



Worldwide prevalence and associated risk factors of obstructive sleep apnea: a meta-analysis and meta-regression

Anna Beatriz de Araujo Dantas¹ · Flávio Magno Gonçalves^{2,3} · Agnes Andrade Martins^{1,3} · Giorvan Ânderson Alves⁴ · José Stechman-Neto^{2,3} · Camila de Castro Corrêa^{3,5} · Rosane Sampaio Santos^{2,3} · Weslania Viviane Nascimento^{3,6} · Cristiano Miranda de Araujo^{2,3} · Karinna Veríssimo Meira Taveira^{1,3}

Received: 18 August 2022 / Revised: 6 March 2023 / Accepted: 8 March 2023
© The Author(s), under exclusive licence to Springer Nature Switzerland AG 2023

Abstract

Purpose This study aimed to identify the prevalence of obstructive sleep apnea (OSA) and associated risk factors globally. **Methods** Six databases and registrations and three grey databases were explored for observational field research. Independently and impartially paired reviewers selected research, gathered data, and evaluated the methodological quality. Heterogeneity was investigated using subgroup analysis and meta-regression following the moderating variable in a meta-analysis of proportions with a random-effects model. The critical appraisal instrument developed by the Joanna Briggs Institute was used to evaluate the listed studies' methodology. The certainty of the evidence was evaluated using the GRADE tool.

Results A total of 8236 articles were collected during the database search, resulting in 99 articles included for qualitative synthesis, and 98 articles were included for the meta-analysis. The estimated combined prevalence of OSA was 54% [CI 95% = 46–62%; I² = 100%]. Mean age, percentage of moderate-severe cases, and the sample's body mass index (BMI) did not affect the heterogeneity that was already present when meta-regressed ($p > 0.05$). Ninety-one studies were deemed to have a low risk of bias, while eight were deemed to have a moderate risk. For OSA prevalence outcomes, the GRADE criteria were considered very low.

Conclusion Approximately half of the people worldwide have OSA. High BMI, increasing age, and male gender are described as risk factors in the literature, but these covariates do not affect pre-existing heterogeneity.

Keywords Obstructive sleep apnea · Risk factors · Prevalence · Systematic review

Introduction

Recurrent bouts of partial or complete blockage of the upper airway during sleep constitute obstructed sleep apnea (OSA). When a person sleeps, the pharyngeal region, a portion of the upper airway, may constrict or collapse [1, 2]. Etiologies includes neuromuscular, anatomical, mechanical, and chemical factors [3, 4] and can be classified as mild, moderate, or severe [5]. Polysomnography (PSG) is the reference standard for its diagnosis [6, 7].

Oxygen desaturation, excessive sleepiness, poor attention, sleep awakenings, fragmented sleep, exhaustion, and changes in blood pressure and heart rate can all be symptoms of OSA [1, 2]. These symptoms can affect the patient's quality of life and increase the risk of road traffic accidents and depression [3].

Age, gender, and obesity are all recognized risk factors for OSA [4–7]. In addition, this condition is also associated

This study was presented at the 30th Brazilian Congress of Speech Therapy in 2022 (city of João Pessoa, Paraíba) with its abstract having been published in Portuguese in the annals of the Congress at: <https://na01.safelinks.protection.outlook.com/?url=https%3A%2F%2Fp.sbfa.org.br%2Fcongressosbfa30anos%2F&data=05%7C01%7C%7C8197c4d19ac248c5ceea08db1e49d583%7C84df9e7fe9f640afb435aaaaaaaaaaaa%7C1%7C0%7C638137074704263785%7CUnknown%7CTWFpbGZsb3d8eyJWljiMC4wLjAwMDAiLlJlZ2ludWZliLlCBjTiI6IklhaWwiLCJXVCi6Mn0%3D%7C3000%7C%7C%7C&sdata=IkjCI2SY2DI6Y7IHM42ek2aR615w3YpiRcDFJ%2Fg8IIA%3D&reserved=0>

✉ Karinna Veríssimo Meira Taveira
karinnataveira@narsm.com.br; karinna.taveira@ufrn.br

Extended author information available on the last page of the article

with cardiovascular diseases [8], such as hypertension, congestive heart failure, nocturnal arrhythmias, atrial fibrillation, stroke, and type 2 diabetes [9]. Given that OSA is a public health issue because of its high incidence and correlation with higher mortality and morbidity, it is important to understand these relationships [10]. In order to comprehend the effects of OSA on the population and to develop public interventions, greater knowledge is necessary.

A recent systematic review addressed the frequency of OSA in the general population [7]. However, the search strategy was conducted through March 2016. Furthermore, this review used only two databases and did not use grey literature. Thus, there is a need to develop an updated systematic review with a broader search strategy to provide a qualitative and quantitative synthesis of the subject. Therefore, this study aimed to systematically review the literature on OSA prevalence and its associated risk factors worldwide.

Methods

Protocol

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) standards were followed while reporting this systematic review [11].

Eligibility criteria

This study used the PECOS strategy [12] in this analysis to specify the following criteria for choosing studies: global population, denoted by the symbol “P”; Obstructive sleep apnea exposure (E); Comparison (C): Since this is prevalence research, there is no comparison; Obstructive sleep apnea prevalence and risk factors comprise the outcomes (O); Cross-sectional studies comprise the study design (S).

This research sought to determine the prevalence of OSA in the global population. Thus, we included studies that evaluated the global population, performed PSG for the OSA diagnosis, and measured the apnea-hypopnea index (AHI). There were no exclusion criteria related to language, year of publication, gender, or ethnicity.

Studies were excluded based on the following criteria: (1) Studies that did not evaluate the prevalence of obstructive sleep apnea; (2) Studies that did not perform home or complete PSG for OSA diagnosis, such as studies assessing OSA only with questionnaires or exclusively based on clinical diagnosis without testing; (3) Studies that did not provide AHI; (4) Studies that reported the prevalence of central sleep apnea; (5) Hospital-based studies and studies assessing OSA prevalence among patients with resistant hypertension, chronic kidney disease, heart failure, and stroke/neurological disease since OSA prevalence is much higher in these

groups than in the global population; (6) Reviews, letters, books, conference abstracts, case reports, case series, opinion articles, technical articles, and guidelines; (7) Studies for which the full text were not available, even after requesting full texts from the authors.

Information sources and search strategies

The Medical Subject Headings (MeSH) and the Health Sciences Descriptors were used to choose the descriptors (DeCS). The search strategy used a word combination (Appendix 1), with the word combinations and search phrases modified in each database as deemed necessary. In the following electronic databases, Embase, Latin American and Caribbean Literature on Health Sciences (LILACS), LIVIVO, PubMed (including MEDLINE), Scopus, and Web of Science, we performed the search in May 2021 and updated it in April 2022. In order to find the grey literature, we also ran searches on Google Scholar, OpenGrey, and ProQuest Dissertation and Theses. In order to locate papers missed by the searches in the electronic databases, we manually searched the references of the chosen manuscripts. In order to find unpublished or potentially relevant studies, experts with recognized expertise in the field were contacted by e-mail. The articles were imported into a reference manager program (EndNote X7.0.1, Thomson Reuters) to remove duplicate studies.

Study selection

Initially, for calibration, the reviewers considered the eligibility requirements and used them on a small sample of retrieved papers to gauge inter-examiner agreement. Two investigators (ABAD and FMG) independently carried out a methodical analysis of the study titles and abstracts after attaining an acceptable degree of agreement (the Kappa Coefficient of Agreement 0.70) (phase 1). Subsequently, phase 2 involved the two reviewers reading the entire manuscripts independently. In the event of a disagreement, a third author (AAM) was consulted. We used the Rayyan website (<http://rayyan.qcri.org>) to blind the reading of references and allow this to occur independently and discreetly in both phases.

Data collection process and data listing

Two investigators (ABAD and FMG) independently collected the following information from the studies: author, year of publication and country, sample characteristics, OSA definition criteria, OSA prevalence, and associated risk factors.

Risk of study bias

The methodological quality was evaluated using the Joanna Briggs Institute Critical Appraisal (JBI) tool for

cross-sectional studies [13]. Using criteria that the reviewers have qualified with responses like “yes”, “no”, “uncertain”, or “not applicable”, it was possible to pinpoint the causes of bias through this procedure. The percentage of affirmative responses to the evaluation instrument’s questions was used to categorize each study. When up to 49% of responses were “yes”, the risk of bias was considered high. When 50 to 69% of responses were “yes”, moderate, and when more than 70% of responses were “yes”, low [14]. The JBI tool’s criteria were examined separately by two investigators (ABAD and FMG). The third and fourth investigators (AAM and KVMT) were consulted in case of disagreements. RevMan 5.4 software created the figures (Review Manager 5.4, The Cochrane Collaboration).

Summarization measures

The primary outcome included the prevalence of OSA, which was reported using relative or absolute frequencies and their accompanying 95% confidence intervals (95% CI).

Summary of the results

A meta-analysis of proportions with a random effect model was performed using the DerSimonian and Laird estimator to calculate the Tau-squared [15]. Using the Higgins inconsistency index (I^2) [16], heterogeneity was evaluated. The Freeman-Tukey double arcsine transformation method ensured the data followed an approximately normal [17] distribution. In order to estimate the weight of the studies included in the analysis, the effect size of each study was weighted using the inverse variance method, calculating the estimate based on the inverse proportion of the study variance [17]. Confidence intervals of 95% CI were considered and calculated using the Clopper-Pearson method. All analyses and graphics were made using the statistical software Rstudio version 1.2.1335 (Rstudio Inc, Boston, USA).

Assessment of reporting bias

The evaluation of the existence of publication bias was performed using the Egger test. In order to assess the influence of studies with low statistical power and higher probability of type-2 error, we conducted a sensitivity analysis removing studies with insufficient sample size to ensure adequate statistical power. Thus, we used the estimates obtained by the global effect for sample calculation, considering an infinite population, a 5% margin of error, and a 95% confidence level. We considered the following variables for subgroup analyses to assess the effect of potential confounding and risk factors on heterogeneity in the analysis: age group; country of origin of the sample; degree of OSA included in the sample (mild, moderate, severe, or not reported); initial

screening for eligible patients or PSG performed in the entire sample; and conceptualization used by the authors to define the outcome (OSA or syndrome). Meta-regressions with a random-effects model were also performed to assess the impact of study characteristics on the observed variance between the effect sizes of the studies included in the analysis, considering the following factors: mean sample age, percentage of cases diagnosed as moderate-severe, and mean BMI of the sample. We used a 5% level of significance.

Assessment of the certainty of evidence

The Grading of Recommendations Assessment, Development, and Evaluation (GRADE) standards were used to determine the degree of certainty in the available cumulative evidence [18]. Publication bias, indirect evidence, consistency, and imprecision were assessed by two investigators (AAM and KVMT). Consensus settled disagreements and a third investigator (CMA) was consulted if needed. The online program GRADEpro (McMaster University) generated an evidence profile. The level of evidence was classified as “high”, “moderate”, “low”, or “very low”.

Results

Study selection

The search technique revealed 8236 studies from the six electronic databases and the grey literature (excluding duplicate studies). A total of 199 studies were included for full-text evaluation after phase 1, while 100 studies were eliminated after phase 2. Thus, 99 studies for the qualitative analysis and 98 for the quantitative analysis were included. Figure 1 provides a summary of the full selection procedure. Appendix 2 presents the exclusionary factors.

Characteristics of the studies

All included studies were classified as cross-sectional. Fourteen studies were conducted in the USA [19–32], followed by nine in Brazil [4, 33–40]; eight in China [41–48] and Australia [49–56]; seven in India [57–63]; five in Turkey [64–68], Thailand [69–73], and Spain [74–78]; four in Korea [79–82]; three in Germany [83–85]; and two in Canada [86, 87], Slovakia [88, 89], Japan [90, 91], Belgium [92, 93], Iran [94, 95], Greece [96, 97], and Italy [98, 99]. The remaining 17 were conducted in Sweden [100], Lebanon [101], Mexico [102], Oman [6], Ireland [103], Vietnam [104], Poland [105], Norway [106], South Africa [107], Colombia [108], Malaysia [109], Saudi Arabia [110], Portugal [111], Ecuador [112], Egypt [113], Mongolia [114], and Bangladesh [115] (Fig. 2).

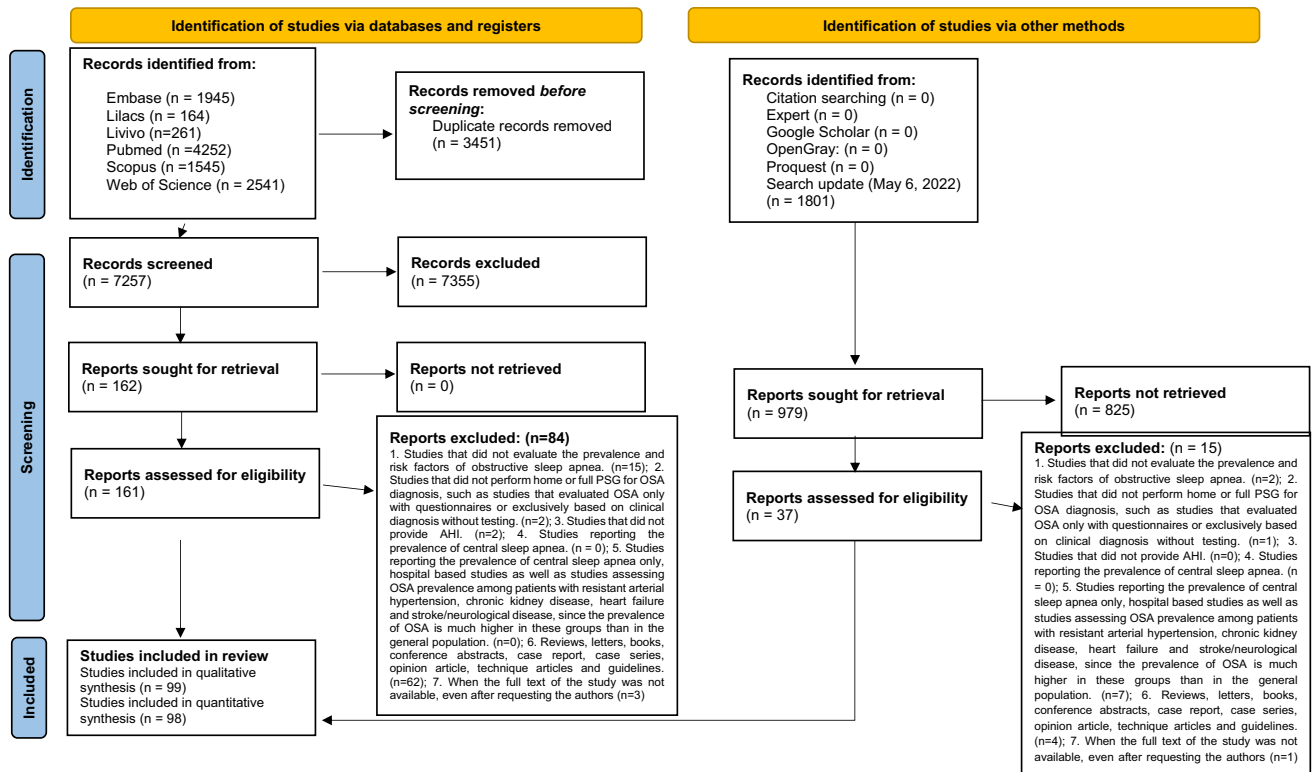


Fig. 1 PRISMA 2020 flow diagram for new systematic reviews which included searches of databases, registers and other sources

The sample size ranged from 22 [50] to 19371 [23] participants. All articles used PSG to identify OSA. However, 21 [23, 32, 57, 60–64, 66–68, 71, 84, 90, 91, 98, 104, 106, 110, 113, 115] used a screening questionnaire primarily aimed at directing high-risk patients to PSG, and two [75, 99] used either PSG or polygraphy. Table 1 provides the detailed characteristics of the included studies.

Estimates of prevalence for each subgroup reported in the 99 studies varied significantly. Table 2 shows the estimates for each subgroup. Adults and the elderly had the highest prevalence of OSA at 85%. When using PSG, the prevalence of OSA was 58%, and patients with mild to moderate OSA were the most prevalent when considering the classification of OSA. In terms of nationality, Mongolia and Taiwan (93%), Greece (82%), and Iran (81%) had the highest prevalence of OSA (Fig. 3).

Risk of study bias

Regarding the overall risk of bias, eight studies [21, 69, 84, 90, 94, 104, 109, 113] were classified as having a moderate risk of bias, and 91 [4, 6, 19, 20, 22–68, 70–83, 85–89, 91–93, 95–103, 105–108, 110–112, 114, 115] were classified as having a low risk of bias. The lack of a thorough description of the study subjects and environments, the

reliability of exposure assessment, and the absence of confounding factor control measures stood out as these studies' methodological flaws. Appendix 3 and Fig. 3 show the results of the risk of bias analysis.

Results of the individual studies

The studies differed in the OSA prevalence, ranging from 12% [79] to 92% [37]. Sixty-one studies used the AHI > 5 to classify the presence of OSA, while the remaining studies used the AHI ranging from 1 to 30.

Among the risk factors for OSA, the most recurrent were body mass index (BMI) [21, 26–28, 32, 35, 38, 43, 45, 46, 50, 52, 55, 57, 61, 66, 67, 72, 80–82, 84, 93, 97, 98, 101, 102, 107, 109, 110, 114], cited by 31 articles, followed by age [26–28, 35, 38–40, 45, 52, 55, 60, 61, 71, 72, 81–83, 97, 100, 101, 107, 109–111, 115], cited by 25 articles; the male gender [24, 26, 27, 35, 47, 51, 52, 55, 60–62, 71, 72, 81–83, 92, 93, 101–104, 110, 115], cited by 24 articles; neck circumference [32, 35, 40, 43, 46, 48, 57, 66, 67, 72, 98, 102, 109, 113, 114], cited by 15 articles; obesity [39, 40, 47, 60, 61, 71, 87, 92, 105, 110, 111, 115], cited by 12; and hypertension [6, 26, 45, 52, 60, 71, 76, 77, 100, 109, 110], cited by 11 articles. In addition, some articles brought alcohol consumption [39, 71] and smoking [60, 71, 97] as risk factors.

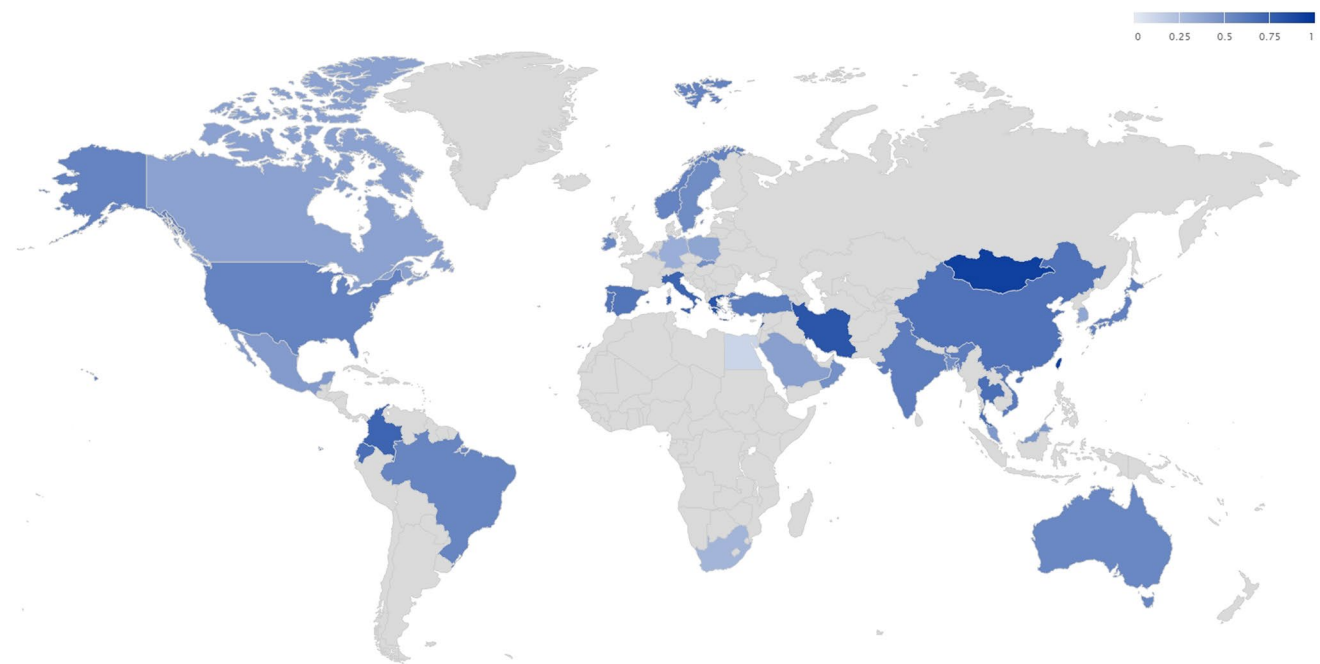


Fig. 2 General heatmap of OSA prevalence and country where the studies were performed

Summary of the results

The analysis included 47634 people, with the OSA prevalence estimated to be 56% [95% CI = 51–61%; $I^2 = 99\%$] (Fig. 3). We calculated the sample size using this obtained prevalence and considering an infinite population, a 5% margin of error, and a 95% confidence level, which required 379 individuals. As a result, studies with fewer than the required sample size were excluded from the analysis. Following this sensitivity analysis, the obtained prevalence was 54% [95% CI = 46–62%; $I^2 = 100\%$]. Following the sample size calculation, 69 studies were excluded from the sensitivity analysis and excluded in the meta-regression.

Reporting bias

The Egger test found no publication bias ($p > 0.05$).

Table 2 shows the prevalence estimates for each subgroup. The mean age, the % of moderate-severe cases, and the BMI of the sample variables did not influence the heterogeneity when meta-regressed ($p > 0.05$).

Level of evidence

The certainty of evidence was very low according to the GRADE system. Due to the use of non-probability sampling and the insufficient management of confounding variables, the domain “risk of bias” was classified as “severe”. Due to high levels of inconsistency and sources of heterogeneity

that subgroup analysis and meta-regression had failed to detect, the “inconsistency” domain was “very severe”. When prevalence was evaluated as a secondary endpoint, the subject of “indirect evidence” was considered “very serious”. Publishing bias was considered undetectable because a thorough search of the subject’s literature, including grey literature, was conducted. Also, we discovered no potential conflicts of interest in the included studies (Table 3).

Discussion

This systematic review assessed the evidence on OSA prevalence and its associated risk factors worldwide. When a meta-analysis only included studies with sufficient sample sizes, the prevalence of OSA was 54%. In addition, the qualitative synthesis revealed that high BMI, growing age, and male gender were the risk factors most frequently identified. While these findings are similar to those of a systematic review previously published by Senaratna and colleagues [7]. However, there were significant differences in the age groups, diagnostic criteria, and cut-off levels of the indexes used to report sleep apnea in the research.

In order to have a better understanding of OSA, it is critical to emphasize that its etiology is multifactorial [116, 117]. At least four major characteristics or phenotypes are known to play a role in OSA pathogenesis [118]. A study conducted in Oman sought to assess the relationship between BMI, age, and gender disparities in the prevalence

Table 1 Characteristics of the included studies ($n=99$)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Abdel et al, 2022, Egypt ¹¹⁶	160 (all male)	48.7 (5.6)	OSAS screening (160) and PSG (42)	AHI > 5	42	Weight, neck circumference, systolic blood pressure, PaO ₂ and PaCO ₂ are the most useful predictors of suspecting OSAS.
Abraham et al, 2021, USA ²²	456 263 male 193 female	5.7 (3.2)	PSG	AHI ≥ 2	252	Children with large tonsil size had 3.18 times the odds of OSA Patients who reported their race as White had 0.28 times the odds of OSA; Patients with more severe symptoms had 1.72 times the odds of OSA compared to patients with only snoring
Akkoyunlu et al, 2013, Turkey ⁶⁷	241 (all male)	42 (9.9)	OSAS screening (241) and PSG (42)	AHI ≥ 5 + symptoms (sleep attacks or EDS, unsatisfying sleep, fatigue or insomnia, witnessed heavy snoring and/or breathing pauses referred by the partner) OR AHI ≥ 15 AHI > 15	General: 34 AHI 5-15: 11 AHI 15-30: 9 AHI > 30: 14	Has not been reported
Al-Abri et al, 2011, Oman ¹⁰	608 405 male 203 female	Male: 45.3 (12.1) Female: 39.8 (13.2)	PSG	AHI ≥ 15 AHI > 15	General: 300 Male: 232 Female: 68	In men, the prevalence of hypertension, IHD and hypothyroidism. In women, the prevalence of COPD / asthma, IHD, diabetes and hypothyroidism.
Anuntasree et al, 2005, Thailand ⁷²	755 371 male 384 female	10.2 (0.5)	OSA screening (755) and PSG (10)	AHI ≥ 1	10	Has not been reported
Arora et al, 2022, USA ²⁴	170 (all male)	53.7 (3.1)	PSG	AHI ≥ 15	58	BMI

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Arazola-Cortés et al, 2017, Mexico ¹⁰⁵	51 23 male 28 female	54.1 (12.1)	PSG	Mild OSA: AHI \geq 5 and < 15 Moderate OSA: AHI \geq 15 and < 30 Severe OSA: AHI \leq 30	General: 22 Mild OSA: 10 Moderate OSA: 6 Severe OSA: 6	Gender (male), BMI and neck perimeter
Bains et al, 2020, USA ²⁵	456 256 male 193 female	5.7 (3.2)	PSG	Severe OSA: AHI > 10	66	African American race patients had 3.7 times the odds of severe OSA compared to with race patients (95% CI, 1.2-10.8). Patients aged 2 to 3 years had 2.2 times the odds of severe OSA compared to children aged 4 to 6 years (95% CI, 1.2-4.0). The mean OHAI was significantly higher in patients: -With symptoms of chronic snoring compared with those with no chronic snoring ($p=0.011$). -Who had apnea during sleep compared with those without sleep apnea ($p=0.008$).
Bariş et al, 2017, Turkey ⁶⁸	131 62 female 69 male	101.9 (59.2) months	PSG	Mild OSA: AHI > 1 and < 5 Moderate OSA: AHI > 5 and < 10 Severe OSA: AHI > 10	General: 44 Mild OSA: 13 Moderate OSA: 8 Severe OSA: 23	
Benn et al, 2021, Australia ⁵²	82 41 male 41 female	45.5	PSG	Mild OSA: AHI \geq 5 and < 15 Moderate OSA: AHI \geq 15 and < 30 Severe OSA: AHI \leq 30	General: 72 Mild: 16 Moderate: 15 Severe: 41	Has not been reported
Berger et al, 2012, USA ²⁶	19,371	Male 41.2 (11.2) Female 41.2 (9.7)	OSA screening (19371) and PSG (2429)	Possible OSA: AHI \geq 5 and \leq 10 Definite OSA: AHI > 10	Higher-risk drivers OSA -Definite OSA: 1424 -Possible OSA: 253 Lower-risk drivers OSA -Definite 156 -Possible OSA: 38	Has not been reported

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Bruyneel et al, 2010, Belgium ⁹⁵	2,093	Patients under age 40: 34 (5) Patients over age 40: 56 (10)	PSG	AHI > 5	General: 781 Patients under age 40: 121 Patients over age 40: 660	Compared to >40 years OSA patients, younger patients were more often male, African and obese (BMI>30).
Caia et al, 2020, Australia ⁵³	22 (all male)	23.8 (3.6)	Home-based PSG	Mild OSA: AHI \geq 5 and < 15 Moderate OSA: AHI \geq 15 and < 30 Severe OSA: AHI \geq 30	General: 10 Mild OSA: 7 Moderate OSA: 3 Severe OSA: 0	Increased BMI and higher skinfold thickness
Campanholo et al, 2022, Brazil ³⁶	701 317 male 384 female	50.2 (13.3)	PSG	AHI \geq 15	270	Has not been reported
Celikhisar et al, 2019, Turkey ⁶⁹	965 (all male)	45.1 (5.5)	OSA screening (965) and PSG (110)	Mild OSA: AHI 5 -15 Moderate OSA: AHI 15 - 30 Severe OSA: AHI >30	General: 69 Mild OSA: 35 Moderate OSA: 20 Severe OSA: 14	BMI values and neck circumference.
Celikhisar et al, 2020, Turkey ⁷⁰	1450 (all male)	42.1 (5.5)	OSAS screening (1450) and PSG (162)	Mild OSA: AHI 5 -15 Moderate OSA: AHI 15 - 30 Severe OSA: AHI >30	General: 127 Mild OSA: 39 Moderate OSA: 35 Severe OSA: 53	BMI, neck circumference and waist/hip ratio
Chau et al, 2003, China ⁴⁴	62 36 male 26 female	6.8 (3.6)	PSG	AHI >1	22	Snoring every night
Chen et al, 2021, China ⁴⁵	93 (all male)	28.8 (3.0)	PSG	AHI \geq 5	19	Has not been reported
Cho et al, 2013, Korea ⁸²	746 252 male 494 female	59.3 (7.2)	PSG	Moderate OSA: AHI 15 > 30 Severe OSA: AHI \geq 30	General: 90	Has not been reported
Chuanprasitkul et al, 2021, Thailand ⁷³	189 119 male 70 female	Normal group: 81.1 (28.1) months OSA group: 79.0 (31.4) months	PSG	Mild OSA: has not been reported Moderate OSA: OAH \geq 5 and < 10 Severe OSA: OAH \geq 10	General: 167 Mild: 62 Moderate: 47 Severe: 58	Arousal index were predictors of pediatric OSA
Coussa-Koniski et al, 2020, Lebanon ¹⁰⁴	663 506 male 157 female	60.8 (16.0)	PSG	Mild OSA: AHI 5 - 15 Moderate OSA: AHI 15 - 30 Severe OSA: AHI > 30	General: 526 Mild : 99 Moderate : 200 Severe : 227	Gender (male), age and BMI class.

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Dashzeveg et al, 2021, Mongolia ¹¹⁷	205 105 male 100 female	48.7 (12.6)	PSG	Mild OSA: AHI \geq 5 and \geq 15 Moderate OSA: AHU > 15 and < 30 Severe OSA: AHI \geq 30	General: 191 Mild : 50 Moderate : 41 Severe : 100	BMI, neck/waist circumference, and blood pressure
de Menezes Júnior et al, 2021, Brazil ³⁷	119 (all male)	35.3	PSG	Mild OSA: AHI \geq 5 and \leq 15 Moderate and severe OSA: AHI \geq 15	Mild OSA: 45 Moderate and severe OSA: 55	Has not been reported
Deegan et al, 1996, Ireland ¹⁰⁶	250 201 male 49 female	OSA group: 50 (11) Non-OSA group: 45 (13)	PSG	AHI \geq 15	136 (54%)	Gender (male). Female patients with OSA were more likely to be postmenopausal.
Del Bruto et al, 2021, Ecuador ¹⁵	180 64 male 116 female	71.8 (7.2)	PSG	Mild OSA: 5-15 Moderate-to-severe OSA: >15	Mild OSA: 83 Moderate-to-severe OSA: 38	Has not been reported
Domany et al, 2018, USA ²³	92 44 male 48 female	4 - 11 years	PSG	OI > 2	General: 40 Preterm children: 26 Term children: 14	The risk for OSA could be stratified according to controller gain, plant gain, cardiorespiratory coupling and gestational age.
Dorkova et al, 2010, Slovakia ⁹¹	non-Roma subjects: 137 Roma subjects: 23. 114 male 46 female	50.0 (1.1)	PSG	AHI > 5	General: 100 Roma subjects: 21 non-Roma subjects: 89	Roma background and WC were independent predictors of AHI ($p < 0.001$)
Dostálová et al, 2020, Slovakia ⁹²	42 35 male 7 female	64.6 (7.5)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: \geq 30	General: 34 Mild OSA: 13 Moderate and Severe OSA: 21	Has not been reported
Duarte et al, 2017, Brazil ³⁸	456 290 male 166 female	43.7 (12.5)	PSG	Mild OSA: AHI \geq 5 and < 15 Moderate-Severe OSA: AHI \geq 15 and < 30 Severe OSA: AHI \geq 30	General: 357 Moderate-Severe OSA: 237 Severe OSA: 130	Age, BMI, neck circumference, prevalence of OSA was higher in males.
Durán et al, 2001, Spain ⁸⁰	2148 1050 male 1098 female	OSA high risk: -Male: 52.0 (9.9) -Female: 55.5 (11.3) OSA low risk: -Male: 47.4 (10.8) -Female: 46.9 (10.5)	OSA Screening (2148) and PSG (555)	AHI \geq 10	OSA High-risk: 187 OSA low-risk: 15	Hypertension

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Duong-Quy et al, 2018, Vietnam ¹⁰⁷	667 374 male 293 female	44 +12	OSA screening and PSG	Mild OSA: 5-14 Moderate OSA: 15-30 Severe OSA: >30	General: 57 Mild OSA: 22 Moderate OSA: 19 Severe OSA: 16	Male
Drakou et al, 2021, Greece ⁹⁹	272 200 male 72 female	52.9	PSG	Mild OSA: 5-14 Moderate OSA: 15-29 Severe OSA: ≥ 30	General: 206 Mild OSA: 46 Moderate OSA: 75 Severe OSA: 85	Has not been reported
Ferreira-Santos et al, 2021, Portugal ¹¹⁴	318	61 (11)	PSG	Mild OSA: 5-14 Moderate OSA: 15-29 Severe OSA: ≥ 30	General: 203 Mild OSA: 111 Moderate OSA: 50 Severe OSA: 42	There was an increasing prevalence of symptoms and comorbidities, the latter describing older and obese patients, and a substantial increase in some comorbidities, suggesting their beneficial use as combined predictors
Fietze et al, 2018, Germany ⁸⁶	1,208 652 male 556 female	54	PSG	Mild-to-severe OSA: ≥ 5 Moderate-to-severe OSA: ≥ 15 ; Severe OSA: ≥ 30	Mild-to-severe OSA: 555 (45.9%) Moderate-to-severe OSA: 256 (21%) Severe OSA: 93 (8%)	Men, aged 60 years or older. Gender differences decreased with increasing age
Foster et al, 2017, USA ²⁷	209 108 male 101 female.	34.3 (± 8.5)	PSG	AHI > 5	General: 142 (68%) Male: 92 (85.2%) Female: 50 (49.5%)	Men
Franklin et al, 2013, Sweden ¹⁰³	399 female	48 (46–49)	PSG	OSA: ≥ 5 Moderate OSA: ≥ 15 Severe OSA: ≥ 30	General: 200 (50%) Mild OSA: 124 (31%) Moderate OSA: 57 (14%) Severe OSA: 23 (5.9%)	Overweight, age, hypertension, witnessed apnoea and snoring
Garbarino et al, 2016, Italy ¹⁰¹	283 male professional truck drivers	42.3 (± 8.3)	OSA Screening and PSG (139)	Mild OSA: 5-14 Moderate OSA: 15-30 Severe OSA: >30	General: 101 (36%) Mild OSA: 44 (16%) Moderate OSA: 33 (12%) Severe OSA: 24 (8%)	BMI, neck circumference and neck-chin angle
Ghosh et al, 2020, India ⁶⁰	1000	51,4 (12,3)	OSA Screening (1000) and PSG (80)	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: >30	General: 75 (36%) Mild OSA: 44 (16%) Moderate OSA: 33 (12%) Severe OSA: 24 (8%)	Increased BMI and neck circumference
Ghoshal et al, 2010, India ⁶¹	714 590 male 124 female	41 - 80	PSG	AHI > 5	604 (84.59%)	Has not been reported

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Godoy et al, 2022, Brazil ³⁹	62 11 male 51 female	73	PSG	Mild OSA: AHI \geq 5 and \leq 15 Moderate OSA: AHI > 15 and \leq 30 Severe OSA: AHI > 30	General: 45 Mild OSA: 17 Moderate OSA: 18 Severe OSA: 10	Has not been reported
Gregório et al, 2008, Brazil ⁴⁰	38 17 male 17 female	8.4 (4.0)	PSG	Mild OSA: 1-5 Moderate OSA: 5-10 Severe OSA: > 10	General: 35 (92%) Mild OSA: 16 (42.1%) Moderate OSA: 11 (28.9%) Severe OSA: 8 (22.1%)	Snoring, bruxism, nasal obstruction and restless legs during sleep. It is present in 75% of the examined patients with severe disorder.
Guscoth et al., 2021, Australia ⁵⁴	753 (all male)	60.8 (10.9)	PSG	Mild OSA: 10-19 Moderate OSA: 20-29 Severe OSA: \geq 30	General: 396 Mild OSA: 196 Moderate OSA: 107 Severe OSA: 93	OSA metrics were positively associated with serum triglyceride levels in men with a normal WC
Han et al, 2021, Korea ⁸³	103 (all male)	44.3 (8.1)	PSG	Moderate and severe OSA: > 15	73	BMI and cumulative flight time
Heraganahally et al, 2019, Australia ⁵⁵	3078 1972 male 1767 female	51.1	PSG	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: > 30	General: 2569 (83.7%) Mild OSA: (28.4%) Moderate: (22.3%) Severe: (33.0%)	Male, age and higher BMI. Among individuals with severe AHI, hypertension, heart disease and diabetes.
Heraganahally et al, 2021, Australia ⁵⁶	340 173 male 167 female	47 (38-57)	PSG	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: > 30	General: 297 Mild OSA: 77 Moderate OSA: 69 Severe OSA: 151	Has not been reported
Hu et al, 2019, China ⁴⁶	196 161 male 35 female	18 - 70	PSG	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: \geq 30	General: 168 Mild OSA: 28 Moderate OSA: 31 Severe OS: 109	Gender, BMI and neck circumference.
Ioachimescu et al, 2020, USA ²⁸	500 400 male 100 female	52.5 (41.8–62.5)	PSG	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: \geq 30	General: 423 Mild OSA: 133 Moderate OSA: 135 Severe OSA: 155	Has not been reported

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Ioannidou et al, 2021, Greece ¹⁰⁰	3791 2678 male 1113 female	57.2 (13.6)	PSG	Mild OSA: 5-15 Moderate OSA: 15-30 Severe OSA: ≥ 30	General: 3271 Mild OSA: 534 Moderate OSA: 918 Severe OSA: 1819	Age, BMI, the number of cigarettes/day, pack/ Years, and Fagerstrom Test for Nicotine Dependence were higher in patients with more severe OSA with more prevalent CVD comorbidities
Jalilolghadr et al, 2022, Iran ⁹⁷	112 73 male 39 female	5.8	PSG	Mild OSA: 1 - 4 Moderate OSA: 5 - 10 Severe OSA: > 10	General: 99 Mild OSA: 14 Moderate OSA: 19 Severe OSA: 66	Has not been reported
Jia et al, 2021, China ⁴⁷	107 63 male 44 female	31.7 (11.9)	PSG	Mild OSA: 5-15 Moderate OSA: 15 - 30 Severe OSA: ≥ 30	General: 63 Mild OSA: 25 Moderate OSA: 20 Severe OSA: 18	Increased risk for EPVS in OSA is a potential contributor to increased stroke risk in OSA
Jung et al, 2017, USA ²⁹	352 194 male 158 female	58	PSG	Mild OSA: 5-14 Moderate OSA: ≥ 15	General: 267 Mild OSA: 126 Moderate OSA: 141	Male sex, age, BMI, and presence of hypertension.
Kanda et al, 2018, Brazil ⁴¹	255 137 male 118 female	55.7 (10.6)	PSG	AHI ≥ 15	91	Female, older and with higher BMI
Katz et al, 2014, Canada ⁸⁹	222 121 male 101 female	12.1 (7.0 - 17.9)	PSG	AHI > 5 or OAI ≥ 1	47	-
Kim et al, 2004, Korea ⁸⁴	457 309 male 148 female	Male with snoring: 49.1 Male without snoring: 48.4 Female with snoring: 54.3 Female without snoring: 48.8	PSG	AHI ≥ 5	160	WHR, BMI, and systolic blood pressure men aged 50 to 59 years women increased with age
Klawe et al, 2005, Poland ¹⁰⁸	G1: 21 fast-rotating shift worked police officers G2: 21 subjects age-matched (control group)	G1: 47.1 (3.2) G2: 42.0 (2.8)	PSG	AHI ≥ 5	Police officers: 8 Control volunteers: 8	Overweight or obesity
Kobayashi et al, 2013, Japan ⁹³	152	6-12	OSA Screening (152) and PSG (8)	AHI ≥ 5	6	Has not been reported

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Koyama et al, 2012, Brazil ⁴²	745 All male	35.6 (9.6)	PSG	Mild OSA: 5–15; Moderate OSA: 15–30; Severe OSA: >30	General: 261 Mild OSA: 405 Moderate OSA: 188 Severe OSA: 151	Obese, older, time employed as a shift worker, and had consumed more alcohol.
Lee et al, 2019, Korea ⁸⁵	105 56 male 59 female	40.6 (11.6)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: ≥ 30	General: 37 Mild OSA: 22 Moderate OSA: 9 Severe OSA: 6	Older, BMI and predominant with male.
Li et al, 2014, China ⁴⁸	350 302 males 48 females	42.9 (11.0)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: ≥ 30	General: 315 Mild OSA: 44 Moderate OSA: 57 Severe OSA: 214	Age, BMI, likelihood and hypertension.
Liu et al, 2020, China ⁴⁹	4297 3416 male 881 female	Male: 40 Female: 48 (non-OSA); 50 (OSA)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: ≥ 30	General: 3679 Mild OSA: 713 Moderate OSA: 716 Severe OSA: 2250	BMI, neck circumference, WC, hip circumference and waist-to-hip ratio
McArdle et al, 2020, Australia ⁵⁷	935 459 male 476 female	22.2	PSG	Mild-to-severe OSA: ≥ 5 Moderate-to-severe OSA: ≥ 15;	General: 192 Moderate-severe OSA: 34	Has not been reported
Morell-Garcia et al, 2021, Spain ⁷⁷	168 Children -50 male -34 female Adults: has not been reported	Children: 8.1 Adults: has not been reported	PSG	Children -Mild OSA: AHI ≥ 1 and ≤ 5 -Moderate and severe OSA: AHI ≥ 5 Adults AHI ≥ 5	Children -Mild OSA: 28 -Moderate and severe OSA: 18 Adults: 51	The prevalence of fathers with OSA increases with the children's severity in the group of children with moderate-severe OSA. Having moderate-severe pediatric OSA were more than 4 times higher among children with a father with OSA
Mosharraf-Hossain et al, 2015, Bangladesh ¹¹⁸	2250	39.8 (9.4)	OSA Screening (2250) and PSG (120)	OSAH: AHI >5 and OSASH: AHI >5 + excessive daytime sleepiness	OSAH: -Snoring sample: 29 -non-snoring sample: 1 OSASH: -Snoring sample: 9 -non-snoring sample: 0	Male gender, age, obesity (defined by a high BMI), waist/hip ratio

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Mungo et al, 2021, Belgium ⁹⁶	264 131 male 133 female	18-30 years	PSG	OSAH: AHI > 5	49	Atypical depression, male, snoring, presence of at least one cardiometabolic alteration, mass index body weight greater than 30 kg/m ² and ferritin greater than or equal to 150 g/L
Mysliwiec et al, 2013, USA ³⁰	725 676 male 49 female	35.5 (8.6)	PSG	Mild OSA: 5-15 Moderate to severe OSA: > 15)	General OSA: 390 Mild OSA: 207 Moderate to severe OSA: 183	Older military personnel and males were more likely, an elevated BMI and the absence of an anxiety diagnosis were also associated with moderate-to-severe OSA
Neruntarat et al, 2010, Thailand ⁷⁴	2685	43.4 (9.3)	OSA screening (2685) and PSG (196- 102 snorers and 94 non-snorers)	AHI > 5	Snorers: 40 non-snorers: 2	Male gender, age >45 years, obesity defined by BMI of >25 kg/m ² , alcohol ingestion, smoking, and hypertension.
Pien et al, 2014, USA ¹⁰⁹	105 woman	26.7 (7.2)	PSG	General OSA: AHI ≥ 5 Mild OSA: 5-14 Moderate OSA: 15-29 Severe OSA: ≥ 30	First semester: 11 Third semester: 28 -Mild OSA: 23 -Moderate OSA: 4 -Severe OSA: 1	First trimester BMI and maternal age were significantly associated with third trimester OSA.
Powers et al, 2009, USA ³²	62 59 male 3 female	40.5 (7.8)	PSG	Mild or moderate OSA: AHI ≥ 5 and < 30	41	Has not been reported
Quintana-Gallego et al, 2004, Spain ⁷⁸	1745 1386 male 359 female	Male: 52.9 (11) Female: 57.8 (9.5)	PSG OR polygraphy	AHI > 10	General: 1166 -Male: 970 -Female: 196 Overnight PSG: 233 Nap PSG: 342 Polygraphy: 591	Has not been reported
Randby et al, 2013, Norway ¹⁰⁹	16,302 479 after screening 273 male 206 female	47.9	OSA Screening (16,3052) and PSG (479)	Mild to moderate OSA: 5 - 29.9 Severe OSA: ≥ 30	266	OSA was an independent predictor of impaired endothelial function in women
Ratnavadivel et al, 2009, Australia ⁵⁸	253 169 male 84 female	Non OSA: 47.5 (1.9) OSA: 54.1 (1.1)	PSG	AHI ≥ 15	171	This group was significantly older, comprised relatively more males, and had a higher BMI

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Reddy et al, 2009, India ⁶³	2505 1263 male 1242 female	41 (9)	Snorers screening (2505) and PSG (365- 287 snorers and 73 non-habitual snorers)	AHI ≥ 5	Habitual snorers: 94 Non-habitual snorers: 3	Increasing age, male gender, socioeconomic status score, habitual smoking, hypertension, obesity, high WHR, PPNC > 90, and high subscapular and suprailliac skinfold thicknesses Has not been reported
Reddy et al, 2014, India ⁶²	425 children 389 male 36 female	2 months - 18 years	PSG	Mild OSA: 1-5 Moderate OSA: 5-10 Severe OSA: ≥ 10	General OSA: 211 Mild OSA: 99 Moderate OSA: 34 Severe OSA: 78	Has not been reported
Roche et al, 2021, South Africa ¹¹⁰	75 22 male 53 male	66.1 (10.7)	PSG	AHI ≥ 15	22	Participants were older, had higher BMI, higher WC, higher prevalence of Mets and greater CMR score
Sadeghniaat-Haghighi et al, 2015, Iran ⁹⁸	603 451 male 152 female	45.8 (12.7)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: ≥ 30	General OSA: 438 Mild OSA: 124 Moderate OSA: 114 Severe OSA: 201	Has not been reported
Sasai-Sakuma et al, 2016, Japan ⁹⁴	2389 male	48.2 (7.6)	OSA Screening (2389) and PSG (422)	AHI ≥ 15 or AHI $\geq 5 + ESS \geq 11$	General: 235 AHI ≥ 15 : 224 AHI $\geq 5 + ESS \geq 11$: 11	Has not been reported
Scarlata et al, 2013, Italy ¹⁰²	254 154 male 100 female	65.8 (12.1)	Cardio-respiratory polygraphy (219) or PSG (35)	Mild OSA: 10 - 20 Moderate OSA: 20-30 Severe OSA: ≥ 30	General OSA: 141 Mild OSA: 46 Moderate 35 Severe 60	Has not been reported
Selvadurai et al, 2019, Canada ⁹⁰	113 children 75 male 38 female	No-OSA: 2.2 (0.6) OSA: 2.2 (0.5)	Overnight PSG	Mild OSA: OAHl ≥ 1.5 to <5 Moderate OSA: OAHl ≥ 5 to <10 Severe OSA: OAHl ≥ 10	General: 66 Mild OSA: 13 Moderate-to-severe OSA: 53	Overweight and obese individuals
Sharma et al, 2006, India ⁶⁴	-	42.7 (9.0)	OSA Screening (2150) and PSG (151)	AHI > 5	General: 38 Habitual snorers: 36 Non habitual snorers: 2	Male gender, age 45 years, obesity defined by a BMI of 25 kg/m ² , and WHR of 0.80 in women and 0.95 in men
Singh et al, 2017, India ⁶⁵	1512 1026 male 485 female	42.6 (11.2)	OSA Screening (1512) and PSG (311)	AHI ≥ 5	General: 62 OSA high-risk: 44 OSA low-risk: 18	Gender (males)

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Sogut et al, 2005, Turkey ⁷¹	1198 582 male 616 female	8.1 (1.9)	OSA Screening (1198) and PSG for snoring children (28)	AHI > 3	11	Has not been reported
Tapia et al, 2016, USA ⁵⁹	197 114 male 83 female	5–12	PSG	AHI ≥ 2	19	OSAS is associated with chorioamnionitis and multiple gestation in ex-preterm children.
Tawaranurak et al, 2019, Thailand ⁷⁵	929 622 male 307 female	18 - 85	PSG	Mild OSA: AHI ≥ 5 and < 15 Moderate OSA: AHI ≥ 15 and ≤ 30 Severe OSA: AHI > 30	General OSA: 795 Mild OSA: 215 Moderate OSA: 167 Severe OSA: 413	Male gender, age ≥ 50 years, BMI > 25 kg/m ² , neck circumference > 40 cm and waist to height ratio > 0.6
Togei et al, 2013, Brazil ⁴³	1,042	Non-OSA: 37.3 (12.3) Mild OSA: 48.3 (13.6) Moderate-to-severe OSA: 53.3 (13.3)	PSG	Mild OSA: AHI 5 - 14.9 Moderate to severe OSA: AHI ≥ 15	General: 396 Mild OSA: 221 Moderate-to-severe: 175	Older, more obese, and had greater mean values for waist and neck circumferences as well as for lipid levels
Tripathi et al, 2017, India ⁶⁶	552	35 - 55	OSA Screening (552) and PSG (120)	AHI > 5	57	Lifestyle stress, sleep deprivation, and upper airway inflammation
Tufik et al, 2010, Brazil ⁸	104 469 male 573 female	42 (14)	PSG	AHI ≥ 5 + at least one of the following complaints: loud snoring, daytime sleepiness, fatigue, and breathing interruptions during sleep. OR AHI ≥ 15	General OSA: 398 Mild OSA: 222 Moderate-to-severe OSA: 176	Has not been reported
Urschitz et al, 2010, Germany ⁸⁷	1144 583 male 561 female	9.6 (0.7)	OSA Screening (1144) and Home PSG (183)	AHI ≥ 1	OSA high-risk: 20 OSA low-risk: 2	BMI, history of allergy, a composite questionnaire score, and two oximetry-based criteria.
Valenza et al, 2012, Spain ⁷⁹	1008 804 male 205 female	18 - 80 years	PSG	AHI ≥ 5	893	Hypertension, diabetes and cardiac pathology
Veeravigrom et al, 2016, Thailand ⁷⁶	166 111 male 55 female	8.6 (4.6)	PSG	Mild OSA: 1 - 5 Moderate OSA: 5 - 10 Severe OSA: ≥ 10	General: 154 Mild OSA: 43 Moderate OSA: 44 Severe OSA: 67	Has not been reported

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Vega-Osorio et al, 2020, Colombia ¹¹¹	575 273 male 302 female	51.0 (13.7)	PSG	Mild OSA: 5 - 14 Moderate OSA: 15 - 29 Severe OSA: ≥ 30	General: 420 Mild OSA: 142 Moderate OSA: 169 Severe OSA: 109	Has not been reported
Wahida et al, 2013, Malaysia ¹¹²	289	45.4 (7.0)	PSG	Mild OSA: AHI ≥ 5 and < 15 Moderate OSA: AHI ≥ 15 and < 30 Severe OSA: AHI ≥ 30	General: 128 Mild OSA: 83 Moderate OSA: 26 Severe OSA: 19	Age, BMI, neck circumference and hypertension
Wahner-Roedler et al, 2007, USA ³³	522 267 male 255 female	57 years (range, 18-76 years)	PSG	AHI ≥ 15 or AHI ≥ 5 and < 15 + insomnia, excessive daytime sleepiness, mood disorder, impaired cognition, hypertension, ischemic heart disease, or history of stroke.	406	Has not been reported
Wali et al, 2017, Saudi Arabia ¹¹³	2682 1286 male 1396 females	40.7 (6.5)	Snoring screening (2682) and PSG (375)	AHI ≥ 5	235	Male gender, age ≥ 50 years, obesity as defined by a BMI ≥ 30 kg/m ² , and a history of hypertension
Wiebracht et al, 2017, USA ³⁴	342 182 male 160 female	11.3 (2.4)	PSG	Mild OSA: AHI ≥ 1 and < 5 Moderate OSA: AHI ≥ 5 and < 10 Severe OSA: AHI ≥ 10	General OSA: 170 Mild OSA: 109 Moderate OSA: 34 Severe OSA: 27	Has not been reported
Xie et al, 2011, EUA ³⁵	1890 1731 male 159 female	43.7 (11.5)	OSA screening (1833) and PSG (65)	Mild OSA: AHI ≥ 5 and ≤ 15 Moderate OSA: AHI < 15 and < 30 Severe OSA: ≥ 30	General OSA: 51 Mild OSA: 22 Moderate OSA: 20 Severe OSA: 31	Higher BMI, higher neck circumference, and lower minimum oxygen saturation
Xu et al, 2006, China ⁵¹	50	7.8 (3.2)	PSG	AHI > 5	31	Observable apnea during sleep, nocturnal enuresis, intrusive naps, mouth breathing, enlarged tonsils, and radiologic features of upper airway narrowing due to adenoid hypertrophy

Table 1 (continued)

Author, years, country	Sample size and type (male; female)	Mean (SD) age or range	Diagnostic methods	OSA or OSAS definition	Results number	Factor risk
Xu et al, 2020, China ⁵⁰	1578 1060 male 518 female	OSAS children: 5.7 non-OSAS children: 6.0	PSG	AHI ≥ 1	1008	Snoring ≥ 3 months, male gender, preterm birth, breastfeeding, obesity, Neck circumference ≥ 30 cm, Neck /height ratio ≥ 0.25 , tonsillar and adenoid hypertrophy.
Zacharias et al, 2021, Germany ⁸⁸	529 247 male 282 female	52.2	PSG	Mild OSA: 5 - 15 Moderate OSA: 15 - 30 Severe OSA: ≥ 30	General OSA: 209 Mild OSA: 125 Moderate OSA: 52 Severe OSA: 32	Has not been reported
Zamarrón et al, 2006, Spain ⁸¹	187 147 male 40 female	57.9 (12.8)	PSG	AHI ≥ 10	111	COPD

OSA, obstructive sleep apnea; OSAS, obstructive sleep apnea syndrome; PSG, polysomnography; AHI, apnea-hypopnea index; EDS, excessive daytime sleepiness; IHD, ischemic heart disease; COPD, chronic obstructive pulmonary disease; OI, obstructive apnea-hypopnea index; BMI, body mass index; OHA1, obstructive hypopnea apnea index; CVD, cardiovascular disease; EPVS, enlarged perivascular space; WHR, waist-hip ratio; PPNC, percent predicted neck circumference; WC, waist circumference; CMR, cardiovascular magnetic resonance

of Obstructive Sleep Apnea Syndrome (OSAS) in the country's population. They discovered that the male gender and obesity pose a greater risk than the female gender in the same age group. However, age is the most important predictor in women, with postmenopausal women being at a higher risk [6]. In this review, no single risk factor identified in the included articles explained the heterogeneity in the meta-regression analysis, indicating the possibility of multifactorial risk influencing this outcome.

The BMI was the most commonly reported risk factor in the studies, followed by age and gender, and BMI was reported more frequently in men. Other factors found in fewer studies included hypertension, cigarette smoking, and alcohol consumption. These findings are consistent with the findings of other studies, which identified male gender, older age, and higher BMI as risk factors [4, 5, 7]. Several included studies showed increasing OSA prevalence with increasing age, being significant for countries with older or aging populations. Therefore, the predominance of BMI and the male gender should be considered, particularly when planning risk-reduction interventions. Positive associations were found between OSA and waist-to-hip ratio, larger neck circumference, hypertension, smoking, snoring, and sleepiness [5]. Additionally, ethnicity was considered a risk factor in three studies: one linked white race to a higher likelihood of OSA [19]; another found that African Americans had a higher likelihood of having severe OSA than white patients [22]; and the third found that African patients had a higher likelihood of having a higher AHI [92].

Ninety-eight of the 99 articles chosen were eligible for meta-analysis and were divided into subgroups to increase the reliability of the analysis. When the age group was considered in the subgroup analysis, the prevalence of OSA in adults was 57%. However, the prevalence was higher in the elderly subgroup, at 69%, with the lowest prevalence in the subgroup of samples made up entirely of children, with a 50% estimate. A study of Chinese children [119] found a prevalence of OSAS ranging from 4.8 to 40.3%, depending on the cut-off point used, with male gender, obesity, and amygdala size associated with OSAS [119].

Regarding the classification of OSA, a subgroup analysis was performed concerning the severity of diagnosed OSA using the following classifications: "mild", "mixed", "severe", "mild to moderate", "moderate to severe", and "not classified". The classifications with the highest prevalence occurred in "mild to moderate" (69%), followed by "mixed" (68%), and with the lowest prevalence in "severe" (14%). Another subgroup pertains to the included countries, in which we found that 34 countries had at least one study that sought to

Table 2 Estimates for each subgroup

<i>Variant</i>	Sample number	<i>Prevalence by subgroup</i>	<i>IC95%</i>	<i>I</i> ²
Age group (mean age)	47634	0.56	[0.51; 0.61]	99%
Children and teenagers (2.2–11.3)	4881	0.51	[0.38; 0.64]	98%
Adults (22.2–66.1)	41797	0.57	[0.51; 0.62]	99%
Elderly (71.8–73)	242	0.69	[0.63; 0.74]	0%,
Adults and elderly (not subdivided)	714	0.85	[0.82; 0.87]	Not applied
PSG	47370	0.57	[0.51; 0.62]	99%
Triage	6413	0.52	[0.41; 0.63]	99%
Total PSG	40957	0.58	[0.52; 0.64]	99%
Type	47634	0.56	[0.51; 0.61]	99%
Syndrome	13242	0.52	[0.42; 0.61]	99%
OSA	34392	0.58	[0.52; 0.64]	99%
Country	47634	0.56	[0.51; 0.61]	99%
Egypt	42	0.12	[0.04; 0.26]	
USA	6156	0.56	[0.44; 0.68]	99%
Turkey	473	0.60	[0.40; 0.78]	95%
Oman	608	0.49	[0.45; 0.53]	
Mexico	51	0.43	[0.29; 0.58]	
Australia	5649	0.54	[0.30; 0.78]	100%
Belgium	2357	0.28	[0.16; 0.41]	97%
Brazil	4460	0.55	[0.36; 0.73]	99%
China	6733	0.65	[0.46; 0.81]	99%
Korea	1411	0.37	[0.15; 0.63]	99%
Thailand	1314	0.67	[0.22; 0.98]	99%
Lebanon	663	0.79	[0.76; 0.82]	
Mongolia	205	0.93	[0.89; 0.96]	
Ireland	250	0.54	[0.48; 0.61]	
Non-reported	160	0.78	[0.49; 0.97]	87%
Ecuador	180	0.67	[0.60; 0.74]	
Slovakia	42	0.50	[0.34; 0.66]	
Greece	4063	0.82	[0.70; 0.91]	95%
Spain	3498	0.64	[0.51; 0.76]	98%
Vietnam	93	0.61	[0.51; 0.71]	
Germany	1869	0.33	[0.16; 0.53]	97%
Portugal	318	0.64	[0.58; 0.69]	
Sweden	399	0.51	[0.46; 0.56]	
Italy	370	0.73	[0.38; 0.96]	98%
India	1987	0.59	[0.29; 0.87]	100%
Iran	715	0.81	[0.64; 0.93]	93%
Canada	335	0.39	[0.08; 0.76]	98%
Japan	430	0.57	[0.50; 0.63]	2%
Poland	42	0.38	[0.24; 0.54]	0%
Bangladesh	60	0.50	[0.37; 0.63]	
Norway	479	0.56	[0.51; 0.60]	
Singapore	425	0.50	[0.45; 0.55]	
Africa	75	0.29	[0.19; 0.41]	
Taiwan	166	0.93	[0.88; 0.96]	
Colombia	575	0.73	[0.69; 0.77]	
Malaysia	289	0.44	[0.38; 0.50]	
Saudi Arabia	692	0.40	[0.17; 0.66]	97%
OSA classification	47370	0.57	[0.51; 0.62]	99%

Table 2 (continued)

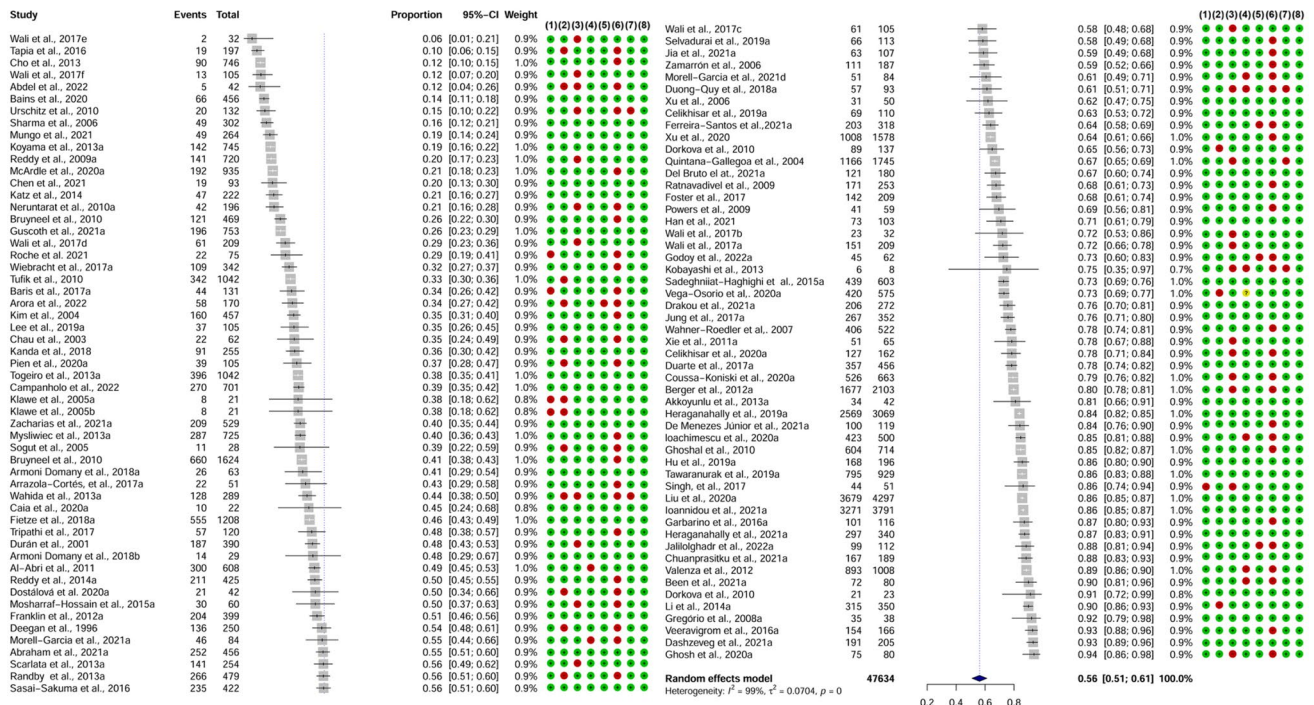
<i>Variant</i>	Sample number	<i>Prevalence by subgroup</i>	<i>IC95%</i>	<i>I²</i>
Mild	725	0.40	[0.36; 0.43]	Not applied
Mild to moderate	59	0.69	[0.57; 0.81]	Not applied
Moderate to severe	1019	0.37	[0.08; 0.73]	99%
Severe	465	0.14	[0.11; 0.18]	Not applied
Mixed	27041	0.68	[0.61; 0.73]	99%
No classification	18070	0.46	[0.39; 0.54]	99%
Sample origin	47634	0.56	[0.51; 0.61]	99%
Drivers	2145	0.45	[0.00; 0.99]	99%
PSG Lab	31460	0.65	[0.59; 0.71]	99%
Long distance drivers	42	0.81	[0.66; 0.91]	
Military personal (PSG Lab)	954	0.47	[0.27; 0.68]	91%
Rugby league athletes	22	0.45	[0.24; 0.68]	
General population	9290	0.40	[0.31; 0.50]	98%
Heavy equipment users	110	0.63	[0.53; 0.72]	
Bus drivers	162	0.78	[0.71; 0.84]	
Truck drivers	116	0.87	[0.80; 0.93]	
Airplane drivers	103	0.71	[0.61; 0.79]	
Unclear	107	0.59	[0.49; 0.68]	
Policemen	21	0.38	[0.18; 0.62]	
Age matched group	21	0.38	[0.18; 0.62]	
Railroad workers	745	0.19	[0.16; 0.22]	
Pregnant women	1040	0.28	[0.13; 0.45]	92%
Shift workers (iron ore mining)	119	0.84	[0.76; 0.90]	
Parents of children	84	0.61	[0.49; 0.71]	
Public transport drivers	422	0.56	[0.51; 0.60]	

investigate the relationship between OSA and their population. Among all the continents, Asia had the most countries, with 13, followed by Europe, with 11. However, the continents with the most studies performed were Asia, with 35, followed by America, with 29. These numbers emphasize the importance of conducting additional studies in Africa and Oceania, which total three countries and nine studies aimed at understanding the relationship between this population and OSA.

The most significant strengths of this systematic review are related to the fact that the included studies were global in scope, with no language restrictions and no age restrictions. In addition, only studies that used PSG to diagnose OSA were considered. A thorough search of six databases and the grey literature was also conducted. Nonetheless, it is worth noting that the GRADE system assigned a very low level of certainty of evidence. The explanation stems from the studies' risk of bias assessment, which revealed that eight had a moderate risk of bias. In addition, these studies lacked clarity regarding the criteria used for OSA diagnosis and outcomes and uncontrolled confounding factors.

Regarding diagnosis criteria, some studies used a screening questionnaire on the study population before selecting a convenience sample and performing the PSG on them alone. Furthermore, other studies were based on a specific population with a convenience sample, which may have biased the estimate. Due to their morbidity profiles and other phenomena, such as the work effect, this convenience sample of drivers, pilots, athletes, military personnel, pregnant women, and others did not represent the general population. Subgroup analyses were performed to stratify categorical confounders that could overestimate the estimates generated in some population groups.

The study was also limited by the PSG conducted. Sleep studies are classified as type I, type II, type III, or type IV, with type I currently being considered the standard reference diagnostic test for OSA because it is conducted in a sleep laboratory and observed by a technician, while the other studies are unattended [120]. Nonetheless, some studies that used PSG did not report the type of sleep study used or conducted the other types, which may have acted as a confounding factor for the



Joanna Briggs Institute Critical Assessment Checklist (Cross-sectional Studies)

1. Were the inclusion criteria in the sample clearly defined?
2. Were the study subjects and the scenario described in detail?
3. Was the exposure measured in a valid and reliable way?
4. Were objective and standard criteria used to measure the condition?
5. Were confounding factors identified?
6. Were strategies for deal with confounding factors stated?
7. Were the outcomes measured in a valid and reliable way?
8. Was appropriate statistical analysis used?

Judgement
 ● High
 ● Unclear
 ● Low

Fig. 3 Forest plot of the meta-analysis of the prevalence of OSA displaying risk-of-bias judgements for each study included

estimates generated, given that type I has a higher cost than the others.

The importance of the current study lies in raising awareness of the high prevalence of OSA in the global population and across age groups, highlighting the need for public initiatives for discovery and for effective treatment. Continuous positive airway pressure (CPAP) is the standard treatment for OSA [121]. CPAP provides adequate upper airway stability through pressurization. Benefits may include improvement in cardiovascular comorbidities, decrease in excessive daytime sleepiness, and improvement in cognitive function and quality of life [121].

Additional treatment options can be used alone or combined, considering the condition's multifactorial etiology. These include using a mandibular advancement device [121], performing oropharyngeal exercises, and engaging in functional exercises that involve breathing, chewing, swallowing, and speaking [122]. However, studies on the long-term effects of speech-language therapy combined with functional and oropharyngeal exercises are still lacking. In contrast, CPAP

use and the mandibular advancement device have been shown to successfully treat OSA in follow-ups [123–125].

Conclusion

Approximately half of the people have OSA worldwide. Although the literature cites high BMI, increased age, and male gender as risk factors, these covariates did not influence the heterogeneity existing by meta-regression. Based upon the level of certainty, the results of this investigation should be with caution. We suggest studies with better methodological qualities.

Other information

Registration

The PROSPERO (International Prospective Registry of Systematic Reviews - Center for Reviews and Dissemination University of York-CRD42021265627) website has the protocol for this systematic review listed.

13. Moola S, Munn Z, Tufanaru C, Aromataris E, Sears K, Sfetcu R, et al. Chapter 7: Systematic reviews of etiology and risk . In: Aromataris E, Munn Z (Editors). JBI Manual for Evidence Synthesis JBI 2020.
14. Campos LGN, Pedrosa BH, Cavalcanti RVA, Stechman-Neto J, Gadotti IC, de Araujo CM et al (2021) Prevalence of temporomandibular disorders in musicians: a systematic review and meta-analysis. *J Oral Rehabil* 48:632–42. <https://doi.org/10.1111/joor.13150>
15. Dersimonian R, Laird N (1986) Meta-Analysis in Clinical Trials. *Control Clin Trials* 7:177–88
16. Higgins JPT, Thompson SG (2002) Quantifying heterogeneity in a meta-analysis. *Stat Med* 21:1539–58. <https://doi.org/10.1002/sim.1186>
17. Barendregt JJ, Doi SA, Lee YY (2013) Meta-analysis of prevalence. *J Epidemiol Community Health* 67:974–8. <https://doi.org/10.1136/jech>
18. Schünemann H, Brożek J, Guyatt G, Oxman A (2013) Handbook for grading the quality of evidence and the strength of recommendations using the GRADE. Available from <https://gdt.gradepro.org/app/handbook/handbook.html#h.svwngs6pm0f2>
19. Abraham EJ, Bains A, Rubin BR, Cohen MB, Levi JR (2021) Predictors of a normal sleep study in healthy children with sleep disordered breathing symptoms. *Annals of Otolaryngology, Rhinology and Laryngology* 130:1029–35. <https://doi.org/10.1177/0003489421990156>
20. Domany KA, Hossain MM, Nava-Guerra L, Khoo MC, McConnell K, Carroll JL et al (2018) Cardioventilatory control in preterm-born children and the risk of obstructive sleep apnea. *Am J Respir Crit Care Med* 197:1596–603. <https://doi.org/10.1164/rccm.201708-1700OC>
21. Arora SK, Powell TA, Foster SN, Hansen SL, Morris MJ (2022) Exercise capacity remains supernormal, though mildly reduced in middle-aged military personnel with moderate to severe obstructive sleep apnea. *Sleep and Breathing*. <https://doi.org/10.1007/s11325-022-02587-1>
22. Bains A, Abraham E, Hsieh A, Rubin BR, Levi JR, Cohen MB (2020) Characteristics and frequency of children with severe obstructive sleep apnea undergoing elective polysomnography. *Otolaryngology - Head and Neck Surgery (United States)* 163:1055–60. <https://doi.org/10.1177/0194599820931084>
23. Berger M, Varvarigou V, Rielly A, Czeisler CA, Malhotra A, Kales SN (2012) Employer-mandated sleep apnea screening and diagnosis in commercial drivers. *J Occup Environ Med* 54:1017–25. <https://doi.org/10.1097/JOM.0b013e3182572e16>
24. Foster SN, Hansen SL, Capener DC, Matsangas P, Mysliwiec V (2017) Gender differences in sleep disorders in the US military. *Sleep Health* 3:336–41. <https://doi.org/10.1016/j.sleh.2017.07.015>
25. Ioachimescu OC, Shirine Allam J, Samarghandi A, Anand N, Fields BG, Dholakia SA et al (2020) Performance of peripheral arterial tonometry–based testing for the diagnosis of obstructive sleep apnea in a large sleep clinic cohort. *J Clin Sleep Med* 6:1663–74. <https://doi.org/10.5664/jcsm.8620>
26. Jung Y, Junna MR, Mandrekar JN, Morgenthaler TI (2017) The National Healthy Sleep Awareness Project Sleep Health Surveillance Questionnaire as an Obstructive Sleep Apnea Surveillance Tool. *J Clin Sleep Med* 13:1067–74. <https://doi.org/10.5664/jcsm.6724>
27. Mysliwiec V, McGraw L, Pierce R, Smith P, Trapp B, Roth BJ (2013) Sleep disorders and associated medical comorbidities in active duty military personnel. *Sleep* 36:167–74. <https://doi.org/10.5665/sleep.2364>
28. Pien GW, Pack AI, Jackson N, Maislin G, Macones GA, Schwab RJ (2014) Risk factors for sleep-disordered breathing in pregnancy. *Thorax* 69:371–7. <https://doi.org/10.1136/thoraxjnl-2012-202718>
29. Powers CR, Frey WC (2009) Maintenance of wakefulness test in military personnel with upper airway resistance syndrome and mild to moderate obstructive sleep apnea. *Sleep and Breathing* 13:253–8. <https://doi.org/10.1007/s11325-009-0245-7>
30. Wahner-Roedler DL, Olson EJ, Loehrler LL, Sood A (2007) Gender-specific differences in a patient population with obstructive sleep apnea-hypopnea syndrome. *Gend Med* 4:329–38
31. Wiebracht ND, He S, Cotton C, Meinzen-Derr J, Shott G, Smith DF et al (2017) Polysomnographic oxygen saturation findings for preteen children versus adolescents. *Otolaryngology - Head and Neck Surgery (United States)* 158:187–93. <https://doi.org/10.1177/0194599817733687>
32. Xie W, Chakrabarty S, Levine R, Johnson R, Talmage JB (2011) Factors associated with obstructive sleep apnea among commercial motor vehicle drivers. *J Occup Environ Med* 53:169–73. <https://doi.org/10.1097/JOM.0b013e3182068ceb>
33. Campanholo M de AT, Caparroz F de A, Vidigal T de A, Kenchian CH, Andersen ML, Tufik S, et al. Assessment of laryngopharyngeal reflux and obstructive sleep apnea: a population-based study. *Laryngoscope* 2022. <https://doi.org/10.1002/lary.30061>.
34. de Menezes Júnior LAA, Fajardo VC, do Nascimento Neto RM, de Freitas SN, de Oliveira FLP, Pimenta FAP, et al. Diagnostic accuracy of the Berlin questionnaire and the NoSAS score in detecting risk for obstructive sleep apnea in rotating shift workers. *Sleep and Breathing* 2021. <https://doi.org/10.1007/s11325-021-02446-5>.
35. de Duarte RL, M, Fonseca LB de M, Magalhães-Da-Silveira FJ, da Silveira EA, Rabahi MF. (2017) Validação do questionário STOP-Bang para a identificação de apneia obstrutiva do sono em adultos no Brasil. *J Brasileiro de Pneumologia* 43:456–63. <https://doi.org/10.1590/s1806-37562017000000139>
36. Godoy PH, dos Santos Nucera APC, de Paiva Colcher A, de Andrade JE, da Silveira Barroso Alves D. (2022) Screening for obstructive sleep apnea in elderly: performance of the Berlin and STOP-Bang questionnaires and the Epworth Sleepiness Scale using polysomnography as gold standard. *Sleep Science* 15:203–8. <https://doi.org/10.5935/1984-0063.20220020>
37. Gregório PB, Athanazio RA, Bitencourt Almir Galvão Vieira, Neves FBCS, Terse R, Hora F. Symptoms of obstructive sleep apnea-hypopnea syndrome in children. *J Bras Pneumol*;34:356–61.
38. Kanda GM, Priore NC, Toledo C, Shimizu RN, Arata YP, Gonzaga C et al (2018) Perfil clínico e laboratorial de pacientes com e sem apneia obstrutiva do sono Clinical and laboratory profile of patients with or without obstructive sleep apnea Endereço para correspondência. *Rev Soc Bras Clin Med* 16:108–20
39. Koyama RG, Esteves AM, Oliveira e Silva L, Lira FS, Bittencourt LRA, Tufik S, et al (2012) Prevalence of and risk factors for obstructive sleep apnea syndrome in Brazilian railroad workers. *Sleep Med* 13:1028–32. <https://doi.org/10.1016/j.sleep.2012.06.017>
40. Togeiro SM, Carneiro G, Ribeiro Filho FF, Zanella MT, Santos-Silva R, Taddei JA, et al. Consequences of obstructive sleep apnea on metabolic profile: a population-based survey. *Obesity* 2013;21. <https://doi.org/10.1038/oby.2012.146>.
41. Chau KW, Ng KK, Kwok L, Chow Y, Ho JCS, Ng DKK. Clinical risk factors for obstructive sleep apnoea in children FHKAM (Paed) currently in private practice. vol. 44. 2003.
42. Chen Y, Metz JE, Gao H, Gao X. Association between obstructive sleep apnea and periodontitis in Chinese male adults: A cross-sectional study. *J Prosthet Dent* 2021.
43. Hu Y, yuan, Yu Y, Wang Z bin, Liu C, Cui Y hua, Xiao W min. (2019) Reliability and validity of simplified

- Chinese STOP-BANG Questionnaire in diagnosing and screening obstructive sleep apnea hypopnea syndrome. *Curr Med Sci* 39:127–33. <https://doi.org/10.1007/s11596-019-2010-x>
44. Jia Y, Liu C, Li H, Li X, Wu J, Zhao Y et al (2021) Enlarged perivascular space and its correlation with polysomnography indicators of obstructive sleep apnea. *Nat Sci Sleep* 13:863–72. <https://doi.org/10.2147/NSS.S305465>
 45. Li Z, Du L, Li Y, Huang L, Lei F, Yang L et al (2014) Characterization of primary symptoms leading to Chinese patients presenting at hospital with suspected obstructive sleep apnea. *J Thorac Dis* 6:444–51. <https://doi.org/10.3978/j.issn.2072-1439.2014.02.08>
 46. Liu Y, Huang W, Zou J, Xu H, Qian Y, Zhu H et al (2020) Sea level nocturnal minimal oxygen saturation can accurately detect the presence of obstructive sleep apnea in a population with high pretest probability. *Sleep and Breathing* 25:171–9. <https://doi.org/10.1007/s11325-020-02014-3>
 47. Xu Z, Wu Y, Tai J, Feng G, Ge W, Zheng L, et al. Risk factors of obstructive sleep apnea syndrome in children. *Journal of Otolaryngology - Head and Neck Surgery* 2020;49. <https://doi.org/10.1186/s40463-020-0404-1>.
 48. Xu Z, Cheuk DKL, Lee SL (2006) Clinical evaluation in predicting childhood obstructive sleep apnea. *Chest* 130:1765–71. <https://doi.org/10.1378/chest.130.6.1765>
 49. Benn E, Wirth H, Short T, Howarth T, Heraganahally SS (2021) The Top End Sleepiness Scale (TESS): a new tool to assess subjective daytime sleepiness among indigenous Australian adults. *Nat Sci Sleep* 13:315–28
 50. Caia J, Halson SL, Scott A, Kelly VG (2020) Obstructive sleep apnea in professional rugby league athletes: an exploratory study. *J Sci Med Sport* 23:1011–5. <https://doi.org/10.1016/j.jsams.2020.04.014>
 51. Guscoth LB, Appleton SL, Martin SA, Adams RJ, Melaku YA, Wittert GA (2021) The association of obstructive sleep apnea and nocturnal hypoxemia with lipid profiles in a population-based study of community-dwelling Australian men. *Nat Sci Sleep* 13:1771–82. <https://doi.org/10.2147/NSS.S327478>
 52. Heraganahally SS, Kravavt A, Oguoma VM, Gokula C, Mehra S, Judge D, et al. Sleep apnoea among Australian Aboriginal and non-Aboriginal patients in the Northern Territory of Australia—a comparative study. *Sleep* 2019;43. <https://doi.org/10.1093/sleep/zsz248/5586811>.
 53. Heraganahally SS, Rajaratnam B, Silva SAAS, Robinson N, Oguoma VM, Naing P et al (2021) Obstructive sleep apnoea and cardiac disease among aboriginal patients in the Northern Territory of Australia. *Heart Lung Circ* 30:1184–92. <https://doi.org/10.1016/j.hlc.2021.01.007>
 54. McArdle N, Ward S v, Bucks RS, Maddison K, Smith A, Huang R-C, et al. The prevalence of common sleep disorders in young adults: a descriptive population-based study. *Sleep* 2020;43. <https://doi.org/10.1093/sleep/zsaa072/5819386>.
 55. Ratnavadivel R, Chau N, Stadler D, Yeo A, McEvoy RD, Catchside PG (2009) Marked reduction in obstructive sleep apnea severity in slow wave sleep. *J Clin Sleep Med* 5:519–24
 56. Tapia IE, Shults J, Doyle LW, Nixon GM, Cielo CM, Traylor J et al (2016) Perinatal risk factors associated with the obstructive sleep apnea syndrome in school-aged children born preterm. *Sleep* 39:737–42. <https://doi.org/10.5665/sleep.5618>
 57. Ghosh P, Sapna Varma NK, Ajith VV, Prabha RD, Raj M (2020) Epidemiological study on prevalent risk factors and craniofacial skeletal patterns in obstructive sleep apnea among South Indian population. *Indian J Dent Res* 31:784–90
 58. Ghoshal AG, Sarkar S, Roy DJ, Das RK, Ray M. Polysomnographic profile in a sleep laboratory in Kolkata: a retrospective analysis of 714 cases. *J Assoc Physicians India* 2010;58.
 59. Reddy KR, Lim MT, Lee TJ, Goh DY, Ramamurthy MB (2014) Pediatric polysomnographic studies at a tertiary-care hospital in Singapore. *Indian Pediatr* 51:484–6
 60. Reddy EV, Kadiravan T, Mishra HK, Sreenivas V, Handa KK, Sinha S et al (2009) Prevalence and risk factors of obstructive sleep apnea among middle-aged urban Indians: a community-based study. *Sleep Med* 10:913–8. <https://doi.org/10.1016/j.sleep.2008.08.011>
 61. Sharma SK, Kumpawat S, Banga A, Goel A (2006) Prevalence and risk factors of obstructive sleep apnea syndrome in a population of Delhi. *India. Chest* 130:149–56. <https://doi.org/10.1378/chest.130.1.149>
 62. Singh A, Prasad R, Garg R, Kant S, Hosmane GB, Dubey A et al (2017) A study to estimate prevalence and risk factors of obstructive sleep apnea syndrome in a semi-urban Indian population. *Monaldi Archives for Chest Disease* 87:40–8. <https://doi.org/10.4081/monaldi.2017.773>
 63. Tripathi A, Bagchi S, Singh J, Pandey P, Tripathi S, Gupta NK (2017) Lifestyle and occupational stress: a potential risk factor for obstructive sleep apnea in nonobese male subjects. *J Prosthodont* 27:716–21. <https://doi.org/10.1111/jopr.12627>
 64. Akkoyunlu ME, Altın R, Kart L, Atalay F, Örnek T, Bayram M, et al. Investigation of obstructive sleep apnoea syndrome prevalence among long-distance drivers from Zonguldak, Turkey. *Multidisciplinary Respiratory Medicine* 2013;8.
 65. Barış HE, Gökdemir Y, Eralp EE, Baş İkiçoğlu N, Karakoç F, Karadağ B et al (2017) Clinical and polysomnographic features of children evaluated with polysomnography in pediatric sleep laboratory. *Turk Pediatri Ars* 52:23–9. <https://doi.org/10.5152/TurkPediatriArs.2017.4218>
 66. Celikhisar H, Ilkhan GD. The association of obstructive sleep apnea syndrome and accident risk in heavy equipment operators. *Medicina (Lithuania)* 2019;55. <https://doi.org/10.3390/medicina55090599>.
 67. Celikhisar H, Ilkhan GD. 2020 Association of presence and severity of obstructive sleep apnoea syndrome with accident risk in city bus drivers. *J Pak Med Assoc* 70:2184–9. <https://doi.org/10.47391/JPMA.435>.
 68. Sogut A, Altın R, Uzun L, Ugur MB, Tomac N, Acun C et al (2005) Prevalence of obstructive sleep apnea syndrome and associated symptoms in 3–11-year-old Turkish children. *Pediatr Pulmonol* 39:251–6. <https://doi.org/10.1002/ppul.20179>
 69. Anuntaseree W, Kuasirikul S, Suntornlohanakul S (2005) Natural history of snoring and obstructive sleep apnea in Thai school-age children. *Pediatr Pulmonol* 39:415–20. <https://doi.org/10.1002/ppul.20207>
 70. Chuanprasitkul C, Veeravigrom M, Sunkonkit K, Tansrirattana-wong S, Sritippayawan S (2021) Incidence / predictors of pediatric obstructive sleep apnea with normal oximetry. *Pediatrics International* 63:1376–80. <https://doi.org/10.1111/ped.14663>
 71. Neruntarat C, Chantapant S (2010) Prevalence of sleep apnea in HRH Princess Maha Chakri Srinthorn Medical Center Thailand. *Sleep and Breathing* 15:641–8. <https://doi.org/10.1007/s11325-010-0412-x>
 72. Tawaranurak K, Leelasawatsuk P, Chaiyarakjirakun V (2019) Prevalence, risk factors and clinical manifestation of patients suspected as having obstructive sleep apnea in Songklanagarind Hospital sleep center. *J Health Sci Med Res* 37:305–12. <https://doi.org/10.31584/jhsmr.201965>
 73. Veeravigrom M, Desudchit T (2016) Prevalence of Sleep Disorders in Thai Children. *Indian J Pediatr* 83:1237–41. <https://doi.org/10.1007/s12098-016-2148-5>
 74. Morell-García D, Peña-Zarza JA, Sanchís P, Piérola J, de la Peña M, Bauça JM et al (2021) Polysomnographic characteristics of snoring children: a familial study of obstructive sleep apnea

- syndrome. *Arch Bronconeumol* 57:387–92. <https://doi.org/10.1016/j.arbres.2020.01.006>
75. Quintana-Gallego E, Carmona-Bernal C, Capote F, Sánchez-Armengol Á, Botbol-Benhamou G, Polo-Padillo J et al (2004) Gender differences in obstructive sleep apnea syndrome: a clinical study of 1166 patients. *Respir Med* 98:984–9. <https://doi.org/10.1016/j.rmed.2004.03.002>
 76. Valenza MC, Valenza G, Muñoz-Casaubon T, Botella-López M, Puenteledura EJ, Arroyo-Morales M et al (2012) Epidemiology of sleep-related complaints associated with obstructive sleep apnea, insomnia and non-restorative sleep in an at-risk population in Granada Spain. *Sleep Biol Rhythms* 10:222–30. <https://doi.org/10.1111/j.1479-8425.2012.00565.x>
 77. Durán J, Esnaola S, Rubio R, Izutueta Á (2001) Obstructive sleep apnea-hypopnea and related clinical features in a population-based sample of subjects aged 30 to 70 yr. *Am J Respir Crit Care Med* 163:685–9. <https://doi.org/10.1164/ajrccm.163.3.2005065>
 78. Zamarrón C, Hornero R, del Campo F, Abásolo D, Alvarez D (2006) Heart rate regularity analysis obtained from pulse oximetric recordings in the diagnosis of obstructive sleep apnea. *Sleep and Breathing* 10:83–9. <https://doi.org/10.1007/s11325-005-0049-3>
 79. Cho ER, Kim H, Seo HS, Suh S, Lee SK, Shin C (2013) Obstructive sleep apnea as a risk factor for silent cerebral infarction. *J Sleep Res* 22:452–8. <https://doi.org/10.1111/jsr.12034>
 80. Han SH, Lee GY, Hyun W, Kim Y, Jang JS. Obstructive sleep apnea in airline pilots during daytime sleep following overnight flights. *J Sleep Res* 2021;30. <https://doi.org/10.1111/jsr.13375>.
 81. Kim JK, In KH, Kim JH, You SH, Kang KH, Shim JJ et al (2004) Prevalence of sleep-disordered breathing in middle-aged Korean men and women. *Am J Respir Crit Care Med* 170:1108–13. <https://doi.org/10.1164/rccm.200404-519OC>
 82. Lee J, Choi JH, Kim SG (2019) Prevalence and characteristics of subjects with obstructive sleep apnea among adults with insomnia disorder. *Sleep Med Res* 10:108–12. <https://doi.org/10.17241/smr.2019.00409>
 83. Fietze I, Laharnar N, Obst A, Ewert R, Felix SB, Garcia C, et al. Prevalence and association analysis of obstructive sleep apnea with gender and age differences – Results of SHIP-Trend. *J Sleep Res* 2018;28. <https://doi.org/10.1111/jsr.12770>.
 84. Urschitz MS, Brockmann PE, Schlaud M, Poets CF (2010) Population prevalence of obstructive sleep apnoea in a community of German third graders. *Eur Respir J* 36:556–68. <https://doi.org/10.1183/09031936.00078409>
 85. Zacharias HU, Weihs A, Habes M, Wittfeld K, Frenzel S, Rashid T et al (2021) Association between obstructive sleep apnea and brain white matter hyperintensities in a population-based cohort in Germany. *JAMA Netw Open*. <https://doi.org/10.1001/jamanetworkopen.2021.28225>
 86. Katz SL, Vaccani JP, Barrowman N, Momoli F, Bradbury CL, Murto K (2014) Does neck-to-waist ratio predict obstructive sleep apnea in children? *J Clin Sleep Med* 10:1303–8. <https://doi.org/10.5664/jcs.m.4284>
 87. Selvadurai S, Voutsas G, Propst EJ, Wolter NE, Narang I (2019) Obstructive sleep apnea in children aged 3 years and younger: rate and risk factors. *Paediatrics and Child Health (Canada)* 25:432–8. <https://doi.org/10.1093/pch/pxz097>
 88. Dorkova Z, Sopkova Z, Tkacova R (2010) “CEM” risk factors and severity of obstructive sleep apnoea in central European Roma and non-Roma patients referred for a diagnostic polysomnography. *Int J Public Health* 55:429–34. <https://doi.org/10.1007/s00038-010-0146-3>
 89. Dostálová Š, Šusta M, Nepožitek J, Peřinová P, Přihodová I, Ibarburu Lorenzo V et al (2020) Polysomnographic findings in individuals over 50 years of age lacking subjective signs of sleep disturbance. *Ceska a Slovenska Neurologie a Neurochirurgie* 83:57–63. <https://doi.org/10.14735/amcsnn202057>
 90. Kobayashi R, Miyazaki S, Karaki M, Hoshikawa H, Nakata S, Hara H, et al. Obstructive sleep apnea in Asian primary school children. *Sleep and Breathing* 2013;18. <https://doi.org/10.1007/s11325-013-0909-1>.
 91. Sasai-Sakuma T, Kikuchi K, Inoue Y (2016) Cross-sectional study of obstructive sleep apnea syndrome in Japanese public transportation drivers its prevalence and association with pathological objective daytime sleepiness. *J Occup Environ Med* 58:455–8. <https://doi.org/10.1097/JOM.0000000000000692>
 92. Bruyneel M, Ameze L, Ninane V (2010) Sleep apnea syndrome in a young cosmopolite urban adult population: risk factors for disease severity. *Sleep and Breathing* 15:543–8. <https://doi.org/10.1007/s11325-010-0398-4>
 93. Mungo A, Hein M, Lanquart JP, Loas G (2021) Atypical depression as a risk factor for obstructive sleep apnea syndrome in young adults. *Encephale*. <https://doi.org/10.1016/j.encep.2021.02.018>
 94. Jalilolghadr S, Taghiloo M, Parsarad E, Taherahmadi M, Taherahmadi H. Evaluation of sleep-disordered breathing in children and adolescents referred to the sleep ward of Qazvin children’s hospital during 2014–2019. *Int J Pediatr* 2022;10. <https://doi.org/10.22038/IJP.2021.58958.4598>.
 95. Sadeghniai-Haghighi K, Montazeri A, Khajeh-Mehrzi A, Ghajarzadeh M, Alemohammad ZB, Aminian O et al (2015) The STOP-BANG questionnaire: reliability and validity of the Persian version in sleep clinic population. *Quality of Life Research* 24:2025–30. <https://doi.org/10.1007/s1136-015-0923-9>
 96. Drakou T, Steiropoulos P, Saroglou M, Georgopoulou A, Kazis D, Papagiannopoulos S et al (2021) The presence of insomnia and depression contributes to the acceptance of an initial treatment trial of continuous positive airway pressure therapy in patients with obstructive sleep apnea. *Sleep and Breathing* 25:1803–12. <https://doi.org/10.1007/s11325-020-02266-z>
 97. Ioannidou D, Kalamaras G, Kotoulas SC, Pataka A. Smoking and obstructive sleep apnea: Is there an association between these cardiometabolic risk factors?—gender analysis. *Medicina* 2021;57. <https://doi.org/10.3390/medicina57111137>.
 98. Garbarino S, Guglielmi O, Campus C, Mascialino B, Pizzorni D, Nobili L et al (2016) Screening, diagnosis, and management of obstructive sleep apnea in dangerous-goods truck drivers: to be aware or not? *Sleep Med* 25:98–104. <https://doi.org/10.1016/j.sleep.2016.05.015>
 99. Scarlata S, Pedone C, Curcio G, Cortese L, Chieruro D, Fontana D et al (2013) Pre-polysomnographic assessment using the Pittsburgh Sleep Quality Index questionnaire is not useful in identifying people at higher risk for obstructive sleep apnea. *J Med Screen* 20:220–6. <https://doi.org/10.1177/0969141313511591>
 100. Franklin KA, Sahlin C, Stenlund H, Lindberg E (2013) Sleep apnoea is a common occurrence in females. *Eur Respir J* 41:610–5. <https://doi.org/10.1183/09031936.00212711>
 101. Coussa-Koniski ML, Saliba E, Welty FK, Deeb M. Epidemiological characteristics of obstructive sleep apnea in a hospital-based historical cohort in Lebanon. *PLoS One* 2020;15. <https://doi.org/10.1371/journal.pone.0231528>.
 102. Arrazola-Cortés E, Hernández-Cervantes J, González-Pérez B, Sauri-Suárez S, Berenice López-Hernández L, Toledo-Lozano CG et al (2017) Polysomnography-based diagnosis in Mexican adult patients with Obstructive Sleep Apnea Syndrome (OSAS) clinical suspicion. *Neuroendocrinol Lett* 38:449–54
 103. Deegan PC, McNicholas WT (1996) Predictive value of clinical features for the obstructive sleep apnoea syndrome. *Eur Respir J* 9:117–24. <https://doi.org/10.1183/09031936.96.09010117>

104. Duong-Quy S, Dang Thi Mai K, Van Tran N, Nguyen Xuan Bich H, Hua-Huy T, Chalumeau F et al (2018) Étude de la prévalence du syndrome d'apnées obstructives du sommeil au Vietnam. *Rev Mal Respir* 35:14–24. <https://doi.org/10.1016/j.rmr.2017.10.006>
105. Klawe JJ, Laudenska A, MiEkwiec I, Tafil-Klawe M (2005) Occurrence of obstructive sleep apnea in a group of shift worked police officers. *J Physiol Pharmacol* 56:115–7
106. Randby A, Namtvedt SK, Hrubos-Strøm H, Einvik G, Somers VK, Omland T (2013) Sex-dependent impact of OSA on digital vascular function. *Chest* 144:915–22. <https://doi.org/10.1378/chest.12-2283>
107. Roche J, Rae DE, Redman KN, Knutson KL, von Schantz M, Gómez-Olivé FX et al (2021) Impact of obstructive sleep apnea on cardiometabolic health in a random sample of older adults in rural South Africa: building the case for the treatment of sleep disorders in underresourced settings. *J Clin Sleep Med* 17:1423–34. <https://doi.org/10.5664/jcsm.9214>
108. Vega-Osorio PA ro;, Orozco-Porras LD, Castro-Bonilla NL, Patiño-Ríos VH, Camacho PA. The analysis of polysomnographic variables and their relation with severity of obstructive sleep apnea and hypopnea syndrome. *Acta Otorrinolaringol Cir Cabeza Cuello* 2020;48:69–77.
109. Wahida AB, Ilhamah O, Suffian AM, Aimi MF, Norlen M, Wong SV (2013) Obstructive sleep apnea among commercial vehicle drivers in Malaysia: issues and initiatives. *Health N Hav* 05:80–6. <https://doi.org/10.4236/health.2013.58a2012>
110. Wali SO, Abalkhail B, Krayem A (2017) Prevalence and risk factors of obstructive sleep apnea syndrome in a Saudi Arabian population. *Ann Thorac Med* 12:88–94. <https://doi.org/10.4103/1817-1737.203746>
111. Ferreira-Santos D, Rodrigues PP. Enhancing obstructive sleep apnea diagnosis with screening through disease phenotypes: algorithm development and validation. *JMIR Med Inform* 2021;9. <https://doi.org/10.2196/25124>.
112. del Brutto OH, Mera RM, Castillo PR, Recalde BY, Costa AF. Previously diagnosed obstructive sleep apnea is not associated with increased risk of SARS-CoV-2 infection in community-dwelling older adults living in a highly endemic setting. *Clin Neurol Neurosurg* 2021;205. <https://doi.org/10.1016/j.clineuro.2021.106639>.
113. Abdel Dayem A, Madkour A, Abdel-Fattah E, Abdelazeem M. Prevalence of obstructive sleep apnea in a sample of Egyptian railway drivers. *Egypt J Chest Dis Tuberc* 2022;71.
114. Dashzeveg S, Oka Y, Purevtogtokh M, Tumurbaatar E, Lkhagasuren B, Luvsannorov O, et al. Obstructive sleep apnea in a clinical population: Prevalence, predictive factors and clinical characteristics of patients referred to a sleep center in mongolia. *Int J Environ Res Public Health* 2021;18. <https://doi.org/10.3390/ijerph182212032>.
115. Mosharraf-Hossain A, Ahmed K, Islam M, Chakraborty R (2015) A Community study of obstructive sleep apnea hypopnea syndrome (OSAHS) in middle-aged Bangladeshi population. *Bangladesh Med Res Counc Bull* 41:13–8
116. Eastwood PR, Malhotra A, Palmer LJ, Kezirian EJ, Horner RL, Ip MS et al (2010) Obstructive sleep apnoea: from pathogenesis to treatment: current controversies and future directions. *Respirology* 15:587–95. <https://doi.org/10.1111/j.1440-1843.2009.01699.x>
117. Malhotra A, White DP (2002) Obstructive sleep apnoea. *Lancet* 360:237–45
118. Eckert DJ (2018) Phenotypic approaches to obstructive sleep apnoea – new pathways for targeted therapy. *Sleep Med Rev* 37:45–59. <https://doi.org/10.1016/j.smrv.2016.12.003>
119. Li AM, So HK, Au CT, Ho C, Lau J, Ng SK et al (2010) Epidemiology of obstructive sleep apnoea syndrome in Chinese children: a two-phase community study. *Thorax* 65:991–7. <https://doi.org/10.1136/thx.2010.134858>
120. Kapur VK, Auckley DH, Chowdhuri S, Kuhlmann DC, Mehra R, Ramar K et al (2017) Clinical practice guideline for diagnostic testing for adult obstructive sleep apnea: an American academy of sleep medicine clinical practice guideline. *J Clin Sleep Med* 13:479–504. <https://doi.org/10.5664/JCSM.6506>
121. Li P, Ning X-H, Lin H, Zhang N, Gao Y-F, Ping F (2020) Continuous positive airway pressure versus mandibular advancement device in the treatment of obstructive sleep apnea: a systematic review and meta-analysis. *Sleep Med* 72:5–11. <https://doi.org/10.1016/j.sleep.2020.03.015>
122. Ieto V, Kayamori F, Montes MI, Hirata RP, Gregório MG, Alencar AM et al (2015) Effects of oropharyngeal exercises on snoring: a randomized trial. *Chest* 148:683–91. <https://doi.org/10.1378/chest.14-2953>
123. Alessandri-Bonetti A, Bortolotti F, Moreno-Hay I, Michelotti A, Cordaro M, Alessandri-Bonetti G, et al. Effects of mandibular advancement device for obstructive sleep apnea on temporomandibular disorders: a systematic review and meta-analysis. *Sleep Med Rev* 2019;48. <https://doi.org/10.1016/j.smrv.2019.101211>.
124. Vecchierini MF, Attali V, Collet JM, d'Ortho MP, Goutorbe F, Kerbrat JB et al (2021) Mandibular advancement device use in obstructive sleep apnea: ORCADES study 5-year follow-up data. *J Clin Sleep Med* 17:1695–705. <https://doi.org/10.5664/jcsm.9308>
125. Uniken Venema JAM, Doff MHJ, Joffe-Sokolova D, Wijkstra PJ, van der Hoeven JH, Stegenga B et al (2020) Long-term obstructive sleep apnea therapy: a 10-year follow-up of mandibular advancement device and continuous positive airway pressure. *J Clin Sleep Med* 16:353–9. <https://doi.org/10.5664/JCSM.8204>

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Springer Nature or its licensor (e.g. a society or other partner) holds exclusive rights to this article under a publishing agreement with the author(s) or other rightsholder(s); author self-archiving of the accepted manuscript version of this article is solely governed by the terms of such publishing agreement and applicable law.

Authors and Affiliations

Anna Beatriz de Araujo Dantas¹ · Flávio Magno Gonçalves^{2,3} · Agnes Andrade Martins^{1,3} · Giorvan Anderson Alves⁴ · José Stechman-Neto^{2,3} · Camila de Castro Corrêa^{3,5} · Rosane Sampaio Santos^{2,3} · Weslania Viviane Nascimento^{3,6} · Cristiano Miranda de Araujo^{2,3} · Karinna Veríssimo Meira Taveira^{1,3} 

Cristiano Miranda de Araujo
cristianoaraujo@narsm.com.br

¹ Department of Morphology – Center of Biosciences, Federal University of Rio Grande do Norte (UFRN), BR 101- Lagoa Nova, Natal RN - 59072-970, Brazil

² Tuiuti University of Paraná (UTP), Curitiba, Brazil

³ Center for Advanced Studies in Systematic Review and Meta-Analysis (NARSM), Curitiba, Brazil

⁴ Federal University of Paraíba (UFPB), João Pessoa, Brazil

⁵ Planalto University Center of the Federal District (UNIPLAN), Brasília, Brazil

⁶ Universitat Autònoma de Barcelona, Barcelona, Spain