Effects of remifentanil-propofol combined with dexmedetomidine on cognitive dysfunction in elder patients after ureteroscopic holmium laser lithotripsy: A double-blind randomized controlled trial

Fangjun Wang (wfjlxy006@nsmc.edu.cn)
North Sichuan Medical University

Dan Xie
Affiliated Hospital of North Sichuan Medical College

Hongchun Xu
Affiliated Hospital of North Sichuan Medical College

Qin Ye
Affiliated Hospital of North Sichuan Medical College

Le Wu
Affiliated Hospital of North Sichuan Medical College

Xiaopei Gao
Affiliated Hospital of North Sichuan Medical College

Research

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Abstract

BACKGROUND

Clinical study indicated that infusion of dexmedetomidine without loading dose intraoperative provided smooth and haemodynamically stable emergence, and improved quality of recovery with fewer postoperative side effects and analgesic requirements 9 . The objective is to determine remifentanil-propofol combined with dexmedetomidine during general anesthesia would decrease the incidence and severity of postoperative emergence agitation, anxiety and depression, and without effect on the cognitive dysfunction in elder patients.

METHODS

120 elder patients scheduled for ureteroscopic holmium laser lithotripsy were randomly allocated to PR group administered the normal saline, and PRD group administered dexmedetomidine 0.4 µg.Kg⁻¹.h⁻¹ intravenously after induction of anesthesia, and stopped 30 minutes before the end of surgery. The primary outcome was the scores of richmond agitation sedation, mini mental state examination, state-trait anxiety inventory, zung self-Rating depression scale, and the arabic numeral memory. The secondary outcome was the duration of surgery, and time to spontaneous respiration, recovery and extubation.

RESULTS

The dosage of propofol and remifentanil decreased more significantly in PRD group (P< 0.001). The RASS scores in the PRD group was significantly lower than in PR group at t 1-3 (P< 0.001). The MMSE scores were lower at T 1 - 2 in two groups (P< 0.001). Compared to PR group, the ZSDS scores and STAI scores at T 1 - 2 were lower in PRD group (P<0.005). The recalled arabic numbers were lower at T 2 in PR group (P< 0.001).

CONCLUSION

Dexmedetomidine administration could reduce both the dosage of remifentanil and propofol during surgery, and the incidence and severity of postoperative emergence agitation, anxiety and depression in elderly patients.

Trial registration

the Chinese Clinical Trial Registry, ChiCTR1900021254, Registered 3 February 2019, http://www.chictr.org.cn/ChiCTR1900021254.

Introduction

Postoperative cognitive dysfunction (POCD) is a common complication after surgery that adversely affects the social independence, quality of life, and mortality of patients1. Approximately 12% of
apparently previously cognitively well patients undergoing anaesthesia and noncardiac surgery will develop symptoms of cognitive dysfunction after their procedure\(^2\). In particular, the incidence of POCD is much higher for elderly Surgical patients\(^3\). The Risk factors including preoperative impairment in neurocognitive function, old age, metabolic disturbances, duration/type of surgery, hypoxemia, use of certain anesthetics, and pain are implicated in contributing to POCD\(^4\). As yet no effective treatment for POCD, the prevention of occurrence or reduction the incidence of POCD is more important.

The incidence of POCD was significantly higher in elderly patients undergoing laparoscopic cholecystectomy anaesthetized with sevoflurane or isoflurane compared to propofol\(^5\). Ekmekci P et al reported that Propofol-remifentanil is better than meperidine-midazolam concerning cognitive function in sedation for colonoscopy\(^6\). Propofol-remifentanil allows earlier cognitive recovery compared to Propofol-dexmedetomidine\(^7\). These results showed that maybe Propofol-remifentanil is a good choice of anesthesia for elderly Surgical patients.

However, both propofol and remifentanil are rapid onset, short duration, and rapid revival, it leads to early postoperative catheter-related bladder discomfort (CRBD) and earlier demand for postoperative analgesics\(^7,8\). The clinical study indicated that infusion of dexmedetomidine without loading dose intraoperative provided smooth and haemodynamically stable emergence, and improved quality of recovery with fewer postoperative side effects and analgesic requirements after nasal surgery\(^9\). Thus, we postulated that intraoperative infusion of dexmedetomidine without loading dose would decrease the incidence and severity of early postoperative CRBD, and have little effect on the cognitive dysfunction in elderly patients undergoing ureteroscopic holmium laser lithotripsy anaesthetized with Propofol-remifentanil. Therefore, we designed this study to test the hypothesis that remifentanil-propofol combined with or without dexmedetomidine would have the same effects on postoperative cognitive function in elderly patients after ureteroscopic holmium laser lithotripsy.

**Methods**

Following the approval by the Ethics Committee of the affiliated hospital of north sichuan medical college, we obtained the written informed consent from all the participants for this randomized prospective clinical trial conducted at the affiliated hospital of north sichuan medical college, on patients with the upper urinary tract calculi. This prospective, double-blind, randomized controlled study was registered at the Chinese Clinical Trial Registry (http://www.chictr.org.cn/; Registration number ChiCTR1900021254).

One hundred and twenty adult ASA I-II patients between 60 and 75 years of age undergoing ureteroscopic holmium laser lithotripsy were enrolled in the study between February 2019 and September 2019. Patients scheduled for elective ureteroscopic holmium laser lithotripsy under general anesthesia, and fasted for 12h for solid food, and 6 h for clear liquids before the study were included in the study. The exclusion criteria were as follows: adverses response to propofol, remifentanil or dexmedetomidine, presence of cardiovascular disease, endocrine disease, liver or kidney dysfunctions, recent smoking,
history of chronic use of alcohol, sedatives, opioid, cognitive dysfunction or paracenesthesia and change of surgical.

Patients were divided randomly into two groups, using sealed envelopes indicating the allocation, to receive intravenous dexmedetomidine 0.4 µg.Kg\(^{-1}\).h\(^{-1}\) (PRD group, \(n=60\)) or equal volume of normal saline (PR group, \(n=60\)) after general anesthesia induction. Randomization was performed by an anesthesiologist who was not responsible for Surgical anesthesia of the patients or data collection. The study drugs were administered by an anesthetic nurse while the anesthesiologist responsible for the patient did not know what they were.

Preoperative visits and communications with patients and their relatives were conducted the day before surgery. The patients were familiar with the questionnaires which included mini-mental state examination (MMSE), State-Trait Anxiety Inventory (STAI), Zung Self-Rating Depression Scale (ZSDS), and remembered five random arabic numbers.

Patients enrolled in the study were premedicated with an intramuscular injection of atropine (0.5 mg), 30 minutes before the induction of anesthesia. When the patients arrived in the operating room (\(T_0\)), recalled the arabic numbers (RAM) which remembered the day before surgery and Counted the correct number, then the Mini Mental State Examination (MMSE), State-Trait Anxiety Inventory (STAI), and Zung Self-Rating Depression Scale (ZSDS) were applied. Routine monitoring included electrocardiography, noninvasive blood pressure (systolic blood pressure, mean arterial pressure and diastolic blood pressure), heart rate, respiratory rate, pulse oximetry, end-tidal CO2, the bispectral index and temperature was started.

Patients were induced with intravenous propofol 2 mg/kg, remifentanil 2µg/kg, Cisatracurium 0.15mg/kg. Following the endotracheal tube was inserted, controlled mechanical ventilation were adjusted to maintain an end-tidal carbon dioxide concentration of 40 to 45 mmHg. Immediately after induction of anesthesia, the dexmedetomidine 0.4 µg.Kg\(^{-1}\).h\(^{-1}\) was infused intravenously in dexmedetomidine group, while equal volume of normal saline was administrated in control group. Anesthesia was maintained with plasma target concentration of propofol 2 ~ 3 µg/ml, remifentanil 4 ~ 6 ng/ml, and the value of the bispectral index was maintained between 40 and 60 during surgery. Cisatracurium was given intraoperatively if required. The administration of Cisatracurium, dexmedetomidine or placebo, and propofol-remifentanil were stopped 45,30, and 5 minutes before the end of surgery respectively. Bradycardia (heart rate below 50 beats/min) and hypertension (SBP below 90 mmHg) were treated with atropine (0.5mg) and ephedrine (5 mg) intravenously, respectively.

The patients were extubated after spontaneous respiration (tidal volume > 6 ml/kg, respiratory rate > 13/min), a train-of-four (TOF) ratio \(\geq 0.9\), SpO\(_2\) > 90% under air inspiration, and BIS > 80. The duration of surgery, and time to spontaneous respiration, recovery and extubation (time from stopping administration of propofol-remifentanil to spontaneous respiration, recovery and extubation respectively) were recorded. Patients were transferred to the post-anesthesia care unit (PACU) after extubation. \(O_2\) was applied at 5L
min⁻¹ via a nasal catheter. The PACU emergence agitation score was evaluated 10 minutes (t₁), 20 minutes (t₂), 30 minutes (t₃) and 60 minutes (t₄) after extubation by an anesthetic nurse blinded to the study using the Ramsay sedation scale (RASS: +4, combative; +3, very agitated; +2, agitated; +1, restless; 0, alert and calm; -1, drowsy; -2, light sedation; -3, moderate sedation; -4, deep sedation; -5, unarousable)¹⁰. Emergence agitation (EA) was defined as any RASS score ≥ +2, with severe EA defined as RASS ≥ +3. When the modified Aldrete score >9, the patients were transferred to the surgical ward¹¹. The duration of stay in the PACU was recorded.

The MMSE, STAI, ZSDS, and RAM were applied at 3h (T₁), 6h (T₂), 24h (T₃), 48h (T₄) and 72h (T₅) postoperatively.

**Statistical analysis**

A previous study¹² showed that the mean±SD value of MMSE scores evaluated at 6h postoperatively in patients anesthetized with intravenous propofol-remifentanil was 24.3±2.3, and POCD was considered according to the criteria of MMSE score reductions of ≥1 ± standard deviation. We calculated that a sample size of 62 patients was required in each group at a power of 90%, with a two-sided significance level of 0.05 by an independent t-test. To account for a 10% dropout rate, we included 68 patients in each group. We thus planned to enroll 136 subjects in this study.

Statistical analyses were performed using SPSS22.0 program. Results are expressed as the mean ± standard deviation (SD). One-way analysis of variance (ANOVA) was used to compare mean differences between groups for demographic data (age and weight), time of operation, time to spontaneous respiration, time to recovery and time to extubation, and dosage of propofol and remifentanil administered. Two-way ANOVA, followed by post hoc tests, was used to analyze the scores of MMSE, STAI, ZSDS, and RAM. The sex ratio, ASA physical status, levels of education and scores of RASS were analysed using the X² or Fisher exact tests. P < 0.05 was considered statistically significant.

**Results**

One hundred and thirty-four patients were screened for eligibility, four patients with hypertension were excluded, three patients declined to participate and two patients were Cancelled the surgical procedure. 125 patients were subsequently allocated to two groups. Surgical plan changed in 5 patients during operation. A total of one hundred and twenty patients completed the study and were analysed (Fig. 1). There was no differences between groups regarding demographics (Table 1). The duration of surgery, time to spontaneous respiration and the length of stay in the PACU were similar between groups. The dosage of propofol or remifentanil was significantly lower in PRD group compared with PR group (p < 0.001). The time to recovery and tracheal extubation time were delayed greater in PRD group (p < 0.001) (Table 2).
The Richmond Agitation Sedation scores are shown in table 3. The RASS scores in the PRD group was significantly lower than in PR group at $t_1$, $t_2$, $t_3$ ($P<0.001$). The incidence of EA in the PRD group was significantly lower than in PR group at $t_1$ (10.0% [6/54] vs 40.0% [24/36]), $t_2$ (3.3% [2/58] vs 40.0% [24/36]), $t_3$ (0.0% [0/60] vs 23.3% [14/46]). The Mini Mental State Examination scores are shown in table 4. The MMSE scores were lower at $T_1$ and $T_2$ compared to $T_0$ in two groups. But the MMSE scores were similar between the two groups at $T_0$, $T_1$, $T_2$, $T_3$, $T_4$, and $T_5$. The Zung Self-Rating Depression Scale scores are shown in table 5. The ZSDS scores in PR group were higher compared to PRD group at $T_1$ and $T_2$ ($P<0.001$), and the ZSDS scores were similar between the two groups at $T_0$, $T_3$, $T_4$, and $T_5$. The ZSDS scores were higher at $T_1$ and $T_2$ compared to $T_0$ in PR group ($P<0.001$), and there were no differences in ZSDS scores at $T_0$, $T_1$, $T_2$, $T_3$, $T_4$, and $T_5$ in PRD groups ($P=0.359$). The State-Trait Anxiety Inventory scores are shown in table 6. The STAI scores were higher at $T_1$ and $T_2$ compared to $T_0$ in PR group, and the STAI scores were higher at $T_1$ compared to $T_0$ in PRD group. The STAI scores at $T_1$ and $T_2$ were lower in PRD group compared to PR group ($P<0.005$). The recalled arabic numbers are shown in table 7. The recalled arabic numbers were lower at $T_1$ and $T_2$ in PR group and at $T_1$ in PRD group compared to $T_0$ ($P<0.001$). The recalled arabic numbers were lower at $T_2$ in PR group compared to PRD group ($P<0.001$).

**Discussion**

The results of this clinical trial showed that a combination of intravenous dexmedetomidine 0.4 $\mu$g.Kg$^{-1}$.h$^{-1}$ resulted in lower incidence and severity of postoperative emergence agitation in elder patients undergoing ureteroscopic holmium laser lithotripsy. Intravenous dexmedetomidine reduced the use of remifentanil and propofol during operation. Although there was no difference between groups in influence on postoperative cognitive dysfunction, the patients anaesthetized with remifentanil-propofol combined with or without dexmedetomidine had transient postoperative cognitive dysfunction. The duration of anxious and depression after operation were longer in the patients treated without intravenous dexmedetomidine.

Postoperative cognitive dysfunction (POCD) is one of the most common postoperative complications in elderly patients and is associated with an increased risk of death in the first year after surgery$^{13}$. Advancing age, multiple surgeries, duration of anesthesia and acute postoperative pain have been implicated as the risk factors for POCD$^{14,15}$. The older surgical patients have probable cognitive impairment preoperatively, and the impairment is associated with development of delirium postoperatively$^{16}$. We previously found that the incidence of postoperative cognitive dysfunction was higher in elderly patients with inhalational anaesthesia compared to total intravenous anaesthesia$^{17}$. Both anesthetic and pain management strategies do appear to influence the risk of cognitive dysfunction after elective joint arthroplasty, and perioperative pain may be a risk factor for postoperative delirium$^{18-20}$. In the present study, we selected patients undergoing ureteroscopic holmium laser lithotripsy with minor acute postoperative pain, thus avoiding the effects of postoperative pain and postoperative analgesic drugs on POCD in elderly patients. Our study showed that the MMSE scores were decreased
significantly 3th hour after surgery, and restored to pre-anesthesia level 24th hour after surgery in two
groups. This Suggests that postoperative cognitive dysfunction was temporary during TIVA with
remifentanil and propofol given by TCI with or without dexmedetomidine in elderly patients undergoing
ureteroscopic holmium laser lithotripsy. In contrast, İlvan G et al. 19 reported that the TIVA method did not
affect postoperative early cognitive functions in either old or young patients who underwent lumbar disk
surgery. The time to apply the MMSE postoperatively might be responsible for the differences. Because
they applied the MMSE 24h after surgery and found no one experienced POCD, and the MMSE scores
were decreased significantly 3th hour and restored 24th hour after surgery in our study. Therefore, their
results could not demonstrate that no transient cognitive dysfunction was occurred within 24 hours
postoperatively. Our results were consistent with previous studies that propofol might cause POCD21,22
and different with the results reported by Zhang D et al.23 that propofol might be able to inhibit the
inflammatory response in central nervous system and improve POCD. This indicated that prolonged
propofol-based general anesthesia may have an effect on the central nerve system, and short-term
infusion propofol during TIVA would yield a minimal influence on postoperative cognitive function.

Dexmedetomidine is a new α2 adrenergic receptor agonist with a short half-life (about two hours), and it
has a dose-dependent sedative and analgesic effects and no negative effects on respiration24. A meta-
analysis done by Tan JA et al. found that dexmedetomidine could induce sedation without increasing
delirium 25. These findings are compatible with the results of our study that dexmedetomidine has no
effects on postoperative cognitive function in elderly patients anaesthetized with remifentanil and
propofol. Several studies have demonstrated that dexmedetomidine may benefit cognitive function in
elderly patients due to its neuroprotective effect and inhibiting inflammation24, 26, 27. In our study, we
didn’t detect the Plasma inflammatory factors, and the effect of dexmedetomidine on plasma
inflammatory mediators in elderly patients was unclear. Tan JA et al.25 reported that the risk of
bradycardia was significantly higher when both a loading dose and high maintenance doses (>0.7
microg/kg/h) were used. In the present study, no patients required intervention for bradycardia or
hypotension. This may be due to the intraoperative infusion of dexmedetomidine without loading dose
and lower maintenance doses (0.4 microg/kg/h < 0.7 microg/kg/h).

Emergence agitation develops in the early phase of general anesthesia recovery and is observed more
frequently in ENT (ear, nose, and throat) surgical patients 28. EA is characterized by agitation, confusion,
disorientation, and possible violent behavior 28, 29. EA can cause hemorrhage, falling out of the bed, self-
tubation, removal of catheters, even injury to the patient or medical staff 30. The present study
describes a higher incidence of EA in PR group compared to PRD group. This indicated that intravenous
dexmedetomidine after induction of anaesthesia significantly reduced the incidence and severity of EA
postoperatively in elderly patients undergoing TIVA with remifentanil and propofol for ureteroscopic
holmium laser lithotripsy. The independent risk factors for EA such as younger age, recent smoking,
sevoflurane anesthesia, postoperative pain of NRS≥ 5, presence of a tracheal tube, and presence of a
urinary catheter were identified 28. In our study, postoperative catheter-related bladder discomfort(CRBD)
was more complained by elderly patients in PR group. This suggested that the intraoperative intravenous dexmedetomidine could decrease the incidence and severity of early postoperative CRBD. Our results were consistent with previous studies that intraoperative administration of dexmedetomidine is a safe and effective practice for the prevention of CRBD after lumbar microdiscectomy and can reduce postoperative pain. The lower incidence of EA in PRD group in present study may be attributed to the decreased incidence and severity of early postoperative CRBD.

In our study, both the ZSDS scores and the STAI scores were increased at 3th hour and restored 12th hour after operation in PR group. However, only the STAI scores were increased at 3th hour and restored 6th hour postoperatively in PRD group. This indicated that intraoperative infusion of dexmedetomidine could attenuate the severity of postoperative anxiety and depression in elderly patients. Ingrid Rundshagen reported that an anxious, depressed basal mood have been identified as further risk factors for POCD. Then the administrated of dexmedetomidine may benefit to the postoperative cognitive function by decreasing incidence and severity of postoperative anxiety and depression.

There are limitations to our study. We didn't detect the Plasma inflammatory factors in our study. Many studies have reported that the protective effect of dexmedetomidine on cognitive function due to its inhibiting inflammation. The patients selected in our study had almost no postoperative pain, so the effects of remifentanil-propofol combined with or without dexmedetomidine on cognitive dysfunction in elderly patients with postoperative pain were unclear.

In conclusion, the present study showed that intravenous dexmedetomidine could provide clinically relevant benefits in elderly patients undergoing ureteroscopic holmium laser lithotripsy surgery. Dexmedetomidine administration could reduce the dosage of remifentanil and propofol during surgery. Although dexmedetomidine had no effect on postoperative cognitive dysfunction, it could reduce the incidence and severity of postoperative emergence agitation, anxiety and depression in elderly patients.

**Declarations**

Ethics approval and consent to participate: All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Following the approval by the Ethics Committee of the affiliated hospital of north sichuan medical college, we obtained the written informed consent from all the participants for this randomized prospective clinical trial conducted at the affiliated hospital of north sichuan medical college, on patients with the upper urinary tract calculi.

Consent for publication: There was no details, images, or videos relating to an individual person in this manuscript.

Trial registration: The Chinese Clinical Trial Registry, ChiCTR1900021254, Registered 3 February 2019, http://www.chictr.org.cn/ChiCTR1900021254.
Availability of data and materials: we presented the study datasets in additional supporting files (PRD-PR study data. XLSX).

Assistance with the study: none.

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Conflicts of interest: none.

IRB Information: This study was approved by the IRB at the Affiliated Hospital of North Sichuan Medical College (IRB 2019ER(R)025), Contact: IRB specialist; Jiemei Huang, Phone: 86-0817-2262124 and written informed consent had been obtained from all subjects participating in the trial. This study was registered prior to patient enrollment at the Chinese Clinical Trial Registry (Principal investigator: Fangjun Wang, Date of registration: February 3 2016), Registry URL: http://www.chictr.org.cn/; Registration number: ChiCTR1900021254.

Presentation: none.

Authors’ contributions

FJ W obtained funding for the research. All authors have contributed to the design of the study. HC X, Q Y, D X and L W were involved in the study design, the experiments performance, the data collection. XP G wrote the manuscript and reviewed and revised by FJ W and DX. All authors have edited, reviewed and approved the final version.

Author details

the Department of Anaesthesiology, Affiliated Hospital, North Sichuan Medical College, Nanchong, China

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Tables

The Patient characteristics are shown in Table 1. Patient characteristics were similar between the two groups, (Table 1).

**Table 1** Demographic data.

| Patient characteristics          | PR group | PRD group | F/X² values | P values |
|---------------------------------|----------|-----------|-------------|----------|
| Sex                              | n=60     | n=60      | F/X² values | P values |
| male /female                    | 35/25    | 33/27     | 4           | 0.2615   |
| level of education               |          |           |             |          |
| illiterate / Primary school literacy | 27/33    | 29/31     | 2           | 0.1573   |
| Age(y)                           |          |           |             |          |
|                                 | 66.7±4.1 | 65.6±3.4  | 3.287       | 0.125    |
| Weight(kg)                       |          |           |             |          |
|                                 | 56.5±5.4 | 55.8±5.9  | 0.465       | 0.496    |
| ASA(I/II)                        |          |           |             |          |
|                                 | 20/40    | 18/42     | 2           | 0.1573   |

Values are mean±SD, number of patients. ASA: American Society of Anesthesiologists; PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine.

The clinical characteristics are shown in Table 2. The duration of surgery, time to spontaneous respiration and the length of stay in the PACU were similar in two groups. The dosage of propofol or remifentanil was significantly higher in PRD group compared with PR group (P<0.001). The time to recovery and tracheal extubation time were delayed greater in group PRD compared with group PR (P<0.001).

**Table 2** Clinical characteristics in two groups
Clinical characteristics | PR group n=60 | PRD group n=60 | F values | P values
--- | --- | --- | --- | ---
duration of surgery (minute) | 93.4±10.9 | 94.1±10.9 | 0.124 | 0.725
dosage of propofol (milligram) | 517.0±75.8 | 335.8±40.8* | 265.928 | 0.000
dosage of remifentanil (microgram) | 486.0±53.2 | 439.4±32.7* | 33.407 | 0.000
time to spontaneous respiration (minute) | 16.4±2.2 | 17.1±2.8 | 2.375 | 0.126
time to recovery (minute) | 20.0±2.4 | 24.5±3.6* | 64.195 | 0.000
tracheal extubation time (minute) | 21.2±2.7 | 25.4±3.6* | 53.371 | 0.000
PACU stay time (minute) | 66.2±5.2 | 67.9±6.5 | 2.544 | 0.113

Values are mean±SD. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine; PACU: postanesthesia care unit.*p < 0.001 vs. PR group.

The Richmond Agitation Sedation scores are shown in Table 3. The RASS scores in the PRD group was significantly lower than in PR group at t1, t2, t3 (P < 0.001). The incidence of EA in the PRD group was significantly lower than in PR group at t1 (10.0% [6/54] vs 40.0% [24/36]), t2 (3.3% [2/58] vs 40.0% [24/36]), t3 (0.0% [0/60] vs 23.3% [14/46]), (Table 3).

**Table 3** The Richmond Agitation Sedation scores at 10, 20, 30 and 60 minutes after extubation in two groups.

| Time | Group | 0 | 1 | 2 | 3 | \(X^2\) | \(P\) values |
|------|-------|---|---|---|---|-------|----------|
| t1   | PR    | 0 | 10| 26| 14| 10 | 42.329 | 0.000*   |
|      | PRD   | 12| 25| 17| 6 | 0   |
| t2   | PR    | 0 | 17| 19| 16| 8  | 54.262 | 0.000*   |
|      | PRD   | 21| 23| 14| 2 | 0   |
| t3   | PR    | 5 | 13| 28| 14| 0  | 48.415 | 0.000*   |
|      | PRD   | 14| 38| 8 | 0 | 0   |
| t4   | PR    | 7 | 36| 17| 0 | 0  | 2.930  | 0.231    |
|      | PRD   | 7 | 42| 9 | 0 | 0   |

Number of patients. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine. t1:10 minutes after extubation, t2:20 minutes after extubation, t3:30 minutes after extubation, t4:60 minutes after extubation. *P<0.001 vs. group PR.

The Mini Mental State Examination scores are shown in Table 4. The MMSE scores were lower at T1 and T2 compared to T0 in two groups (P<0.001). The MMSE scores were similar between the two groups at T0, T1, T2, T3, T4 and T5 (P>0.05).

**Table 4** The Mini Mental State Examination scores at T0, T1, T2, T3, T4 and T5 in two groups.

|      | PR    | PRD   | F values | P values |
|------|-------|-------|----------|----------|
| T0   | 25.4±2.2 | 25.5±2.5 | 0.431 | 0.700 |
| T1   | 20.7±2.1* | 21.3±2.5* | 0.423 | 0.183 |
| T2   | 22.8±2.0* | 23.3±2.0* | 0.367 | 0.223 |
| T3   | 25.0±2.2 | 25.2±2.6 | 0.432 | 0.644 |
| T4   | 25.1±2.2 | 25.2±2.4 | 0.421 | 0.664 |
| T5   | 25.3±2.2 | 25.4±2.5 | 0.429 | 0.756 |

\(F\) values: 129.34, 105.55, —, —, —, —. \(P\) values: 0.000, 0.000, —, —, —, —.

Values are mean±SD. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine; T0: before the induction of anesthesia, T1: 3h after surgery, T2: 6h after surgery, T3: 24h after surgery, T4: 48h after surgery, T5: 72h after surgery. *p < 0.001 vs. T0.
The Zung Self-Rating Depression Scale scores are shown in table 5. The ZSDS scores in PRD group were lower compared to PR group at T₁ and T₂ (P<0.001), and the ZSDS scores were similar between the two groups at T₀, T₃, T₄ and T₅. The ZSDS scores were higher at T₁ and T₂ compared to T₀ in PR group (P<0.001), and there were no differences in ZSDS scores at T₀, T₁, T₂, T₃ and T₅ in PRD groups (P=0.359).

Table 5 The Zung Self-Rating Depression Scale scores at T₀, T₁, T₂, T₃, T₄ and T₅ in two groups.

| Group | n  | T₀         | T₁         | T₂         | T₃         | T₄         | T₅         | F values | P values |
|-------|----|------------|------------|------------|------------|------------|------------|----------|----------|
| PR    | 60 | 43.2±2.2   | 48.1±2.5*  | 46.2±2.4*  | 43.6±1.6  | 43.2±1.7  | 43.3±1.9  | 69.06    | 0.000    |
| PRD   | 60 | 42.8±2.4   | 43.4±2.2*  | 43.2±1.7*  | 42.9±2.0  | 42.7±2.2  | 42.8±1.70 | 1.35     | 0.248    |

F values 0.419 0.434 0.379 0.329 0.357 0.333 — —
P values 0.362 0.000 0.000 0.028 0.138 0.136 — —

Values are mean±SD. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine. T₀: before the induction of anesthesia, T₁: 3h after surgery, T₂: 6h after surgery, T₃: 24h after surgery, T₄: 48h after surgery, T₅: 72h after surgery. *P< 0.001 vs. T₀, #P< 0.005 vs. PR group.

The State-Trait Anxiety Inventory scores are shown in table 6. The STAI scores were higher at T₁ and T₂ compared to T₀ in PR group, and the STAI scores were higher at T₁ compared to T₀ in PRD group. The STAI scores at T₁ and T₂ were lower in PRD group compared to PR group (P<0.005).

Table 6 The State-Trait Anxiety Inventory scores at T₀, T₁, T₂, T₃, T₄ and T₅ in two groups.

| Index | Group | n  | T₀         | T₁         | T₂         | T₃         | T₄         | T₅         | F values | P values |
|-------|-------|----|------------|------------|------------|------------|------------|------------|----------|----------|
| S-AI  | PR    | 60 | 38.9±3.4   | 48.6±3.6*  | 45.3±3.7*  | 39.0±3.0  | 39.1±3.1  | 39.1±3.4  | 171.05   | 0.000    |
| PRD   | 60    | 39.5±2.6 | 45.5±2.5*  | 39.9±2.3*  | 39.9±2.3  | 39.5±2.3  | 39.6±2.0  | 36.30     | 0.000    |

F values 0.554 0.561 0.565 0.490 0.504 0.507 — —
P values 0.294 0.000 0.000 0.079 0.509 0.266 — —

| T-AI  | PR    | 60 | 40.4±2.9   | 46.9±3.1*  | 45.9±2.9*  | 40.7±2.5  | 40.5±2.5  | 40.7±2.5  | 74.79    | 0.000    |
| PRD   | 60    | 41.1±2.2 | 45.1±3.1*  | 41.4±1.9*  | 40.8±2.8  | 40.6±2.2  | 40.7±2.4  | 45.99     | 0.000    |

F values 0.468 0.565 0.449 0.490 0.436 0.447 — —
P values 0.179 0.002 0.000 0.812 0.789 1.000 — —

Values are mean±SD. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine. S-AI: State Anxiety Inventory; T-AI: State-Trait Anxiety Inventory. T₀: before the induction of anesthesia, T₁: 3h after surgery, T₂: 6h after surgery, T₃: 24h after surgery, T₄: 48h after surgery, T₅: 72h after surgery. *P< 0.001 vs. T₀, #P< 0.005 vs. PR group.

The recalled arabic numbers are shown in table 7. The recalled arabic numbers were lower at T₁ and T₂ in PR group and at T₁ in PRD group compared to T₀ (P<0.001). The recalled arabic numbers were higher at T₂ in PRD group compared to PR group (P<0.001).

Table 7 The recalled arabic numbers at T₀, T₁, T₂, T₃, T₄, and T₅ in two groups.
| Group  | n  | T₀    | T₁    | T₂    | T₃    | T₄    | T₅    | F values | P values |
|--------|----|-------|-------|-------|-------|-------|-------|----------|----------|
| PR     | 60 | 4.4±0.6 | 2.7±0.7 | 3.1±0.7 | 4.2±0.7 | 4.3±0.6 | 4.3±0.7 | 61.79    | 0.000    |
| PRD    | 60 | 4.3±0.7 | 2.6±0.8* | 4.1±0.7# | 4.3±0.7 | 4.2±0.6 | 4.3±0.7 | 55.09    | 0.000    |
| F values |    | 0.118 | 0.134 | 0.126 | 0.127 | 0.114 | 0.122 | —        | —        |
| P values |    | 0.573 | 0.323 | 0.000 | 0.694 | 0.465 | 1.000 | —        | —        |

Values are mean±SD. PR: Propofol-remifentanil; PRD: Propofol-remifentanil and dexmedetomidine. S-AI: State Anxiety Inventory; T-AI: State-Trait Anxiety Inventory. T₀: before the induction of anesthesia, T₁: 3h after surgery, T₂: 6h after surgery, T₃: 24h after surgery, T₄: 48h after surgery. T₅: 72h after surgery.*p < 0.001vs.T₀. #p < 0.001vs.PR group.

**Figures**
Figure 1

Study flow diagram. One hundred and thirty-four patients were screened for eligibility, six patients with hypertension were excluded, five patients declined to participate and two patients were cancelled the surgical procedure. 125 patients were subsequently allocated to two groups. During operation, five patients were change of surgical. A total of one hundred and twenty patients completed the study.

Supplementary Files

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