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Performance of brief ICF-sleep disorders and obesity core set in obstructive sleep apnea patients

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Abstract

Background: Clinical questionnaires are mainly applied as screening tools for identification of the Obstructive sleep apnea (OSA) patients. Little attention has been paid to assess the body functions and health status of the patients. International Classification of Functioning, Disability and Health (ICF) was designed for better understanding and describing functioning and disability of patients. This study adopted the Brief ICF-Sleep Disorders and Obesity Core Set to evaluate the impairment of functioning and health status of OSA patients.

Methods: Five hundred ninety-two participants were enrolled in this cross-sectional study. Data were collected using Brief ICF-Sleep Disorders and Obesity Core Set Polysomnography was performed and basic characteristics of the patients were recorded.

Results: The scores for the component Body Functions and Code b130, b134, b140, b440, b530, s330, d160, d240, d450 of the two core sets were significantly different among the patients divided by apnea-hypopnea index (AHI) or oxygen saturation (SaO₂) nadir, but the frequency of code s330, d160, d240, d450 was low. The Body Functions component of the both sets were closely related to neck circumference (NC), body mass index (BMI), apnea-hypopnea index (AHI) of the OSA patients. Body Functions of the Brief ICF-Sleep Disorders performed better with a threshold of 4 with sensitivity, specificity and area under the receiver operating characteristic curve (AUC) as 0.62, 0.74, 0.68 (AHI ≥ 5), 0.69, 0.63, 0.66 (AHI ≥ 15), 0.75, 0.56, 0.66 (AHI ≥ 30), 0.56, 0.70, 0.63 (SaO₂ nadir ≤ 90%), 0.67, 0.66, 0.66 (SaO₂ nadir < 85%), 0.71, 0.59, 0.65 (SaO₂ nadir < 80%), separately.

Conclusion: The Body Functions component of both two sets could be an evaluation tool of impairment of body functions for OSA patients. The Brief ICF-Sleep Disorders Body Functions component performed better with a threshold of 4 and might provide a new insight for physicians to assess OSA patients.

Keywords: Obstructive sleep apnea, International classification of functioning, Disability and health, Brief ICF-sleep disorders Core set, Brief ICF-obesity Core set

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Background

Obstructive sleep apnea (OSA) is a disorders that consists of recurrent obstruction of the upper airway during sleep, nocturnal hypoxemia, and excessive daytime sleepiness [1]. Untreated OSA could lead to increased mortality and morbidity [2]. Several studies have proved that OSA is closely associated with negative consequences such as diabetes, stroke, hypertension, neurocognitive dysfunction, and even the increased risks of traffic accidents [3–6]. The golden standard for diagnosing OSA is polysomnography (PSG) [7]. However, its high cost, relative inaccessibility and time consumption could delay the diagnosis and treatment of the OSA patients. What's more, lack of awareness among physicians about the severity of patients could add difficulty to adjust the treatment without PSG. However, none of these questionnaires have investigated the impairment of functioning and health status of the patients.

The World Health Organization's (WHO) International Classification of Functioning, Disability and Health (ICF) represents a comprehensive and universally common framework to understand and describe functioning and disability, by providing lists of essential categories that are related to specific health conditions and health care contexts [8]. Lots of studies have confirmed its validity in different diseases, including schizophrenia, asthma and chronic obstructive pulmonary disease (COPD) [9–11]. Also, ICF could be applied as a unifying model for the rehabilitation strategy [12]. A total of more than 1400 categories are included in the ICF classification, and several health conditions have been developed to facilitate its application, such as ICF Core Set for COPD, obesity, sleep disorders and diabetes, but there is no specific core set of OSA. Since OSA is a disease characteristic of obesity and sleep disorders, we adopted the ICF Core Set for Sleep Disorders and Obesity to evaluate patients' functioning and health status and test validity of them. There are two versions of each ICF Core Set: the Brief and the Comprehensive Core Set. The Brief Core Set represents a minimum standard for collecting functioning data, while the Comprehensive one is supposed to represent the full spectrum of aspects of functioning and environmental factors relevant to the disease and is supposed to guide rehabilitation from a multidisciplinary perspective. Taking clinical practicability into consideration, we chose the Brief ICF Core Set of Sleep Disorders and Obesity for evaluation of patients' functioning and health status.

The Brief ICF-Sleep Disorders Core Set consists 14 categories and 4 components, the latter include Body Functions and Structures, Activities and Participation and Environmental Factors (<https://www.icf-research-branch.org/icf-core-sets-projects2/other-health-conditions/development-of-icf-core-sets-for-sleep>). For Brief-

ICF Obesity Brief Core Set, it concludes 8 categories and 3 components, the latter involve Body Functions, Activities and Participation and Environmental Factors (<https://www.icf-research-branch.org/icf-core-sets-projects2/cardiovascular-and-respiratory-conditions>). These two practical tools that cover the spectrum of symptoms and limitations in the functioning of all patients with sleep disorders or obesity, also considering the environments they live in.

As there was no investigation had confirmed the validity of the ICF Core Sets so far, this study aimed to validate Brief ICF-Sleep Disorders and Obesity Core Set in evaluating impairment of functioning and health status of OSA patients.

Methods

Patients and design

This nine-month cross-sectional study was carried out at Shanghai Zhongshan Hospital Affiliated to Fudan University from Feb 1st to Oct 31st, 2019. Adults aged over 18 years old who came to the respiratory clinic about OSA were enrolled. Exclusion criteria were as follows: (1) diagnosed OSA previously or accepted OSA treatment before; (2) acute or unstable medical or neurological conditions; (3) major psychiatric disorders or severe cognitive impairment.

Eventually 592 study subjects were enrolled and assessed by two ICF-based measurement tools, Brief ICF-Sleep Disorders and Obesity Core Set. Participants received an overnight polysomnography test (Alice-5 Respironics, Pittsburgh, Pennsylvania, USA). Based on the PSG tests, the subjects were classified by AHI or the lowest oxygen saturation (SaO₂ nadir). In this study, the mean number of apneas and hypopneas per hour of sleep (apnea-hypopnea index (AHI)) was calculated, and OSA was diagnosed if AHI was ≥ 5 events per hour [13]. Based on different values of AHI, the subjects were divided into no OSA group (AHI < 5 events/h), mild OSA group ($5 \leq \text{AHI} < 15$ events/h), moderate OSA group ($15 \leq \text{AHI} < 30$ events/h) and severe OSA group (AHI ≥ 30 events/h). According to the level of oxygen saturation (SaO₂ nadir), the participants could also be divided into group A (SaO₂ > 90%), group B ($85\% \leq \text{SaO}_2 \leq 90\%$), group C ($80\% \leq \text{SaO}_2 < 85\%$) and group D (SaO₂ < 80%).

The Sleep Disorders Brief Core Set concludes 4 components and 14 codes, and the Obesity Brief Core Set consists of 8 codes and 3 components. By comparing every category and component of the two ICF core sets among different groups of participants classified by AHI or SaO₂ nadir, we found that several categories and the Body Functions components showed potential as evaluating tools. Considering the frequency of these codes, only code b130, b134, b440 and b530 were suitable. Since these items belong to ICF-Body Functions

components which represent the impairment of physiological and psychological functions of body systems, the Body Functions components could be suitable as an evaluation tool. Correlation of the Body Functions scores with each reported questionnaire and critical parameters of participants was analyzed separately to identify validation, and then the appropriate cut-off point was determined by comparing sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV), and area under the receiver operating characteristic curve (AUC).

The objective of the study and its requirements were explained to the subjects, and all participants provided written informed consent. This study received Ethical approval from the Zhongshan Hospital Institutional Review Board. The specific process can be referred to Fig. 1, and the sociodemographic and clinical characteristics of the participants are provided in Tables 1 and 2.

Measures

In this study, the Brief ICF-Sleep Disorders and Obesity Core Sets were used to assess functioning and disability as well as environmental factors in all subjects. ICF qualifiers were applied to rate the degree of problems in each code of the Body Functions and Structures component and the Activity and Participation component with a generic five-point scale: no

problem (0), mild problem (1), moderate problem (2), severe problem (3), and complete problem (4). Environmental factors were graded by three levels: being a facilitator (-1), having no influence (0), or being a barrier (+1).

The relations of some critical parameters of the participants with ICF Core Set, including AHI, SaO2 nadir, NC and BMI were analyzed to evaluate the validity of the ICF Core Set as a measuring tool of severity of apnea, hypopnea and hypoxia.

Data analysis

The characteristics of the subjects are presented as mean ± standard deviation for continuous variables, or number and percentage for categorical variables. Comparison between two groups was assessed by Student’s t test or Mann-Whitney rank sum test (for continuous variables) and Chi-square test or Fisher’s exact tests (for categorical variables). Comparison among multiple groups were assessed by one-way Analysis of Variance (ANOVA) or Kruskal-Wallis tests (for continuous variables) and Chi-square test or Fisher’s exact tests (for categorical variables). Regression analysis was used to determine the relationship between each one of AHI, BMI, NC, SaO2 nadir, and the two ICF-based measurement tools separately. Sensitivity, specificity, PPV, NPV and AUC of different cut-off versions of

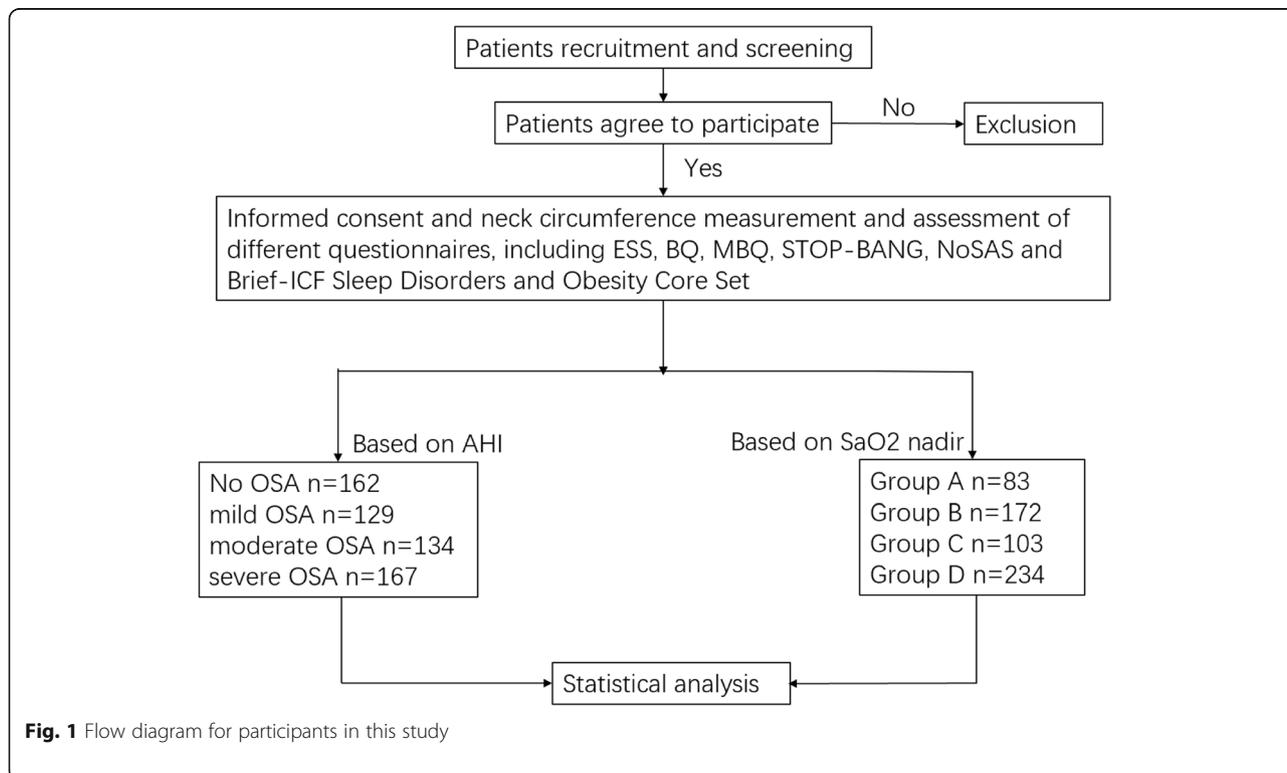


Fig. 1 Flow diagram for participants in this study

Table 1 Clinical characteristics of the study group

	Whole group (n/%)	No OSA (n/%)	OSA (n/%)	p value
Number	592/100	162/100	430/100	
Men	470/79.4	100/61.7	370/86.0	< 0.0001
age	49.21 ± 14.99	44.67 ± 16.43	50.93 ± 14.04	0.0233
BMI [kg/m²]	27.25 ± 4.66	26.13 ± 4.42	27.58 ± 7.05	0.0016
Smoking	263/44.4	46/28.4	217/50.5	< 0.0001
Current smoker	182/30.7	31/19.1	151/35.1	0.0001
Pack-years	19.91 ± 19.85	20.97 ± 23.92	19.69 ± 19.00	0.7435
Quitting smoking	93/15.7	15/9.3	78/18.1	0.0077
Quitting years	7.807 ± 7.716	6.643 ± 8.854	8.033 ± 7.524	0.4775
NC	39.47 ± 3.872	37.49 ± 3.79	39.99 ± 3.82	< 0.0001
Diabetes mellitus	87/14.7	17/10.5	70/16.3	0.0902
Arterial hypertension	268/45.3	51/31.5	217/50.5	< 0.0001
Short of breath	162/27.4	37/22.8	125/29.1	0.1479
AHI	21.00 ± 20.23	1.713 ± 1.522	26.08 ± 19.70	< 0.0001
Max apnea time	43.31 ± 28.11	15.43 ± 17.28	53.99 ± 23.81	< 0.0001
SaO₂ nadir	79.72 ± 10.52	88.19 ± 5.64	76.47 ± 10.15	< 0.0001
Max pulse	109.7 ± 19.08	109.40 ± 20.47	109.8 ± 18.54	0.8531
4% decline of SaO₂ index	20.37 ± 19.06	5.04 ± 7.75	26.50 ± 18.79	< 0.0001
rhinitis	149/25.2	40/24.7	109/25.3	0.9157
pharyngitis	127/21.5	24/14.8	103/24.0	0.0181
nasosinusitis	16/2.7	5/3.1	11/2.6	0.7772
nasal polyp	5/0.8	2/1.2	3/0.7	0.6180
polyp of vocal cord	4/0.7	3/1.9	1/0.2	0.0645
polypectomy	11/1.9	2/1.2	9/2.1	0.7356
Nasal septum deviation	6/1.0	1/0.6	5/1.2	1.00

Data are presented as Mean ± standard deviations. Differences were compared between no OSA and OSA groups
 The data of significant difference were marked in bold.
 NC neck circumference, BMI body mass index, AHI apnea-hypopnea index, SaO₂ oxygen saturation

the Body Functions component scores were calculated including respective confidence intervals. Stata version 15 (Stata Corp, LLC, Texas, USA) and GraphPad Prism (version 6; GraphPad Software, San Diego, California, USA) were used for all statistical analysis and drawing graphs, and the level of significance was set to *p* < 0.05. The data of significant difference were marked in bold.

Results

Clinical characteristics, PSG examination and questionnaire results of the participants

Five hundred ninety-two participants were enrolled in this cross-sectional survey. According to PSG examination results, the subjects were divided into no OSA group (*n* = 162, AHI < 5 events/h), mild OSA group (*n* = 129, 5 ≤ AHI < 15 events/h), moderate OSA group (*n* =

Table 2 PSG examination parameters and questionnaire results of the participants

Category	No OSA(<i>n</i> = 162)	Mild (<i>n</i> = 129)	Moderate(<i>n</i> = 134)	Severe (<i>n</i> = 167)	p value
AHI	1.65 ± 1.54	9.55 ± 3.09	21.87 ± 4.33	48.21 ± 14.19	< 0.0001
Max apnea time	15.43 ± 17.28	36.31 ± 15.86	51.62 ± 18.84	69.35 ± 22.21	< 0.0001
SaO₂ nadir	88.19 ± 5.64	82.76 ± 7.69	79.21 ± 7.98	69.96 ± 9.244	< 0.0001
Max pulse	109.40 ± 20.47	107.80 ± 18.55	109.60 ± 19.15	111.50 ± 18.02	0.25
4% decline of SaO₂ index	5.04 ± 7.75	12.67 ± 7.40	21.41 ± 15.94	40.85 ± 16.9	< 0.0001

Data are presented as Mean ± standard deviations. Differences were compared among the 4 groups
 SaO₂, oxygen saturation. The data of significant difference were marked in bold

134, 15 ≤ AHI < 30 events/h), severe OSA group (n = 167, AHI ≥ 30 events/h) group based on the classification of AHI. Except max pulse, values of AHI, max apnea time, SaO2 nadir and 4% decline of SaO2 index were significantly different among the 4 groups.

Differences of scores for each component of brief ICF-sleep disorders and obesity Core set among groups of participants

Severity of OSA can be measured by both AHI and lowest oxygen saturation (SaO2 nadir). According to the level of SaO2 nadir, the participants could also be divided into group A (n = 83, SaO2 > 90%), group B (n = 172, 85% ≤ SaO2 ≤ 90%), group C (n = 103, 80% ≤ SaO2 < 85%), group D (n = 234, SaO2 < 80%). The Sleep Disorders Brief Core Set concludes 4 components and the Obesity Core Set consists of 3 components. To identify whether the Brief ICF-Sleep and Obesity Core Sets could uncover the difference between participants divided by AHI and SaO2, the scores for each component of the two core sets were compared among the groups, by adding the scores of each category. When the subjects were classified by AHI, the Body Functions and Structures, Activities and Participation of Sleep Disorders Core Set and the Body Functions of Obesity Core Set could tell the difference among the groups (shown in Table 3). But only the scores for the Body Functions component of the two core sets were significantly different among the 4 groups classified by SaO2 nadir (shown in Table 4).

Frequency and extent of the participants’ impairments in different categories of sleep disorders and obesity brief Core set

The Sleep Disorders Brief Core Set concludes 14 codes. To further validate the performance of the categories, 592 participants were divided into 4 groups classified by AHI and SaO2 nadir, evaluating each category of the core set. As S-Tables 1 & 3 show, code b130, b134, b140, b440, s330, d160 and d240 display significant differences among subjects divided by AHI and SaO2 nadir, however, the positive rate of code s330, d160 and d240 was low and code b140 could not tell the difference

when patients were classified by AHI. Compared with the other 3 parts, the categories of Body Functions showed great potential as an evaluation tool to assess the severity of the patients, and the more severe a patient was, the more likely his score was higher. There was no difference among the 4 groups of environmental factors part. Almost all the participants consider their family members, the health systems and policies as benefits.

The Obesity Brief Core Set involves 3 components and 8 codes, and it was applied to assess functioning and disability as well as environmental factors in 4 groups of participants based on AHI and SaO2 nadir classification (shown in S-Tables 2 & 4). The scores for code b130, b530, d240 and d450 were significantly different among the four groups of participants, but code d450 could not tell the difference when patients were divided by SaO2 nadir. Same as the Sleep Disorders Brief Core Set, there was no significant difference in the Environmental Factors component, and the frequency of code d240 and d450 was very low (seen in supplement materials, S-Tables 1, 2, 3 and 4).

Potential for the body functions component of brief ICF-sleep disorders and obesity Core set as tools to evaluate body functioning of OSA patients

In Table 3, the parts of Body Functions and Structures, Activities and Participation of Brief ICF-Sleep Disorders Core Set show potential as an evaluation tool of body functioning and health status when the classification method was AHI. But when the patients were divided by SaO2 nadir, only Body Functions component worked (shown in Table 4). From S-Table 1 and 3, we find out that although the scores of some codes of the part Body Structures and Activities and Participation were significantly different among groups, the frequency was low. Taking all this into consideration, codes for Sleep Functions could be suitable as an evaluation tool of patients’ impairment of functioning. In the same way, the codes of Body Functions of Brief ICF-Obesity Core Set could be potential as a measuring tool, too. Then the scores of Body Functions of the two core sets were compared in

Table 3 Scores of Brief ICF-Sleep Disorder and Obesity Core Set in participants based on AHI classification

Category	No OSA (n = 162)	Mild (n = 129)	Moderate (n = 134)	Severe (n = 167)	p value
ICF-Sleep Body Functions	2.00 ± 1.58	3.30 ± 1.68	3.76 ± 2.09	4.68 ± 2.74	< 0.0001
ICF-Sleep Body Structures	0.23 ± 0.57	0.23 ± 0.60	0.41 ± 0.81	0.63 ± 1.11	0.0010
ICF-Sleep Activities and Participation	0.27 ± 1.00	0.23 ± 0.98	0.34 ± 1.07	0.51 ± 1.36	0.0338
ICF-Sleep Environmental Factors	-2.96 ± 0.25	-2.98 ± 0.24	-2.94 ± 0.34	-2.87 ± 0.50	0.1838
ICF-Obesity Body Functions	2.27 ± 1.23	2.58 ± 1.34	2.45 ± 1.11	3.01 ± 1.40	< 0.0001
ICF-Obesity Activities and Participation	0.38 ± 0.83	0.51 ± 1.18	0.79 ± 1.57	0.71 ± 1.52	0.1934
ICF-Obesity Environmental Factors	-1.98 ± 0.18	-1.98 ± 0.15	-1.96 ± 0.23	-1.95 ± 0.23	0.5767

Data are presented as Mean ± standard deviations. Differences were compared among the 4 groups. The data of significant difference were marked in bold

Table 4 Scores of Brief ICF-Sleep Disorder and Obesity Core Set in participants based on SaO2 classification

Category	A(n = 83)	B(n = 172)	C(n = 103)	D(n = 234)	p value
ICF-Sleep Body Functions	2.65 ± 1.69	3.25 ± 1.93	4.06 ± 2.14	4.80 ± 2.29	< 0.0001
ICF-Sleep Body Structures	0.08 ± 0.37	0.02 ± 0.12	0.03 ± 0.16	0.02 ± 0.13	0.47
ICF-Sleep Activities and Participation	0.29 ± 0.80	0.37 ± 1.22	0.35 ± 1.05	0.50 ± 1.27	0.47
ICF-Sleep Environmental Factors	0.046 ± 0.21	0.068 ± 0.31	0.075 ± 0.27	0.066 ± 0.29	0.89
ICF-Obesity Body Functions	2.05 ± 1.19	2.42 ± 1.26	2.61 ± 1.27	2.92 ± 1.31	< 0.0001
ICF-Obesity Activities and Participation	0.45 ± 1.16	0.74 ± 1.72	0.54 ± 1.22	0.76 ± 1.81	0.44
ICF-Obesity Environmental Factors	0.03 ± 0.17	0.04 ± 0.23	0.04 ± 0.19	0.03 ± 0.16	0.98

Data are presented as Mean ± standard deviations. Differences were compared among the 4 groups. The data of significant difference were marked in bold

patients divided by AHI and SaO2 nadir, which was shown in Fig. 2. Scores of Body Functions of each two groups of the participants were significantly different no matter the classification was by AHI or SaO2. However, the differences in mild and moderate groups or in Group B and C could not be recognized by Body Functions component of ICF-Obesity Core Set.

between BMI and NC and the Body Functions component scores of both core sets, we found significant relations. Then we analyzed the correlation between the Body Functions component and some critical parameters of PSG examination, including AHI and SaO2 nadir, and positive correlation was found (Table 5).

Correlation of different parameters of participants with brief ICF-sleep disorders and obesity core set

Body mass index (BMI) and neck circumference (NC) are proved to be closely related to the morbidity and severity of OSA [14]. By comparing the correlation

Performance of the brief-ICF sleep disorders and obesity body functions in participants classified by AHI or SaO2

The values of sensitivity, specificity, PPV, NPV and AUC were calculated for all cut-off points (mild, moderate and severe OSA), considering AHI or SaO2 (shown in Tables 6 and 7 and Fig. 3). When the patients were

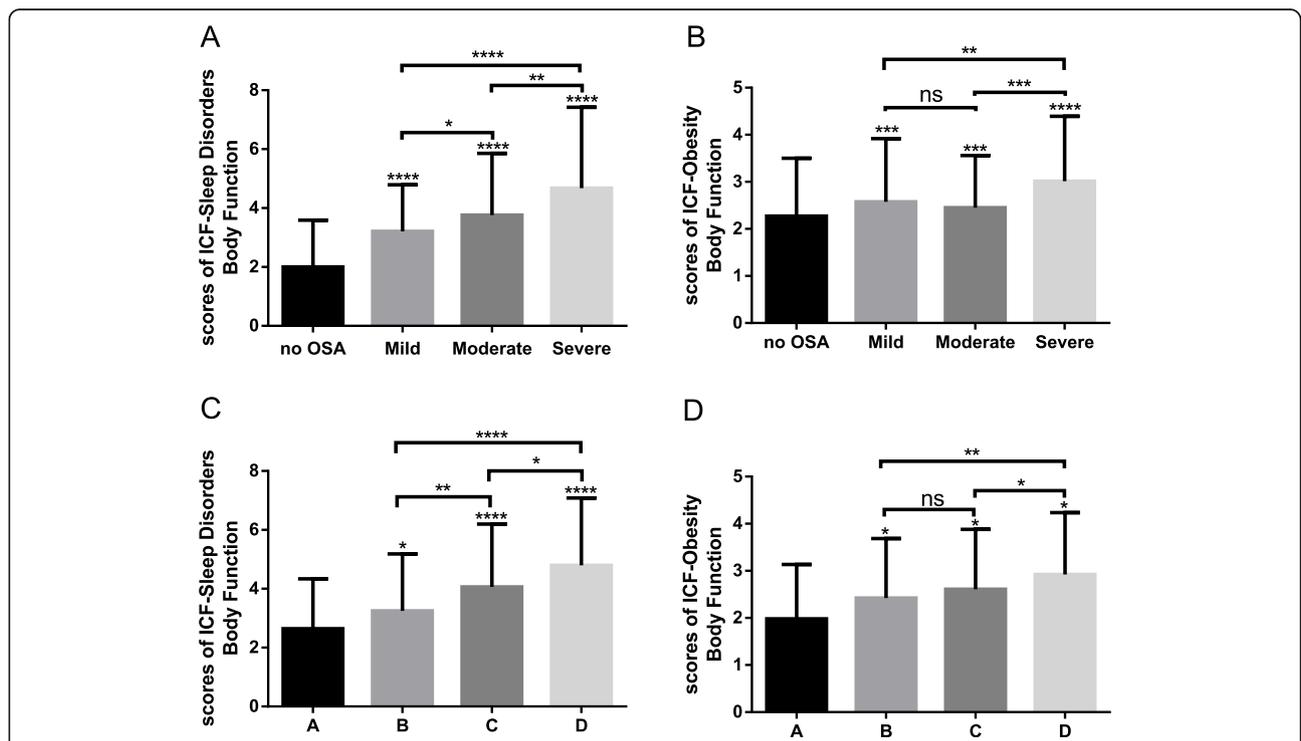


Fig. 2 Scores of Body Functions component among different groups classified by AHI and SaO2 nadir. **a-b** Differences of scores of ICF-Sleep Disorder and Obesity Body Functions between each two groups of participants classified by AHI. **c-d** Differences of scores of ICF-Sleep Disorder and Obesity Body Functions between each two groups of participants classified by SaO2. Patients were classified by AHI in Figure **a-b**, by SaO2 in Figure **c-d**. Statistical significant values are identified with *(α = 0.05), ****p < 0.0001, ***p < 0.001, **p < 0.01, *p < 0.05, ns, not significant

Table 5 Correlation of Participants’ different parameters with the Brief ICF-Sleep Disorder or Obesity Body Functions

Category	β^s	95%CI ^s	β^o	95%CI ^o
AHI	0.053	(0.044–0.063) ****	0.017	(0.011–0.023) ****
BMI	0.089	(0.041–0.14) ****	0.19	(0.17–0.21) ****
NC	0.10	(0.052–0.15) ****	0.13	(0.10–0.16) ****
SaO2 nadir	–0.073	(–0.0911 - -0.055) ****	–0.029	(–0.039 - -0.018) ****
Brief ICF-Obesity or Sleep Disorder Body Functions scores	0.86	(0.73–1.00) ****	0.30	(0.25–0.34) ****

Different superscript indicates different ICF core set. ^s means Brief ICF-Sleep Disorder Core Set Body Functions component. ^o means Brief ICF-Obesity Core Set Body Functions component. β means regression coefficient. CI means confidence interval
 The data of significant difference were marked in bold
 NC neck circumference, SaO2 oxygen saturation

classified by AHI, the threshold of 4 was suitable for Body Functions of ICF-Sleep Disorders with largest AUC and acceptable sensitivity and specificity, while the threshold of 3 of ICF-Obesity Core Set would be more suitable. As Table 6 shows, however the status of the patient changes, from normal to OSA, or mild to moderate, or moderate to severe, the ICF-Sleep Disorders Core Set with a threshold of 4 would be applicable, and whatever severity of hypoxia a patient is, the ICF-Sleep Disorders Core Set behaves better than the Obesity Core Set. When the participants were classified by SaO2, cut-off \geq 4 for ICF-Sleep Disorders Core Set and cut-off \geq 3 for ICF-Obesity Core Set might be suitable for evaluating

severity of hypoxia, considering sensitivity, specificity and AUC. In general, the Body Functions component of the Brief ICF-Sleep Disorders Core Set with cut-off \geq 4 showed better sensitivity, specificity and AUC, and could be more reliable to assess the severity of apnea, hypopnea and hypoxia of OSA patient, as well as impairment of functioning.

Discussion

Multiple studies have reported the application of the screening questionnaires in identifying OSA patients, such as ESS, BQ, MBQ, STOP-BANG and NoSAS questionnaires, however, little attention has been paid to

Table 6 Predictive parameters of various Body Functions score cut-offs for different AHI levels

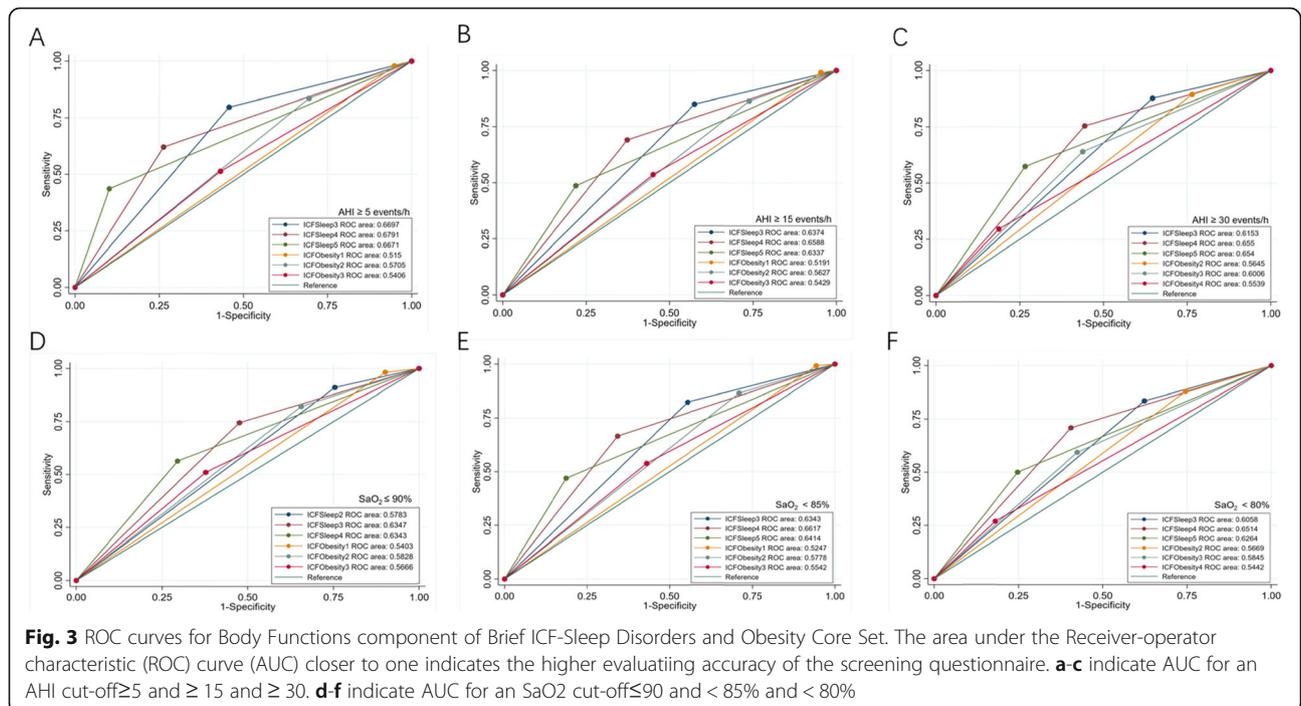
Cut-off points	Sensitivity	Specificity	PPV	NPV	AUC
AHI \geq 5 events/h as the diagnostic criteria for OSA	Prevalence of OSA = 430/592 (72.64%)				
ICF-Sleep \geq 3	0.80	0.54	0.83	0.48	0.67 (0.62–0.72)
ICF-Sleep \geq 4	0.62	0.74	0.87	0.41	0.68 (0.63–0.73)
ICF-Sleep \geq 5	0.44	0.90	0.92	0.36	0.67 (0.63–0.71)
ICF-Obesity \geq 1	0.98	0.05	0.75	0.46	0.52 (0.49–0.54)
ICF-Obesity \geq 2	0.84	0.31	0.77	0.40	0.57 (0.52–0.62)
ICF-Obesity \geq 3	0.51	0.57	0.77	0.29	0.54 (0.49–0.59)
AHI \geq 15 events/h as the diagnostic criteria for OSA	Prevalence of OSA = 301/592 (50.84%)				
ICF-Sleep \geq 3	0.85	0.42	0.58	0.75	0.64 (0.60–0.68)
ICF-Sleep \geq 4	0.69	0.63	0.64	0.68	0.66 (0.62–0.70)
ICF-Sleep \geq 5	0.49	0.68	0.55	0.62	0.63 (0.59–0.68)
ICF-Obesity \geq 1	0.99	0.05	0.50	0.85	0.52 (0.50–0.53)
ICF-Obesity \geq 2	0.86	0.26	0.52	0.67	0.56 (0.53–0.60)
ICF-Obesity \geq 3	0.54	0.55	0.53	0.56	0.54 (0.50–0.59)
AHI \geq 30 events/h as the diagnostic criteria for OSA	Prevalence of OSA = 167/592 (28.21%)				
ICF-Sleep \geq 3	0.88	0.35	0.33	0.89	0.62 (0.58–0.65)
ICF-Sleep \geq 4	0.75	0.56	0.38	0.86	0.66 (0.61–0.70)
ICF-Sleep \geq 5	0.57	0.73	0.44	0.82	0.65 (0.60–0.70)
ICF-Obesity \geq 2	0.89	0.24	0.30	0.86	0.56 (0.53–0.60)
ICF-Obesity \geq 3	0.64	0.56	0.35	0.81	0.60 (0.55–0.65)
ICF-Obesity \geq 4	0.30	0.81	0.37	0.76	0.55 (0.51–0.60)

PPV positive predictive value, NPV negative predictive value, AUC the area under the curve

Table 7 Predictive parameters of the various Body Functions score cut-offs for different SaO2 nadir levels

Cut-off points	Sensitivity	Specificity	PPV	NPV	AUC
SaO2 ≤ 90% as the diagnostic criteria for OSA		Prevalence of OSA = 509/592 (85.98%)			
ICF-Sleep ≥ 2	0.91	0.25	0.89	0.30	0.58 (0.52–0.63)
ICF-Sleep ≥ 3	0.59	0.39	0.86	0.13	0.63 (0.57–0.70)
ICF-Sleep ≥ 4	0.56	0.70	0.92	0.20	0.63 (0.57–0.70)
ICF-Obesity ≥ 1	0.98	0.10	0.88	0.46	0.54 (0.50–0.58)
ICF-Obesity ≥ 2	0.82	0.34	0.89	0.23	0.58 (0.52–0.65)
ICF-Obesity ≥ 3	0.51	0.62	0.90	0.17	0.57 (0.50–0.63)
SaO2 < 85% as the diagnostic criteria for OSA		Prevalence of OSA = 337/592 (56.93%)			
ICF-Sleep ≥ 3	0.82	0.45	0.67	0.65	0.63 (0.59–0.68)
ICF-Sleep ≥ 4	0.67	0.66	0.72	0.59	0.66 (0.62–0.71)
ICF-Sleep ≥ 5	0.47	0.81	0.77	0.53	0.64 (0.60–0.68)
ICF-Obesity ≥ 1	0.99	0.06	0.59	0.85	0.52 (0.51–0.54)
ICF-Obesity ≥ 2	0.87	0.29	0.62	0.62	0.58 (0.54–0.62)
ICF-Obesity ≥ 3	0.54	0.57	0.63	0.48	0.55 (0.51–0.60)
SAO2 < 80% as the diagnostic criteria for OSA		Prevalence of OSA = 234 /592 (39.53%)			
ICF-Sleep ≥ 3	0.84	0.74	0.81	0.77	0.61 (0.57–0.65)
ICF-Sleep ≥ 4	0.71	0.59	0.54	0.75	0.65 (0.61–0.70)
ICF-Sleep ≥ 5	0.50	0.75	0.58	0.69	0.63 (0.58–0.67)
ICF-Obesity ≥ 2	0.88	0.25	0.44	0.76	0.57 (0.53–0.60)
ICF-Obesity ≥ 3	0.59	0.58	0.48	0.68	0.58 (0.54–0.63)
ICF-Obesity ≥ 4	0.27	0.82	0.50	0.63	0.54 (0.50–0.58)

PPV positive predictive value, NPV negative predictive value, AUC the area under the curve



evaluation of the functioning and health status of the OSA patients. OSA results in sleep fragmentation and recurrent hypoxemia and is associated with various adverse consequences that affect not only organ systems, but also quality of life [15]. Hence evaluation of functioning and disability of OSA patients is as important as diagnosis of the disease. ICF is a common framework for understanding and describing functioning and disability. Previous studies have reported that ICF cores sets offered recognized standards for assessing the functioning status of patients in various diseases, such as ADHD, schizophrenia and COPD, and played a critical role in rehabilitation [9, 16–18]. However, there is no specific ICF core set for OSA patients. The most relevant core sets that have been created were sleep disorder and obesity core sets. Using the sleep disorder and obesity ICF core set as guidelines, the given health status can be evaluated, and the management process of a patient can be assessed. Besides that, ICF Core Sets can also be used to rate the content validity of health status measures of the patients. The sleep disorder and obesity core sets were not created primarily for OSA, but the items of the core set could still be potential as evaluation tools for the patients. The aim of our investigation was to select the potential evaluation tools with the two coresets, and it is the first one to confirm the validity of Brief ICF-Sleep Disorders and Obesity Core Set in OSA patients in China.

Five hundred ninety-two participants were evaluated with Brief ICF-Sleep Disorders and Obesity Core Set. Most patients regard environmental factors as facilitators, including family and health systems. Though scores for Structures of Pharynx were significantly different between the groups, the frequency was low, so was the component Activities and Participation. Anatomical and non-anatomical factors are both critical causes of OSA [19]. Different from the anatomical factors, the non-anatomical factors, especially the tonicity of the pharyngeal dilators could not be evaluated by imaging, which resulting in the low detection of impairment of Structure of pharynx. Inability to maintain attention and vigilance is common reported in OSA patients [13], but the attentional deficiency is not common across all patients. It may occur in some, but not all OSA patients [20]. This could be the reason why the scores of code Attention functions and Focusing attention are much higher in the severe group and group D, but the exact number of patients who had the attention problems was low.

Since the significant codes with high frequency belong to the Body Functions component and the latter was statistically different among groups, this part was chosen as the measuring tool to evaluate the impairment of physiological and psychological functions. As an

evaluation tool, specificity and sensitivity should be considered as well as AUC. The ICF-Sleep Disorders Body Functions component with a threshold of 4 proved to be applicable in assessing impairment of functioning of all patients. No matter what classification the patients were, divided by AHI or SaO₂, or what cut-off the AHI or SaO₂ was, cut-off \geq 4 has the largest AUC, more suitable sensitivity and specificity. This means whatever degree of apnea, hypopnea or hypoxia the patient was, the new evaluation method could be applicable to measure the impairment of body functions, or even help physicians adjust the patient's treatment in time without PSG tests according to the scores, but all this need to be validated in the subsequent researches.

Compared with the Body Functions component of ICF-Obesity Core Set, the ICF-Sleep Disorders Body Functions has larger AUC, better sensitivity and specificity. Although obesity is an admitted risk factor for OSA [21], an overweight person did not certainly end up with OSA. Abdominal and neck fat accumulation seem to be more related to OSA than general obesity [22]. That could be reason why the ICF-Obesity Core Set could be applicable in evaluation, but the ICF-Sleep Disorders Core Set performed better.

Current clinical guideline applies AHI as the standard measurement of OSA [13]. However, OSA is a heterogeneous syndrome, influenced by varying risk factors, pathophysiological causes, clinical manifestations and consequences [23]. Hence, relying on AHI alone in diagnosis is not enough [24]. SaO₂ nadir was shown to be closely related to concentration of high-sensitivity C-reactive protein, and lower SaO₂ nadir could lead to higher proinflammatory markers [25]. In the meantime, previous studies mostly concentrate on the performance of questionnaires in subjects classified by AHI, little attention has been paid to the validity of questionnaires in patients with different level of SaO₂ nadir. Our research validated the Brief ICF Core Set in patients classified by AHI as well as SaO₂ nadir. The results show that the Body Functions component proved to be applicable in assessing impairment of functioning in patients with various degrees of AHI and SaO₂ nadir. Meanwhile, the body functions component of the ICF core set positively correlated with AHI and SaO₂ nadir and was significantly different among groups, which means it would be useful for measuring the severity of apnea, hypopnea and hypoxia. In summary, the Body Functions component was proved to be applicable in evaluating the impairment of body functions, and closely relate to severity of apnea, hypopnea and hypoxia. But to confirm the threshold of 4 of the Brief ICF-Sleep Disorders Core Set as a follow-up indicator, more researches of a larger sample need to be done.

Conclusions

This research is the first one to investigate the performance of the Brief ICF-Sleep Disorders and Obesity Core Set in OSA patients classified by AHI and SaO₂ nadir. It reveals the Body Functions component of Brief ICF-Sleep Disorders Core Set closely related to the degree of apnea, hypopnea and hypoxia, and could be applicable in evaluation of impairment of functioning and health status with a threshold of 4. This approach provides a new insight for physicians to evaluate OSA patients and offers an evaluation tool to assess the functioning and health status of them.

Supplementary information

Supplementary information accompanies this paper at <https://doi.org/10.1186/s12931-020-01404-1>.

Additional file 1. S-Table 1 Measuring impairments of the participants classified by AHI with Brief ICF-Sleep Disorders Core Set.

Additional file 2. S-Table 2 Measuring impairments of the participants classified by AHI with Brief ICF-Obesity Core Set.

Additional file 3. S-Table 3 Measuring impairments of the participants classified by SaO₂ nadir with Brief ICF-Sleep Disorders Core Set.

Additional file 4. S-Table 4 Measuring impairments of the participants classified by SaO₂ nadir with Brief ICF-Obesity Core Set.

Abbreviations

OSA: Obstructive sleep apnea; ICF: International Classification of Functioning, Disability and Health; WHO: World Health Organization; NC: Neck circumference; BMI: Body mass index; AHI: Apnea-hypopnea index; COPD: Chronic obstructive pulmonary disease; ANOVA: Analysis of Variance; SaO₂: Oxygen saturation; PPV: Positive predictive value; NPV: Negative predictive value; AUC: Area under the receiver operating characteristic curve; ROC: Receiver-operator characteristic

Authors' contributions

Primary investigation and study design, Liang Xie; Sample acquisition, Liang Xie and Qinhan Wu; Sample processing, Liang Xie, Qinhan Wu and Weiping Hu; Data analysis, Liang Xie, Wenjing Li and Guiling Xiang; Drafting of manuscript, Liang Xie, Shengyu Hao, Chengyao Guo, Hong Jiang; Data interpretation, Xiaodan Wu; Manuscript revision, Liang Xie, Xu Wu and Shanqun Li; Final approval, all.

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Availability of data and materials

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate

The objective of the study and its requirements were explained to the subjects, and all participants provided written informed consent. This study received Ethical approval from the Zhongshan Hospital Institutional Review Board.

Consent for publication

"Not applicable".

Competing interests

The authors declare that they have no competing interests relevant to this work.

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References

- Young T, Palta M, Dempsey J, Skatrud J, Weber S, Badr S. The occurrence of sleep-disordered breathing among middle-aged adults. *N Engl J Med*. 1993; 328:1230–5.
- Gooneratne NS, Richards KC, Joffe M, Lam RW, Pack F, Staley B, Dinges DF, Pack AI. Sleep disordered breathing with excessive daytime sleepiness is a risk factor for mortality in older adults. *Sleep*. 2011;34:435–42.
- Arzt M, Young T, Finn L, Skatrud JB, Bradley TD. Association of sleep-disordered breathing and the occurrence of stroke. *Am J Respir Crit Care Med*. 2005;172:1447–51.
- Elmasry A, Lindberg E, Berne C, Janson C, Gislason T, Awad Tageldin M, Boman G. Sleep-disordered breathing and glucose metabolism in hypertensive men: a population-based study. *J Intern Med*. 2001;249:153–61.
- Teran-Santos J, Jimenez-Gomez A, Cordero-Guevara J. The association between sleep apnea and the risk of traffic accidents. Cooperative group Burgos-Santander. *N Engl J Med*. 1999;340:847–51.
- Quan SF, Wright R, Baldwin CM, Kaemingk KL, Goodwin JL, Kuo TF, Kaszniak A, Boland LL, Caccappolo E, Bootzin RR. Obstructive sleep apnea-hypopnea and neurocognitive functioning in the sleep heart health study. *Sleep Med*. 2006;7:498–507.
- Flemons WW, Douglas NJ, Kuna ST, Rodenstein DO, Wheatley J. Access to diagnosis and treatment of patients with suspected sleep apnea. *Am J Respir Crit Care Med*. 2004;169:668–72.
- Seidel M. The International Classification of Functioning, Disability and Health. *Nervenarzt*. 2005;76:79–90 quiz 91–72.
- Lage SM, Jacome C, Oliveira A, Araujo AG, Pereira DAG, Parreira VF. Validation of the international classification of functioning, disability and health core set for obstructive pulmonary diseases in the perspective of adults with asthma. *Disabil Rehabil*. 2018;42(1):86–92.
- Huang J, Reinhardt JD, Dai R, Wang P, Zhou M. Validation of the brief international classification of functioning, disability, and health core set for obstructive pulmonary disease in the Chinese context. *Chron Respir Dis*. 2019;16:1479973119843648.
- Nuno L, Barrios M, Rojo E, Gomez-Benito J, Guilera G. Validation of the ICF Core sets for schizophrenia from the perspective of psychiatrists: an international Delphi study. *J Psychiatr Res*. 2018;103:134–41.
- Stucki G, Cieza A, Melvin J. The international classification of functioning, disability and health (ICF): a unifying model for the conceptual description of the rehabilitation strategy. *J Rehabil Med*. 2007;39:279–85.
- Kapur VK, Auckley DH, Chowdhuri S, Kuhlmann DC, Mehra R, Ramar K, Harrod CG. Clinical practice guideline for diagnostic testing for adult obstructive sleep apnea: an American Academy of sleep medicine clinical practice guideline. *J Clin Sleep Med*. 2017;13:479–504.
- Whyte A, Gibson D. Adult obstructive sleep apnoea: pathogenesis, importance, diagnosis and imaging. *J Med Imaging Radiat Oncol*. 2019; 64(1):52–66.
- Levy P, Kohler M, McNicholas WT, Barbe F, McEvoy RD, Somers VK, Lavie L, Pepin JL. Obstructive sleep apnoea syndrome. *Nat Rev Dis Primers*. 2015;1: 15015.
- Gomez-Benito J, Guilera G, Barrios M, Rojo E, Pino O, Gorostiaga A, Balluerka N, Hidalgo MD, Padilla JL, Benitez I, Selb M. Beyond diagnosis: the core sets for persons with schizophrenia based on the World Health Organization's international classification of functioning, disability, and health. *Disabil Rehabil*. 2018;40:2756–66.
- Finger ME, Wicki-Roten V, Leger B, Escorpizo R. Cross-cultural adaptation of the work rehabilitation questionnaire (WORQ) to French: a valid and reliable instrument to assess work functioning. *J Occup Rehabil*. 2019;29:350–60.
- Mahdi S, Ronzano N, Knuppel A, Dias JC, Albdah A, Chien-Ho L, Almodayfer O, Bluschke A, Karande S, Huang HL, et al. An international clinical study of ability and disability in ADHD using the WHO-ICF framework. *Eur Child Adolesc Psychiatry*. 2018;27:1305–19.

19. Osman AM, Carter SG, Carberry JC, Eckert DJ. Obstructive sleep apnea: current perspectives. *Nat Sci Sleep*. 2018;10:21–34.
20. Jackson ML, Howard ME, Barnes M. Cognition and daytime functioning in sleep-related breathing disorders. *Prog Brain Res*. 2011;190:53–68.
21. Lam JC, Mak JC, Ip MS. Obesity, obstructive sleep apnoea and metabolic syndrome. *Respirology*. 2012;17:223–36.
22. Oltmanns KM. Abdominal fat and sleep apnea: the chicken or the egg? Response to Pillar and Shehadeh. *Diabetes Care*. 2008;31:e61 author reply e62.
23. Zinchuk A, Yaggi HK. Phenotypic subtypes of OSA: a challenge and opportunity for precision medicine. *Chest*. 2019;157(2):403–20.
24. Gouveris H, Bahr K, Jahn C, Matthias C, Simon P. The apnea-hypopnea index underestimates systemic inflammation in women with sleep-disordered breathing. *J Women's Health (Larchmt)*. 2018;27:920–6.
25. Yilmaz Avci A, Avci S, Lakadamyali H, Can U. Hypoxia and inflammation indicate significant differences in the severity of obstructive sleep apnea within similar apnea-hypopnea index groups. *Sleep Breath*. 2017;21:703–11.

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