Atypical Sleep and Postoperative Delirium in the Cardiothoracic Surgical Intensive Care Unit: A Pilot Prospective Study

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Purpose: Postoperative delirium (POD) is a very common and serious neurological complication in patients admitted to the cardiothoracic surgical intensive care unit (CSICU). We aimed to identify a novel potential sleep-based marker for POD and investigate the relevance between atypical sleep and POD.

Patients and Methods: This was a prospective, observational study of patients admitted to the CSICU between December 2019 and February 2020 at our center. Sleep characteristics from 21:00 on postoperative day 1 to 07:00 on postoperative day 2 were assessed using polysomnography (PSG). POD from the end of PSG monitoring until postoperative day 5 was evaluated using the Confusion Assessment Method for the Intensive Care Unit.

Results: This analysis included 20 patients admitted to the CSICU. The incidence of atypical sleep was 45.0%. Compared to patients without delirium, those with delirium had less delta power, less percentage REM sleep, and a higher proportion of atypical sleep and REM sleep loss ($P < 0.05$).

Conclusion: The presence of atypical sleep and the absence of REM sleep were associated with POD in patients admitted to the CSICU.

Keywords: postoperative delirium, intensive care unit, atypical sleep, cardiac surgery

Introduction

Postoperative delirium (POD), is a common acute disturbance of consciousness occurring postoperatively among 11.5–39.0% of the patients admitted to the cardiothoracic surgical intensive care unit (CSICU), it is associated with prolonged hospital stay, poor outcomes, and increased health costs and mortality.^{1–5} Hence, identifying risk factors for POD is important. Current research shows that POD after cardiac surgery results from a combination of several factors, including advanced age, pre-existing cognitive impairment, previous psychiatric conditions, cerebrovascular disease, mechanical ventilation time, atrial fibrillation, and so forth.^{6,7} However, sleep disturbances, common among CSICU patients, could be a potential risk factor for POD.⁸

Most recent studies have only considered the association between delirium and normal sleep characteristics but not abnormal sleep characteristics. Normal sleep consists of 3 stages: WAKE, non-rapid eye movement (NREM) sleep (further divided into stages N1, N2, and N3), and rapid eye movement (REM) sleep. These can be identified using polysomnography (PSG).⁹ However, these typical sleep characteristics are often absent in patients admitted to the intensive care unit.

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Atypical sleep is an abnormal sleep subtype, characterized by loss of N2 stage markers (K-complexes and sleep spindles).¹⁰ The incidence of atypical sleep has been reported to be 28–50%^{11–14} among awake or lightly sedated patients and 60–85%^{10,15} among deeply sedated patients.

Only limited data exist regarding the association between atypical sleep and delirium, and the causality remains unclear. Previous studies indicate that delirium occurs in 44% of patients with atypical sleep as compared to 18% of patients without atypical sleep; however, whether atypical sleep occurs before delirium onset remains unclear.¹¹ According to a study assessing sleep among 14 patients with mechanical ventilation, typical sleep features were only recognized in 1 patient without delirium.¹⁶ Another study reported no significant difference in the incidence of delirium during the PSG period between atypical and typical sleep.¹² Although the cited studies were important, they did not definitively establish the association between atypical sleep and delirium primarily because these could not establish the sequence of delirium and atypical sleep. Hence, studies assessing the association between atypical sleep and delirium through a rigorous experimental design are lacking.

In this pilot study, we conducted PSG monitoring on the first postoperative night after cardiac surgery using cardiopulmonary bypass (CPB) and assessed delirium daily after PSG monitoring until postoperative day 5 to test the hypothesis that atypical sleep occurs before the onset of and is associated with delirium.

Materials and Methods

Design and Setting

We conducted a pilot prospective study between December 2019 and February 2020 at the CSICU of our hospital. The study was approved by the Ethics Committee of our hospital and is registered at <http://www.chictr.org.cn> (ChiCTR1900023094).

We enrolled 23 patients aged ≥ 18 years and undergoing cardiac surgery using CPB. Patients with the following conditions were excluded: (1) history or current diagnosis of mental illness or neurological disorders; (2) alcoholism (defined using the Alcohol Use Disorders Identification Test Consumption score ≥ 4 in men and ≥ 3 in women);¹⁷ (3) sleep disturbance (defined as a STOP-Bang score of ≥ 3);¹⁸ (4) self-reported use of anticonvulsants, antidepressants, and neuroleptics 48 hours before the study; (5)

contraindications for PSG, including infection, recent trauma, and head or neck surgery; (6) PSG monitoring time ≤ 7 h for various reasons (eg, wire loss or equipment failure); (7) delirium onset after the surgery till the end of monitoring; (8) inability to use the Confusion Assessment Method for the Intensive Care Unit (CAM-ICU) to confirm delirium or non-delirium (the Richmond Agitation-Sedation Scale [RASS ≤ -4]).^{19,20}

Sleep Assessment

A portable 52-channel sleep monitor (BMC Polypro, Beijing) was used for sleep monitoring. Electrode placement conformed to the “International 10–20” System.²¹ Electrodes were placed by a trained technician and included the following: 6 electroencephalogram (EEG) channels (2 frontal channels: F3-M2, F4-M1; 2 central channels: C3-M2, C4-M1; and 2 occipital channels: O1-M2, O2-M1), 2 electrooculogram (E1, E2), and 2 electromyograms (EMG⁺-EMG⁻). PSG was started at 21:00 on postoperative day 1 and was continuously performed until 07:00 on the next day (07:00, morning nursing; 21:00, end of visiting). All records, every 30-second epoch, were scored by a trained sleep technologist and reviewed by 2 sleep medicine physicians. The sleep technologist and sleep medicine physicians were blinded to patient group assignment during the study. If PSG observations conformed with the standard sleep stages, it was scored according to the 2007 American Academy of Sleep Medicine Manual for the Scoring of Sleep and Associated Events. If some epochs of PSG could not be scored using the standard criteria but were consistent with the description of atypical sleep,^{13,15} it was scored as atypical sleep.

For each EEG data set, we selected 100 minutes from a period with minimal care intervention (02:00–03:40); the spectral analysis of the delta (0.5–3.9 Hz), theta (4.0–7.9 Hz), alpha (8.0–13.0 Hz), and beta (13.1–30.0 Hz) bands was performed using fast Fourier transformation of each 4-s epoch. The spectral power estimates were averaged across 6 EEG channels (F3-M2, F4-M1, C3-M2, C4-M1, O1-M2, and O2-M1).

Delirium Assessment

POD was defined as delirium lasting from the end of PSG monitoring till postoperative day 5, ensuring that delirium occurred after sleep monitoring. Patients were screened for delirium from the end of PSG monitoring until postoperative day 5 by trained nurses using RASS and

Table 1 Sleep Characteristics of CSICU Patients

	All Patients (n=20)
Total recording time (min), mean (SD)	506.42±55.11
Total sleep time (min), mean (SD)	235.62±83.24
Sleep efficiency (%), mean (SD)	47.7±18.6
Stage N1 sleep (%), mean (SD)	14.0±7.3
Stage N2 sleep (%), mean (SD)	29.3±20.4
Stage N3 sleep (%), median (IQR)	0.5 (0, 3.3)
REM sleep (%), median (IQR)	0 (0, 2.9)

Abbreviations: REM, rapid eye movement; SD, standard deviation; IQR, interquartile range.

CAM-ICU.^{19,20} To ensure that the sleep data were recorded before delirium and to minimize the effect of the assessment process on patients' sleep at night, patients were screened for delirium at 08:00, 14:00, and 20:00 every day. If patients presented any clinical evidence of delirium (such as agitation or illusion, as observed by the nurses) at other times (including entire sleep monitoring period), they were additionally screened by nurses using RASS and CAM-ICU.

Clinical Data Collection

Patients' sociodemographic characteristics, preoperative status, intraoperative variables, and relevant postoperative data were collected (Table 1). As analgesia and sedation may promote delirium,^{22,23} we also collected analgesia and sedation use information pre- and post-operatively till the end of PSG. Analgesics used were morphine, fentanyl, and dezocine, and sedatives used were dexmedetomidine, midazolam, and chlorpromazine.

Statistical Analysis

Patients were divided into the delirium and non-delirium groups based on whether they developed POD. Continuous variables were presented as mean ± standard deviation (SD; normally distributed data) or medians and interquartile ranges (non-normally distributed data; 25–75% quartiles), and categorical variables were presented as numbers and percentages. The chi-square or Fisher exact test was used to compare qualitative data, as applicable, and the Student *t* or Mann–Whitney *U*-test was used to compare quantitative data, as applicable. SPSS software (version 17.0) was used for data statistical analysis. *P* < 0.05 was considered statistically significant.

Results

Of the 23 patients, 1 was excluded because of delirium onset before the end of PSG monitoring, 2 were excluded because of PSG monitoring lasting ≤7 h (caused by wire disconnection). Of the 20 CSICU patients, 9(45.0%) had atypical sleep (atypical stages 1–3) per the Watson classification.¹⁵ The total atypical sleep duration was 4396 min. A representative EEG of a patient with atypical sleep is shown in Figures 1, and 2 shows the percentage distribution of the sleep stages. Table 1 summarizes the sleep characteristics of all patients.

Figure 3 shows the distribution of POD. No significant differences existed between the groups in baseline characteristics including sociodemographic, preoperative, intraoperative, and postoperative variables (Table 2).

Figure 1 Electroencephalography tracing from an atypical sleep patient. F3, F4, C3, C4, O1, O2 indicate EEG lead locations.

Abbreviations: EMG, electromyogram; LOC and ROC, left and right electrooculogram, respectively.

Figure 2 Sleep stage distribution in CSICU patients.

Abbreviations: NREM, non-rapid eye movement; REM, rapid eye movement.

Patients with delirium had decreased delta power ($P = 0.010$) and percentage REM sleep ($P = 0.011$) and a higher incidence of atypical sleep ($P = 0.005$) and REM sleep absence ($P = 0.028$) (Table 3).

Discussion

To our knowledge, this is the first study assessing the association between atypical sleep and delirium using

Figure 3 Composition ratio of postoperative delirium in different periods of CSICU patients.

PSG through prospectively collected sleep data before delirium onset. Overall, (1) sleep quality was generally poor among all CSICU patients; (2) 45.0% patients (9/20) developed atypical sleep on the first night after cardiac surgery; (3) and patients with delirium had decreased delta power and percentage REM sleep and a higher proportion of atypical sleep and REM sleep absence.

According to the Principles and Practice of Sleep Medicine (Version 6, 2017),²⁴ stage N1, N2, N3, and REM sleep generally constitute about 2%-5%, 45%-55%, 13%-23%, and 20-25% of sleep. However, sleep in CSICU patients is often altered because of the influence of various factors, such as pain, dyspnea, sleep-affecting drugs, ICU environment, and so forth.²⁵ A study by Orr analyzed sleep in 6 patients undergoing open-heart surgery and found that only 1 patient showed electrophysiologic evidence of stage N2 sleep until the 2nd postoperative night, and 5 patients showed a total absence of REM sleep until the 4th postoperative night.²⁶ In another study of 38 patients undergoing coronary artery bypass grafting, the total sleep time reduced to 253.6 ± 94.1 min, with suppressed stage N3 sleep and REM sleep.²⁷ Trends among our data are generally consistent with results of the aforementioned studies, including decreased total sleep time, lower sleep efficiency, and suppressed stage N3 and REM sleep (Table 1).

Severe REM sleep reduction (<6% of total sleep time) is associated with delirium.²⁸ A study by Evans²⁹ including patients undergoing artificial hip replacement showed that reduced total sleep time and reduced non-REM delta power on the 2nd night postoperatively are related to greater POD severity. Similarly, in this study, we found that reduced delta power and absence of REM sleep are associated with POD. Animal studies have shown that REM sleep deprivation cause neurotransmitters imbalance, including increased acetylcholinesterase activity (responsible for acetylcholine degradation) and dopamine levels.^{30,31} Neurotransmitter imbalance (particularly decreased acetylcholine level and increased dopamine level) plays a key role in the development of delirium.³² In contrast, we did not find any statistically significant association between total sleep time and POD. Our inclusion of critically ill patients undergoing cardiac surgery with CPB in this study could explain this difference. Several studies have reported on cardiac surgery affecting sleep characteristics.²⁵ Moreover, sleep in the CSICU is disturbed more often

Table 2 Baseline Characteristics in Non-Delirium Group vs. Delirium Group

	Non-Delirium (n=12)	Delirium (n=8)	t/z/x ²	P
Sociodemographic characteristics				
Age (y), mean (SD)	53.25±15.09	49.38±12.66	0.598	0.557 [†]
Male, n (%)	6 (50.0)	5 (62.5)	-	0.670 [‡]
BMI (kg/m ²), mean (SD)	23.90±4.23	26.52±7.42	-0.855	0.408 [†]
Married, n (%)	11 (91.7)	8 (100.0)	-	> 0.999 [‡]
Education level, n (%)			1.235	0.539 [‡]
Primary school and below	5 (41.7)	2 (25.0)		
Junior and senior high school	5 (41.7)	3 (37.5)		
College and above	2 (16.7)	3 (37.5)		
Smoker, n (%)	6 (50.0)	2 (25.0)	-	0.373 [‡]
Drinker, n (%)	0 (0)	1 (12.5)	-	0.400 [‡]
Preoperative				
Hypertension, n (%)	5 (41.7)	7 (87.5)	-	0.070 [‡]
Diabetes, n (%)	1 (8.3)	0 (0)	-	> 0.999 [‡]
Cerebrovascular disease, n (%)	0 (0)	1 (12.5)	-	0.400 [‡]
Sedation use, n (%)	0 (0)	1 (12.5)	-	0.400 [‡]
Analgesia use, n (%)	8 (66.7)	6 (75.0)	-	> 0.999 [‡]
LVEF (%), mean (SD)	63.3±6.7	63.5±6.0	-0.086	0.932 [†]
Intraoperative				
Operating time (min), mean (SD)	282.58±58.25	312.38±129.94	-0.609	0.558 [†]
CPB time (min), mean (SD)	145.91±43.60	144.43±49.71	0.068	0.946 [†]
Aortic cross-clamp time (min), mean (SD)	71.83±28.78	84.88±29.42	-0.984	0.338 [†]
Postoperative to the end of PSG				
Sedation use, n (%)	3 (25.0)	0 (0)	-	0.242 [‡]
Analgesia use, n (%)	4 (33.3)	3 (37.5)	-	> 0.999 [‡]
APACHE-II score, mean (SD)	28.27±6.65	28.75±2.66	-0.216	0.832 [†]
Length of MV (h), median (IQR)	25.0 (19.0, 148.0)	45.5 (20.3, 98.3)	-0.133	0.894 [*]
Length of ICU stays (d), median (IQR)	5.5 (3.3, 14.8)	5.5 (3.0, 9.8)	-0.428	0.668 [*]

Notes: [†]Student's t-test; ^{*}Mann-Whitney U-test; [‡]Chi-square test; [§]Fisher's Exact Test.

Abbreviations: BMI, body mass index; LVEF, left ventricular ejection fraction; CPB, cardiopulmonary bypass; APACHE, acute physiology and chronic health evaluation; MV, mechanical ventilation; ICU, intensive care unit; SD, standard deviation; IQR, interquartile range.

because of the lights, noise, and a high frequency of medical and nursing activities than in the general ward.

The concept of atypical sleep was introduced by Cooper in 2012, and it is prevalent among ICU patients.¹⁰ A study verifying the feasibility of unattended PSG indicated that 44% of the patients with atypical sleep had delirium compared to 18% of patients with normal sleep; however, this study could not

conclusively establish whether atypical sleep occurs before delirium onset, and the association was not statistically significant.¹¹ The results of our study showed that atypical sleep that occurred before POD is associated with POD. K-complexes and sleep spindle loss are key characteristics of atypical sleep. Higher spindle density has been shown to predict better performance with regard to verbal ability and attention; this indicates

Table 3 Sleep Characteristics in Non-Delirium Group vs. Delirium Group

	Non-Delirium (n=12)	Delirium (n=8)	t/z	P
Total recording time (min), mean (SD)	492.93±59.15	526.65±44.35	-1.371	0.187 [†]
Total sleep time (min), mean (SD)	260.80±83.40	197.85±71.72	1.744	0.098 [†]
Sleep efficiency (%), mean (SD)	54.1±18.9	38.0±14.1	2.048	0.055 [†]
Stage N1 sleep (%), mean (SD)	13.4±6.3	15.1±9.0	-0.508	0.618 [†]
Stage N2 sleep (%), mean (SD)	34.1±23.1	22.2±13.7	1.306	0.208 [†]
Stage N3 sleep (%), median (IQR)	1.3 (0, 3.8)	0 (0, 2.7)	-1.435	0.151 [*]
REM sleep (%), median (IQR)	2.1 (0, 3.2)	0 (0, 0.5)	-2.535	0.011 [*]
Delta (μV^2), median (IQR)	22,866.60 (10,191.84, 36,892.07)	6431.76 (4861.79, 10,907.65)	-2.582	0.010 [*]
Theta (μV^2), mean (SD)	4585.63±2626.11	2894.31±1873.58	1.377	0.206 [†]
Alpha (μV^2), mean (SD)	1198.91±726.27	1121.87±687.34	0.201	0.844 [†]
Beta (μV^2), mean (SD)	165.50±115.65	196.48±107.35	-0.511	0.619 [†]
Absence of stage N3 sleep, n (%)	3 (25.0)	5 (62.5)	–	0.167 [‡]
Absence of REM sleep, n (%)	4 (33.3)	7 (87.5)	–	0.028 [‡]
Atypical sleep, n (%)	2 (16.7)	7 (87.5)	–	0.005 [‡]

Notes: [†]Student's t-test; ^{*}Mann–Whitney U-test; [‡]Fisher's Exact Test.

Abbreviations: REM, rapid eye movement; SD, standard deviation; IQR, interquartile range.

the potential utility of sleep spindles as markers of cognitive functioning.^{33,34} Regarding K-complexes, studies have associated these not only with sleep maintenance but also with executive function.³⁵ Clinical studies have confirmed that worse cognitive performance is an important trigger for delirium.^{36–38} Therefore, we speculate that the association between atypical sleep and POD may be linked to acute impairment in cognitive function caused by K-complex and sleep spindles loss.

Study Limitations

The main limitation of this study is its small sample size that, precluded establishing atypical sleep as an independent risk factor of POD. Some patients underwent emergency surgery such as aortic dissection surgery. Thus, we could not obtain preoperative PSG data of all patients. Preoperative sleep disturbance is a potential risk factor for POD.³⁹ In our study, although we excluded patients with high-risk sleep disturbance (STOP-Bang score ≥ 3) and gathered detailed history of sleep-related disease, all patients with preoperative sleep disturbance could not be excluded. Furthermore, we did not assess daytime sleep characteristics. However, a long-term (24 h) PSG study among ICU patients found that sleep architecture and quality were generally similar between the day and night.¹⁰ Thus, we believe that nighttime studies were sufficient for characterizing the sleep patterns of our patients.

Conclusion

According to our results, atypical sleep and no REM sleep on the first postoperative night are associated with POD. Further large-scale studies are needed to validate these results.

Ethics Approval and Informed Consent

All procedures performed in this study were in accordance with the ethical standards of the Union Hospital, Fujian Medical University and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Informed consent was obtained from all individual participants included in the study.

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Disclosure

The authors report no conflicts of interest in this work.

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