

Nocturnal Mean Oxygen Saturation Is Associated with Secondary Polycythemia in Young Adults with Obstructive Sleep Apnea, Especially in Men

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Objective: Whether the severity of obstructive sleep apnea (OSA) contributes to clinical polycythemia is uncertain, especially in young adults. This study aimed to assess the correlation between untreated OSA and polycythemia, controlling for multiple confounders, and to observe the difference in both genders.

Methods: All participants underwent nocturnal polysomnography. Medical comorbidities, and demographic and laboratory information were also recorded. The relationship between OSA and concomitant polycythemia in both genders was analyzed.

Results: A total of 605 young participants (383 men and 222 women), aged 30.52 ± 7.21 years, were enrolled, with an average body mass index of 32.48 ± 6.06 kg/m². Although 74.4% of patients were diagnosed with OSA, less than 10% had polycythemia. The levels of hemoglobin and hematocrit increased with the severity of OSA; only men with severe OSA had significantly higher hemoglobin, hematocrit, and polycythemia compared with those in the control group ($P < 0.01$). Hemoglobin and hematocrit significantly correlated with mean pulse oxygen saturation (SpO₂) ($P < 0.001$), but the correlation coefficients were weaker in women than in men. In logistic regression analysis, mean SpO₂, but not the apnea-hypopnea index (AHI), was found to be an independent predictor of polycythemia ($P < 0.05$). Areas under the receiver operator characteristic analysis revealed that the cutoff values of hemoglobin and hematocrit were 155.5g/L and 44.6% ($P < 0.001$), respectively, for assessing nocturnal hypoxemia in men with OSA.

Conclusion: Nocturnal mean SpO₂ was an independent predictor of polycythemia in young adults. Mean SpO₂, compared with the AHI, was more associated with polycythemia. Men were more prone to suffer from polycythemia compared with women. Hemoglobin and hematocrit values might have diagnostic utility for assessing nocturnal hypoxia severity of OSA patients, especially in men.

Keywords: obstructive sleep apnea, nocturnal hypoxia, polycythemia, erythropoiesis, biomarker

Introduction

Obstructive sleep apnea (OSA) is a common disease. Almost one billion people worldwide have OSA, with China having the highest number of affected persons, followed by the United States, Brazil, and India.¹ Polycythemia is defined as an absolute increase in hemoglobin (HGB) or hematocrit (HCT).^{2,3} It is classified as primary (e.g., polycythemia vera) or secondary [such as chronic continuous hypoxia (CCH) and renal lesions]. It results in increased circulating levels of erythropoiesis-stimulating factors. Although OSA is characterized by chronic intermittent hypoxia

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(CIH),⁴ rather than sustained hypoxemia, it was assumed that the intermittent hypoxic stress was sufficient to produce secondary polycythemia.^{3,5} Fan et al found that HGB and HCT significantly correlated with the apnea-hypopnea index (AHI) among 1087 male patients, and HCT might be a predictor of OSA severity.⁶ Relevant meta-analyses provided the same point of view.^{5,7} Several animal studies also indicated that red blood cell (RBC) count, HGB, and HCT were much higher after exposure to intermittent hypoxia.^{8–10} After continuous positive airway pressure (CPAP) therapy, RBC count, HGB, and HCT declined in patients with OSA.¹¹ Moreover, Pathak et al analyzed more than 77 million discharges and showed a significant association between OSA and secondary polycythemia with an odds ratio (OR) of 5.90 [95% confidence interval (CI): 5.64–6.17].¹² Therefore, OSA is mentioned as a cause of secondary polycythemia in the current national guidelines on the diagnosis, investigation, and management.¹³ However, other studies pointed out that intermittent nocturnal hypoxemia in OSA did not lead to clinical polycythemia.^{14–17} King et al considered that secondary polycythemia in patients with OSA was caused by a reduction in plasma volume rather than hypoxia and a recent study evaluating this association (n = 1604 participants) reported no significant differences in HGB or HCT among patients with OSA.¹⁸ Therefore, whether OSA severity contributes to clinical polycythemia remains controversial.

A wide variety of factors, including race, age, sex, altitude, and smoking, affect the values of hematological indices.^{19–23} Disparate genetic backgrounds are found among ethnic groups.²⁴ For example, African Americans have slightly lower HGB and HCT compared with Caucasians,¹⁹ and people living at high altitudes develop higher HCT to compensate for low oxygen exposure.²² The differences in hematological profiles between men and women are also well known.^{19–22,25} The values of RBC count, HGB, and HCT increased until adulthood, which were higher in men than in women at all ages,²⁰ and then decreased in the geriatric age.²⁶ People aged more than 50 years were more prone to suffer from anemia of aging.^{21,26} Previous studies generally evaluated predominantly elderly male subjects and failed to define polycythemia well.^{14,15,17,18} The extent to which untreated OSA contributes to clinical polycythemia among young Asian adults remains uncertain.²⁷ Furthermore, OSA is a heterogeneous disorder,²⁸ hypoxia indicates significant differences in OSA patients within similar AHI.²⁹ The cutoff point of hematological indices determining an individual's nocturnal

hypoxia risk of OSA was unclear.⁷ This study aimed to evaluate the association between untreated OSA and polycythemia in a relatively large young sample taking into account established confounders. Moreover, the effects of gender-related differences and the cutoff value of hematological indices were also analyzed.

Methods

Study Population

In this study, consecutive participants who underwent polysomnography (PSG) at the Sleep Disorders Center of Ruijin Hospital affiliated to Shanghai JiaoTong University from September 2009 to September 2017 were enrolled. The demographic characteristics, smoking status, and previous history of chronic diseases were recorded. The body mass index (BMI) was calculated as body weight (in kg) divided by body height (in m) squared. Hypertension was defined as a systolic blood pressure ≥ 140 mmHg, diastolic blood pressure ≥ 90 mmHg, or a clinical diagnosis of hypertension and current prescription of antihypertensive medication.³⁰ The smokers were those regularly consumed 10–20 cigarettes/day for at least 3 years.²³ The inclusion criteria were as follows: 1) participants aged 18–45 years old; and 2) Han Chinese residents of Southeast China with altitude below 10 meters. The exclusion criteria were as follows: patients with sleep disorders other than OSA (e.g., central sleep apnea syndrome, restless leg syndrome, and narcolepsy), previous treatment for OSA (e.g., CPAP, surgery, and oral device), hematologic diseases (e.g., polycythemia vera, other causes of primary/secondary erythrocytosis, anemia, leukemia, and lymphoma), chronic cardiopulmonary diseases except for primary hypertension, liver or kidney disease, neuromuscular disease, malnutrition, chronic alcoholism, malignancy, pregnancy, acute and/or chronic infection, use of medication (such as erythropoietin iron, folate, vitamin B₁₂, and contraceptives), and living at high altitudes in the past. Postmenopausal women with an established diagnosis of polycystic ovary syndrome (PCOS) or suspected of having PCOS based on hyperandrogenemia were also excluded. The study was conducted in accordance with the Declaration of Helsinki and approved by the local ethics committee of Ruijin Hospital ([2018] No.107). Written informed consent was obtained from all participants.

Polysomnographic Evaluation

All the participants underwent PSG (Alice 5, Philips Respironics, Pennsylvania, USA) at the laboratory with

alcohol, caffeine, sedatives, or hypnotics forbidden on that very day. PSG channels included four electroencephalograms, submental electromyogram, two electrooculograms (right and left), two electrocardiography channels, pulse oxygen saturation, oral and nasal airflow, nasal air pressure, thoracic–abdominal respiratory movement, snoring microphone, and body position. Only edited records with more than 4 h of good-quality signals were accepted. Apnea was defined as a respiratory event with a drop in nasal airflow by $\geq 90\%$, and hypopnea was defined as a drop in peak signal excursions by $\geq 30\%$ of pre-event baseline using nasal pressure for at least 10 seconds with a $\geq 4\%$ oxygen desaturation from pre-event baseline. Based on the AHI, the participants were categorized into four groups: control (AHI < 5), mild (AHI, 5–14.9), moderate (AHI, 15–29.9), and severe OSA (AHI ≥ 30) groups. The oxygen desaturation index (ODI) was defined as the total number of at least 4% oxygen desaturations per total sleep time in hours and the mean pulse oxygen saturation (SpO₂) was defined as the mean oxygen saturation during sleep periods at night. The percentage of sleep duration with SpO₂ $< 90\%$ (TS90) was also included.³¹

Laboratory Measurements

Fasting blood samples were taken in the morning after PSG monitoring. Complete blood cell count was analyzed using an automated hematology analysis device (CELL-DYN Ruby, Abbott Laboratories, Illinois, USA). Polycythemia was defined as an HGB level of > 160 g/L in women and > 165 g/L in men, or an HCT level of $> 48\%$ in women and $> 49\%$ in men according to the World Health Organization thresholds.³ In this study, clinical polycythemia defined by HGB (polycythemia–HGB) and HCT (polycythemia–HCT) was described.

Statistical Analysis

Data were analyzed using statistical software (IBM SPSS Statistics for Windows, version 25.0, USA). Continuous variables were reported as means \pm standard deviations (SD), or median with an interquartile range if variables were not normally distributed. Categorical variables were reported as the constituent ratio (%). The significance of the mean differences between groups was assessed using the Student *t*-test and one-way analysis of variance. For measurement data that were heteroscedastic or not normally distributed, the Mann–Whitney *U*-test and Kruskal–Wallis *H*-test were conducted for comparison among groups. The constituent ratio among the groups was

compared using the chi-square test. Spearman correlation coefficients were used to quantify the correlations among RBC count, HGB, HCT, and PSG parameters. Ordinal multivariate logistic regression analysis was performed to determine whether polycythemia was independently related to OSA severity. The effect of various variables on polycythemia–HGB and polycythemia–HCT in women was analyzed with rare events logistic regression analysis using Stata SE version 15.0 (StataCorp, TX, USA). The receiver-operating characteristic (ROC) analysis was used to estimate the predictive validity of HGB and HCT in assessing nocturnal hypoxemia (mean SpO₂ $< 90\%$) and OSA (AHI ≥ 5), and to determine the optimal cutoff value. According to the area under the curve (AUC), the diagnostic values of HGB and HCT were assessed: ≤ 0.5 was considered a chance result; 0.5–0.7, low accuracy; 0.7–0.9, moderate accuracy; and ≥ 0.9 , high accuracy. A *P* value < 0.05 for a two-tailed test was considered statistically significant.

Results

Overall Characteristics of the Participants

In total, 605 participants were included in this study (383 men and 222 women) with an average age of 30.52 ± 7.21 years (range, 18–45 years). The baseline characteristics of the study population are presented in Table 1. The mean BMI was 32.48 ± 6.07 kg/m². No difference was observed between men and women in terms of BMI. Nearly one-quarter of men, but no women, were involved in active smoking. On average, the participants had severe OSA with a mean AHI of 19.70 (range, 0–109.3). Although 450 participants (74.4%) were diagnosed with OSA, only 8.0% (*n* = 36) had polycythemia–HGB and 5.3% (*n* = 24) had polycythemia–HCT. The mean HGB and HCT values were in the clinically normal range in both genders. AHI, ODI, TS90, RBC count, HGB, and HCT were significantly higher, while mean SpO₂ was significantly lower among men compared with women. The occurrence of polycythemia increased in men than in women (*P* < 0.005 ; Table 1).

Correlations Between OSA and Interrelated Parameters of RBC

Table 2 provides a summary of demographic and clinical characteristics in male and female participants according to the OSA severity status.³¹ Patients with OSA were generally older and weighted, and more likely suffered from hypertension. The RBC count, HGB, and HCT

Table 1 Baseline Clinical, Laboratory, and Polysomnographic Data of All Participants

Variables	Total (n = 605)	Men (n = 383)	Women (n = 222)	P Value*
Age (year)	30.52 ± 7.21	32.39 ± 7.19	27.30 ± 6.03	<0.001
BMI (kg/m ²)	32.48 ± 6.07	32.21 ± 6.30	32.95 ± 5.62	0.148
Neck circumference (cm)	41.51 ± 4.01	42.40 ± 3.70	39.97 ± 4.06	<0.001
AHI (events/h)	19.70 (4.70–57.75)	40.00 (11.80–67.60)	6.05 (1.38–16.00)	<0.001
ODI (events/h)	26.30 (6.40–61.40)	43.05 (13.15–69.40)	8.10 (2.35–27.80)	<0.001
Mean SpO ₂ (%)	93.66 ± 4.45	92.78 ± 4.43	95.08 ± 4.12	<0.001
TS90 (%)	2.30 (0.10–21.44)	7.95 (0.40–30.93)	0.10 (0.00–1.90)	<0.001
RBC count (10 ³ /μL)	4.81 ± 0.47	5.00 ± 0.41	4.49 ± 0.40	<0.001
HGB (g/L)	144.17 ± 14.00	150.92 ± 11.16	132.51 ± 10.23	<0.001
HCT (%)	42.55 ± 3.95	44.42 ± 3.17	39.32 ± 2.94	<0.001
Polycythemia-HGB, n (%)	39 (6.45)	34 (8.88)	5 (2.25)	0.001
Polycythemia-HCT, n (%)	26 (4.30)	24 (6.27)	2 (0.90)	0.002
Smoking, n (%)	87 (14.38)	87 (22.72)	0 (0.00)	<0.001
Hypertension, n (%)	134 (22.15)	118 (30.81)	16 (7.21)	<0.001

Notes: Measurement data are presented as mean ± standard deviations or median with inter-quartile range if variables were not normally distributed. *Men vs women. **Abbreviations:** BMI, body mass index; AHI, apnea–hypopnea index; ODI, oxygen desaturation index; Mean SpO₂, mean oxygen saturation; TS90, percentage of sleep duration with SpO₂ < 90%; RBC, red blood cell; HGB, hemoglobin; HCT, hematocrit.

increased with the severity of OSA in both genders. The male patients in the severe OSA group had significantly higher RBC count, HGB, and HCT compared with the other two groups ($P < 0.01$, Table 2), while no difference was found between the male participants in the control group and the mild-to-moderate OSA group. The female participants showed no statistical difference in RBC count, HGB, and HCT among the three groups with respect to OSA measured by the AHI. As shown in Table 3, HCT and HGB significantly correlated with all the PSG parameters related to OSA, such as AHI, ODI, TS90, and mean SpO₂, in male participants. Although mean SpO₂ negatively correlated with the RBC count, HGB, and HCT in women, the correlation coefficients were weaker than those in men. In addition, the AHI was not associated with HGB and HCT among female participants.

Association Between Nocturnal Hypoxemia and Polycythemia Among OSA Patients

Further, the difference between patients with OSA with and without polycythemia was compared, as shown in Table 4. Patients with OSA with polycythemia were found to have higher AHI, ODI, and TS90, as well as lower mean SpO₂. In the subgroup analysis with similar AHI, the mean SpO₂ was still significantly lower in patients with severe OSA and polycythemia compared with those without polycythemia ($P < 0.001$, Table 4). In the logistic regression analysis for the risk factors for

polycythemia in men, mean SpO₂, but not the AHI, was found to be an independent predictor of polycythemia (Table 5). We further conducted logistic regression analysis in men without smokers, which demonstrated that mean SpO₂ was still independently associated with polycythemia–HGB (OR = 0.787; 95% CI: 0.645–0.961; $P = 0.018$) after adjusting for age, BMI, and hypertension. On account of low polycythemia prevalence in women, the logistic regression analysis of rare events showed that the mean SpO₂ was also independently associated with both polycythemia–HGB (OR = 0.848; 95% CI: 0.772–0.932; $P < 0.001$) and polycythemia–HCT (OR = 0.847; 95% CI: 0.747–0.961; $P = 0.010$) in the female participants after adjustment for age, BMI, and hypertension.

Given that the mean SpO₂ mainly affects the occurrence of polycythemia, the association between AHI and nocturnal hypoxia was also observed. The Spearman correlation analysis showed that the AHI significantly negatively correlated with mean SpO₂ in both male ($\rho = -0.711$, $P < 0.001$) and female ($\rho = -0.465$, $P < 0.001$) participants. Furthermore, the mean SpO₂ was significantly lower in both genders in the severe OSA group than in the control and the mild-to-moderate OSA groups ($P < 0.001$, Table 2). Compared with the mild-to-moderate OSA group, the OR of polycythemia–HGB was 2.94 (95% CI: 1.10–7.85; $P = 0.025$) and that of polycythemia–HCT was 12.11 (95% CI: 1.61–91.08; $P = 0.002$) in men in the severe OSA group, but not observed in women.

Table 2 Baseline Clinical, Laboratory, and Polysomnographic Data of Male and Female Participants Stratified by OSA Severity Status

Variables	Men					Women				
	Control (n = 53)	Mild-to-Moderate OSA (n = 110)	Severe OSA (n = 220)	P Value	Control (n = 102)	Mild-to-Moderate OSA (n = 88)	Severe OSA (n = 32)	P Value		
	Age (year)	28.70 ± 7.55	29.86 ± 7.60	34.55 ± 6.06*†	<0.001	25.94 ± 5.28	27.45 ± 5.88	31.19 ± 7.04*†	<0.001	
BMI (kg/m ²)	33.38 ± 5.08	32.85 ± 7.23	31.61 ± 6.02	0.084	31.81 ± 5.06	33.20 ± 5.27	35.90 ± 7.10*†	0.001		
Neck circumference (cm)	41.73 ± 2.52	42.49 ± 3.75	42.53 ± 3.92	0.368	39.00 ± 3.49	40.26 ± 4.15*	42.40 ± 4.59*†	<0.001		
AHI (events/h)	2.00 (0.95–3.15)	14.25 (8.60–22.43)*	64.70 (49.93–74.23)*†	<0.001	1.30 (0.60–2.53)	11.05 (7.20–16.45)*	65.90 (46.53–85.98)*†	<0.001		
ODI (events/h)	4.50 (1.65–10.40)	13.60 (7.10–26.80)*	65.55 (50.65–79.25)*†	<0.001	3.20 (1.00–8.83)	10.60 (5.08–28.93)*	85.10 (52.40–110.50)*†	<0.001		
Mean SpO ₂ (%)	95.88 ± 1.45	95.26 ± 2.33	90.86 ± 4.66*†	<0.001	96.39 ± 1.46	95.72 ± 1.70	88.90 ± 7.98*†	<0.001		
TS90 (%)	0.00 (0.00–0.35)	0.85 (0.10–3.70)*	24.50 (10.30–43.50)*†	<0.001	0.00 (0.00–0.10)	0.30 (0.00–2.58)*	26.35 (4.30–45.30)*†	<0.001		
RBC count (10 ³ /μL)	4.91 ± 0.40	4.93 ± 0.38	5.05 ± 0.42*†	0.009	4.40 ± 0.31	4.54 ± 0.42*	4.64 ± 0.55*	0.004		
HGB (g/L)	148.17 ± 10.66	148.92 ± 9.46	152.59 ± 11.79*†	0.003	132.20 ± 9.16	132.60 ± 10.57	133.25 ± 12.55	0.874		
HCT (%)	43.70 ± 3.09	43.77 ± 2.71	44.91 ± 3.32*†	0.002	39.08 ± 2.63	39.35 ± 2.67	39.98 ± 4.29	0.315		
Polycythemia-HGB, n (%)	2 (3.77)	5 (4.55)	27 (12.27)†	0.025	1 (0.98)	2 (2.27)	2 (6.25)	0.215		
Polycythemia-HCT, n (%)	1 (1.89)	1 (0.91)	22 (10.00)†	0.002	1 (0.98)	0 (0.00)	1 (3.13)	0.275		
Smoking, n (%)	8 (15.09)	22 (20.00)	57 (25.91)	0.174	/	/	/	/		
Hypertension, n (%)	9 (16.98)	28 (25.45)	69 (31.36)*	0.091	7 (6.86)	14 (15.91)*	7 (21.88)*	0.040		

Notes: Data are presented as mean ± standard deviations or median with inter-quartile range if variables were not normally distributed. *P <0.05 vs the control group; †P <0.05 vs the mild-to-moderate OSA group.
Abbreviations: OSA, obstructive sleep apnea; BMI, body mass index; AHI, apnea-hypopnea index; ODI, oxygen desaturation index; Mean SpO₂, mean oxygen saturation; TS90, percentage of sleep duration with SpO₂ < 90%; RBC, red blood cell; HGB, hemoglobin; HCT, hematocrit.

Table 3 Correlations Between Polysomnographic Parameters and RBC Count, HGB, and HCT in Male and Female Participants

Variables	Men			Women		
	RBC Counts	Hemoglobin	Hematocrit	RBC Counts	Hemoglobin	Hematocrit
Age (year)	-0.068, 0.184	0.117, 0.022	0.023, 0.659	-0.048, 0.473	-0.045, 0.504	-0.068, 0.316
BMI (kg/m ²)	0.152, 0.003	-0.067, 0.194	0.019, 0.715	0.228, 0.001	0.007, 0.915	0.097, 0.150
AHI (events/h)	0.212, <0.001	0.211, <0.001	0.212, <0.001	0.226, 0.001	0.028, 0.683	0.094, 0.164
ODI (events/h)	0.238, <0.001	0.182, <0.001	0.200, <0.001	0.285, <0.001	0.132, 0.050	0.193, 0.004
Mean SpO ₂ (%)	-0.344, <0.001	-0.304, <0.001	-0.362, <0.001	-0.303, <0.001	-0.219, 0.001	-0.274, <0.001
TS90 (%)	0.283, <0.001	0.258, <0.001	0.290, <0.001	0.245, <0.001	0.125, 0.064	0.175, 0.009

Notes: Data are Spearman's coefficients (rho, *P* value).

Abbreviations: BMI, body mass index; AHI, apnea-hypopnea index; ODI, oxygen desaturation index; Mean SpO₂, mean oxygen saturation; TS90, percentage of sleep duration with SpO₂ < 90%; RBC, red blood cell.

Contribution of HGB and HCT in Assessing Nocturnal Hypoxemia Among OSA Patients

The ROC curve analysis was performed for the cutoff values of HGB and HCT to predict nocturnal hypoxemia defined with mean SpO₂ < 90% in OSA patients. The cutoff value of HGB was 155.5 g/L in men, with a sensitivity of 60.9%, a specificity of 70.5%, and an AUC of 0.701 (95% CI: 0.628–0.775, *P* < 0.001; **Figure 1A**); and 138.5 g/L in women, with a sensitivity of 60.0%, a specificity of 72.8%, and an AUC of 0.661 (95% CI: 0.498–0.824; *P* = 0.045; **Figure 1B**). The cutoff value of HCT was 44.6% in men, with a sensitivity of 82.8%, a specificity of 57.7%, and an AUC of 0.746 (95% CI: 0.675–0.817, *P* < 0.001; **Figure 1A**); and 41.8% in women, with a sensitivity of 53.3%, a specificity of 84.5%, and an AUC of 0.677 (95% CI: 0.507–0.848, *P* = 0.027; **Figure 1B**). In addition, the AHI of ≥ 5 events/h was used to define OSA for the ROC analysis. The AUC for HGB to identify OSA was 0.574 (*P* = 0.085) in men and 0.506 (*P* = 0.882) in women; the AUC for HCT to identify OSA was 0.565 (*P* = 0.127) in men and 0.539 (*P* = 0.313) in women.

Discussion

To our best knowledge, this study was the first in analyzing a relatively large sample of young adults with untreated OSA to identify the relationship between OSA and polycythemia. Further, the differences in parameters among both genders was compared, and the cutoff points of HGB and HCT determining the nocturnal hypoxia risk of patients with OSA was calculated. This study yielded two key findings. First, nocturnal mean SpO₂, but not the AHI, was an independent predictor of clinical polycythemia after adjusting for known important confounding variables, including age, sex, BMI, smoking, hypertension, and so on. Second,

young male patients with OSA were more prone to suffer from polycythemia compared with female patients.

Several current hematologic guidelines recommend an OSA evaluation for the diagnosis of secondary polycythemia.^{3,13,32} However, clinically significant polycythemia appears uncommon in those with OSA. Hoffstein et al reported 18 of 277 (6.5%) patients with OSA having HCT ≥ 50%;¹⁴ Nguyen et al found 9.59% (119/1241) patients with OSA having HCT ≥ 48%;¹⁸ and Solmaz et al detected only 1 of 335 patients with OSA having polycythemia.¹⁷ The results were not consistent because of the different diagnostic criteria in previous studies, but the prevalence of polycythemia in OSA was low in general. In this study with young OSA subjects, 8.0% had polycythemia–HGB and 5.3% had polycythemia–HCT, consistent with the results of previous studies.

In patients with OSA, airway collapse leads to CIH during sleep,^{4,5} contributing to decreased mean nocturnal hypoxia. The data of this study showed that the mean SpO₂ decreased with the severity of OSA, and mean SpO₂ in the severe OSA group was the lowest among the three groups. Sustained hypoxia, which is reflected by mean SpO₂, leads to the activation of hypoxia-inducible factor-1 (HIF-1), resulting in an increased expression of erythropoietin (EPO) and polycythemia.³³ Previous animal studies^{8–10} reported that CIH exposure could promote the generation of HIF-1 alpha and EPO, leading to significant increases in HGB and HCT. Meanwhile, Kang et al also showed that both CIH- and CCH-exposed rats had significantly higher HCT compared with those exposed to normoxia, but HCT in the CCH group was even much higher than that in the CIH group, which might be due to the longer duration and heavier intensity of hypoxia in CCH.³⁴ Hypoxic effects on HGB and HCT are heavily dependent on the duration and intensity of the hypoxic stimulus.³⁴ Similarly, Winnicki et al

Table 4 Comparison of Clinical, Laboratory, and Polysomnographic Data in OSA Patients with or Without Polycythemia

Variables	Total OSA				Severe OSA				Total OSA				Severe OSA			
	Without Polycythemia-		With Polycythemia-		Without Polycythemia-		With Polycythemia-		Without Polycythemia-		With Polycythemia-		Without Polycythemia-		With Polycythemia-	
	HGB (n = 414)	HGB (n = 36)	P Value	HGB (n = 223)	HGB (n = 29)	P Value	HCT (n = 426)	HCT (n = 24)	P Value	HCT (n = 229)	HCT (n = 23)	P Value	HCT (n = 229)	HCT (n = 23)	P Value	
Age (year)	31.62 ± 7.17	33.53 ± 5.98	0.123	34.07 ± 6.32	34.52 ± 6.05	0.717	31.65 ± 7.17	33.92 ± 5.31	0.057	34.12 ± 6.37	34.09 ± 5.36	0.980	34.12 ± 6.37	34.09 ± 5.36	0.980	
BMI (kg/m ²)	32.45 ± 6.33	33.47 ± 6.85	0.356	31.89 ± 6.18	34.17 ± 7.10	0.068	32.39 ± 6.28	35.02 ± 7.44	0.049	31.88 ± 6.12	34.95 ± 7.59	0.072	31.88 ± 6.12	34.95 ± 7.59	0.072	
AHI (events/h)	34.40	54.50	<0.001	64.70	65.00	0.660	34.40	65.15	<0.001	64.70	65.60	0.720	64.70	65.60	0.720	
	(13.38–66.53)	(36.45–76.38)		(48.90–75.80)	(49.20–82.25)		(13.25–66.18)	(47.40–80.23)		(48.60–75.95)	(50.70–81.10)		(48.60–75.95)	(50.70–81.10)		
ODI (events/h)	42.40	53.40	0.001	68.30	61.20	0.279	42.40	64.30	<0.001	66.85	64.80	0.761	66.85	64.80	0.761	
	(13.10–70.00)	(33.40–75.60)		(51.70–81.35)	(49.00–81.88)		(13.30–69.13)	(49.00–81.98)		(51.18–81.28)	(48.50–83.50)		(51.18–81.28)	(48.50–83.50)		
Mean SpO ₂ (%)	93.20 ± 4.15	87.48 ± 7.87	<0.001	91.15 ± 4.45	86.64 ± 8.18	<0.001	93.15 ± 4.25	85.83 ± 7.70	<0.001	91.12 ± 4.61	85.73 ± 7.87	<0.001	91.12 ± 4.61	85.73 ± 7.87	<0.001	
TS90 (%)	6.60	29.45	<0.001	24.10	37.90	0.018	6.60	44.45	<0.001	24.05	47.30	0.006	24.05	47.30	0.006	
	(0.40–27.10)	(5.55–54.38)		(9.20–42.98)	(18.55–58.55)		(0.50–27.10)	(21.74–63.08)		(9.20–42.90)	(21.38–63.20)		(9.20–42.90)	(21.38–63.20)		
RBC count (10 ⁷ /ul)	4.84 ± 0.42	5.53 ± 0.47	<0.001	4.92 ± 0.40	5.62 ± 0.39	<0.001	4.85 ± 0.43	5.70 ± 0.39	<0.001	4.93 ± 0.40	5.69 ± 0.40	<0.001	4.93 ± 0.40	5.69 ± 0.40	<0.001	
HGB (g/L)	144.30 ± 12.28	170.58 ± 5.57	<0.001	147.37 ± 11.64	171.34 ± 5.86	<0.001	144.94 ± 12.67	172.42 ± 5.82	<0.001	147.89 ± 11.91	172.48 ± 5.94	<0.001	147.89 ± 11.91	172.48 ± 5.94	<0.001	
HCT (%)	42.65 ± 3.38	49.44 ± 3.72	<0.001	43.50 ± 3.20	50.31 ± 2.69	<0.001	42.74 ± 3.42	51.24 ± 2.26	<0.001	43.59 ± 3.19	51.23 ± 2.31	<0.001	43.59 ± 3.19	51.23 ± 2.31	<0.001	

Notes: Data are presented as mean ± standard deviations or median with inter-quartile range if variables were not normally distributed.

Abbreviations: OSA, obstructive sleep apnea; BMI, body mass index; AHI, apnea-hypopnea index; ODI, oxygen desaturation index; Mean SpO₂, mean oxygen saturation; TS90, percentage of sleep duration with SpO₂ < 90%; RBC, red blood cell; HGB, hemoglobin; HCT, hematocrit.

Table 5 Odds Ratio Estimates from Logistic Regression Analysis for the Risk Factors for Polycythemia in Men

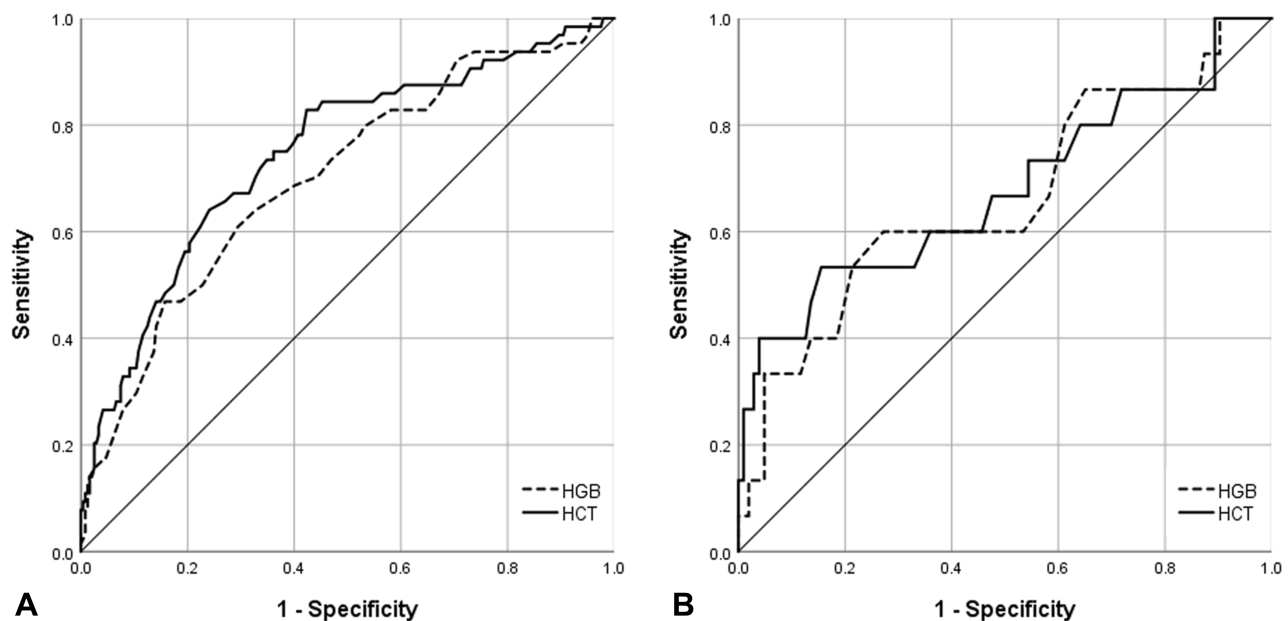
Independent Variables	Men with Polycythemia-HGB			Men with Polycythemia-HCT		
	Odds Ratio	95% Confidence Interval	P Value	Odd Ratio	95% Confidence Interval	P Value
Age (year)	1.006	0.936–1.081	0.874	1.021	0.933–1.117	0.655
BMI (kg/m ²)	1.042	0.966–1.124	0.288	1.070	0.978–1.169	0.140
AHI (events/h)	1.018	0.982–1.056	0.325	1.015	0.972–1.060	0.491
ODI (events/h)	0.982	0.950–1.015	0.290	0.981	0.943–1.021	0.342
Mean SpO ₂ (%)	0.802	0.689–0.934	0.004	0.786	0.665–0.929	0.005
TS90 (%)	0.987	0.947–1.028	0.516	0.994	0.949–1.040	0.784
Smoking	0.426	0.177–1.025	0.057	0.420	0.148–1.197	0.104
Hypertension	1.306	0.525–3.247	0.566	0.809	0.291–2.247	0.684

Abbreviations: BMI, body mass index; AHI, apnea-hypopnea index; ODI, oxygen desaturation index; Mean SpO₂, mean oxygen saturation; TS90, percentage of sleep duration with SpO₂ < 90%; RBC, red blood cell; HGB, hemoglobin; HCT, hematocrit.

reported that EPO increased only in patients with severe OSA.³⁵ In this study, although the RBC counts, HGB, and HCT increased with the severity of OSA, only the severe OSA group had significantly higher RBC counts, HGB, and HCT compared with the control group; this result was consistent with that in the study by Solmaz et al.¹⁷

OSA is a heterogeneous disorder.²⁸ In the subgroup analysis with a similar AHI, patients with polycythemia had significantly lower nocturnal mean SpO₂ compared with those without polycythemia. Although the AHI significantly correlated with mean SpO₂ in both genders, mean SpO₂, but not the AHI, was determined as an independent predictor of polycythemia. As expected, according to the ROC analysis in this study, the AUC of HGB and

HCT in detecting mean SpO₂ < 90% was more than that in detecting the AHI. These results indicated that nocturnal hypoxemia was a stronger predictor of polycythemia compared with OSA severity.² It was assumed that OSA led to polycythemia, which mainly depended on the intensity, frequency, and duration of intermittent hypoxia cycles.³⁴ This might explain why most prior cohort studies^{14,15,17,18} generally failed to demonstrate a clear association between untreated OSA (measured by the AHI) and polycythemia. However, in view of the close relationship between AHI and mean SpO₂, OSA evaluation may still be warranted in those with unexplained polycythemia.¹⁸ Additionally, the cutoff value for HGB and HCT was 155.5g/L and 44.6%, respectively, for assessing nocturnal hypoxemia in male

**Figure 1** ROC curves for HGB and HCT to predict nocturnal hypoxemia defined with as mean SpO₂ < 90% in patients with obstructive sleep apnea. (A) In men; (B) In women.

OSA patients, demonstrating moderate accuracy. This was the first report on the cutoff value of hematological indices in young adults with OSA,⁷ and it was suggested that erythrocytic parameters might serve as useful tools for screening nocturnal hypoxia severity of OSA patients.

In this study, differences in RBC count, HGB, and HCT were found between men and women. Although nocturnal mean SpO₂ independently negatively correlated with RBC count, HGB, and HCT in both genders, the correlation coefficients were weaker in women. Moreover, OSA rarely led to polycythemia in female patients, which was consistent with the results of a previous study.¹⁷ OSA is more common in male patients.³⁶ It was revealed that mean SpO₂ was significantly lower in men compared with women. Men showed higher HGB and HCT levels compared with women because of higher testosterone levels, which facilitate erythropoiesis, and genetic differences in the EPO gene.²⁵ 17 β -estradiol can also influence the expression of hypoxia-inducible genes and decrease hypoxic induction of the expression of the EPO gene.¹⁰ Further, female resilience to CIH-induced upper airway muscle dysfunction existed during acute exposure to severe hypoxic stress, but male patients had a greater loss of muscle performance.³⁷ Moreover, exposure to hypoxic significantly increased the hypoxic ventilatory response in women than in men.³⁸

This study had several limitations. First, given the cross-sectional nature of this single-center study, inferences on the cause-and-effect sequence between nocturnal mean SpO₂ and polycythemia were not possible. Second, the changes in RBC count, HGB, and HCT pre-CPAP and after-CPAP treatment were not investigated. Third, this study was a preliminary research work on young adults without further exploration of fundamental mechanisms. Moreover, further prospective studies are needed to explore the impact of OSA and nocturnal hypoxia on the polycythemia.

Conclusion

In summary, compared with the AHI, nocturnal mean SpO₂ proved to be an independent predictor of polycythemia in young adults. Men were more prone to suffer from polycythemia compared with women. Hematological parameters, such as HGB and HCT, may have diagnostic utility for assessing nocturnal hypoxia severity of patients with OSA, especially in men.

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Disclosure

The authors declare no conflicts of interest in this study.

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