE	p Value	β	SE	SE p Value	g	SE	p Value
Arrival in OR to start of anesthesia	-1.10	0.32	0.001				-1.45	0.39	<0.001						
Surgical time	1.24	4.87	0.799												
Anesthesia time	-4.36	5.19	0.401				-7.10	6.17	0.250				-7.86	4.68	0.093
Extubation time	-5.33	0.41	<0.001				-5.60	0.43	<0.001				-5.34		<0.001
Extubation to departure from OR	-1.21	0.43	0.005				-1.43	0.52	0.006						
Time in OR	-12.00	5.27	0.023												
PACU duration	0.02	1.36	0.990				0.09	1.65	0.958				0.50	1.46	0.732
Total surgical suite time	-11.98	5.49	0.029	-21.66	5.62	<0.001				-20.33	3.66	<0.001			
Fentanyl	39.12	5.02	<0.001	41.60	4.92	<0.001	42.38	5.72	<0.001	43.10	4.95	<0.001	40.88	4.94	<0.001

TARLE 2. Commarisons of surgical status between DES group and TIVA group by multiple variable analyses\*

β = difference between TIVA and DES (DES is reference); SE = standard error of β.
\* Model 1: adjusted sex, age, height, and weight, Model 2: adjusted sex, age, height, veight, weight, and each presented variable; Model 4: adjusted sex, age, height, nevels of fusion, total surgical suite time, and fentanyl; Model 3: adjusted sex, age, height, and each presented variable; Model 4: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 3: adjusted sex, age, height, nevels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical suite time, and fentanyl; Model 5: adjusted sex, age, height, levels of fusion, total surgical surgica

		Model 1			Model 2			Model 3			Model 4			Model 5	
Variable	β	SE	p Value	β	SE	p Value	β	SE	p Value	β	SE	p Value	β	SE	p Value
Max HR during extubation, bpm	-10.92 1.57	1.57	<0.001	-11.58	1.45	<0.001	-11.76	1.45	<0.001	-11.30	1.69	<0.001	-11.00	1.55	<0.001
Max MBP during extubation, mm Hg	-7.87	1.31	<0.001	-8.23	1.26	<0.001	-8.51	1.26	<0.001	-8.75	1.47	<0.001	-8.70	1.35	<0.001
ΔHR, bpm	-12.23	1.57	<0.001												
AMBP, mm Hg	-8.81 1.39	1.39	<0.001												

β = difference between TIVA and DES (DES is reference); SE = standard error of β. \* Model 1: adjusted sex, age, height, and weight, model 2: adjusted sex, age, height, weight, each baseline value, total surgical suite time, and fentanyl; Model 4: adjusted sex, age, height, weight, each baseline value, anesthesia time, extubation time, PACU LOS, and fentanyl; and Model 5: adjusted sex, age, height, weight, each baseline value, anesthesia time, extubation time, PACU LOS, fentanyl, and levels of fusion. p values < 0.05 were considered significant. in anesthesia drugs and surgical procedure (short-duration surgical procedures were predominately investigated in the previous meta-analysis).

The hemodynamics investigated in this study were more stable during extubation. During extubation, heart rate and MBP increased less in the TIVA group than in the DES group. This may be explained because of the use of propofol and the larger dose of fentanyl administered during operations, which reduced stimulation during extubation. Hohlrieder et al.11 found that TIVA is associated with significantly less coughing and reduces hemodynamic response when compared with inhalation anesthesia in elective lumbar disc surgery during emergence from anesthesia. In addition, Wang et al.<sup>21</sup> demonstrated more stable hemodynamics in the TCI patient group and significantly lower mean arterial pressure and faster recovery profiles compared with the fentanyl/DES patient group. Variable doses of intraoperative fentanyl administrated between groups may explain these differences in findings.

A retrospective study design may lead to bias regarding standardization and comparability of study groups. For the purpose of this study, retrospective analysis of data provided a major advantage since anesthesia management was performed by the attending anesthesiologist, according to clinical demands, as opposed to being determined by a study protocol. This study, performed under real clinical conditions, more accurately reflects the clinically relevant benefits that may be expected with the use of new drugs or devices. Finally, comorbidity may play a role in extubation time; however, this could not be investigated in this study owing to lack of patient comorbidity records in the hospital databases.

## Conclusions

Relative to DES, propofol-based TIVA by TCI reduced the mean time to extubation and total surgical time by 5.4 and 11.8 minutes, respectively, and produced more stable hemodynamics during extubation in lengthy lumbar spine surgery. The reduction in extubation and total surgical suite time will have an economic impact through the associated increases in operating room productivity and reduction in labor costs, especially when operating rooms are consistently used for more than 8 hours daily.

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### Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper.

### **Author Contributions**

Conception and design: YH Huang, Lu, Wu. Acquisition of data: YH Huang. Analysis and interpretation of data: Lee, C Lin. Drafting the article: YH Huang, Lu. Critically revising the article: Lu, Wu, BF Lin. Reviewed submitted version of manuscript: YH Huang, Lu, Wu, BF Lin, YS Huang. Approved the final version of the manuscript on behalf of all authors: YH Huang. Statistical analysis: Lee, C Lin. Study supervision: Wu.

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