



Research Article

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An Audit of Sleep Disordered Breathing in Central Sri Lanka, Resource Poor Setting

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To Cite This article: Stephen E Berger, Ph. D, ABPP* and Bina Parekh, Ph. D, A Practical Guide for How to Use the Parekh-Berger Hierarchical Interpretive Phenomenological Analysis Model with Application to Inclusive Research. Am J Biomed Sci & Res. 2026 30(4) AJBSR.MS.ID.003941,

DOI: [10.34297/AJBSR.2026.30.003941](https://doi.org/10.34297/AJBSR.2026.30.003941)

Received: 📅 March 05, 2026; Published: 📅 March 16, 2026

Abstract

Background: Sleep disorders, particularly Obstructive Sleep Apnoea (OSA), significantly impact public health. Polysomnography (PSG) is the gold standard for diagnosis. This study aimed to investigate the clinical indications, diagnostic outcomes, and correlations in patients undergoing overnight PSG at a tertiary care center in Sri Lanka.

Methods: A retrospective descriptive study was conducted on 932 sleep study records from the Respiratory and Research Unit 2 of the National Hospital Kandy between January 2013 and December 2023. Data on demographics, anthropometry (BMI, neck circumference), clinical scores (Epworth Sleepiness Scale - ESS, Mallampati score), and comorbidities were extracted. Statistical analysis was performed using SPSS version 26.0 and Python 3.8.

Results: Among 932 patients (583 males, 62.6%) underwent PSG, OSA was diagnosed in 676 (72.5%), with severe OSA representing the largest subgroup (296, 31.7%), followed by mild (205, 22.0%) and moderate disease (175, 18.8%). The cohort demonstrated a high anthropometric risk profile, with a mean BMI of 30.0 kg/m² and 801 patients (85.9%) classified as overweight or obese using Asia-Pacific criteria. A neck circumference >40 cm was observed in 390 patients (41.8%), while Mallampati class III-IV anatomy was present in 590 (63.3%). Hypertension was the most prevalent comorbidity (430, 46.1%), followed by asthma (231, 24.8%) and diabetes mellitus (219, 23.5%). Nocturnal hypoxemia was prominent, with a mean lowest SpO₂ of 80.6%; 662 patients (71.0%) experienced desaturation <90%, including 351 (37.7%) falling below 80%. Neck circumference correlated strongly with AHI (r=0.45, p<0.001). On multivariate analysis, lowest SpO₂, BMI, male gender, ESS score, neck circumference, hypertension, and habitual snoring independently predicted OSA severity.

Conclusions: OSA is highly prevalent among patients referred for PSG in this setting, often presenting with severe disease and significant comorbidities. Neck circumference is a strong predictor of severity. Implementing screening tools like STOP-BANG in primary care could optimize referral pathways.

Keywords: Polysomnography, Obstructive Sleep Apnoea, Sri Lanka, Sleep Disorders, STOP-BANG

Introduction

Sleep disorders, including Obstructive Sleep Apnoea (OSA), are a growing global health concern associated with significant morbidity, including cardiovascular disease and metabolic dysfunction.

While international data highlights the burden of OSA, there is a paucity of large-scale data regarding the diagnostic patterns and clinical characteristics of sleep disorder patients in Sri Lanka.



Polysomnography (PSG) remains the gold standard for diagnosis, yet access in low-resource settings can be limited. Understanding the local epidemiology is crucial for resource allocation. This study aims to analyze the indications, outcomes, and clinical correlates of patients undergoing PSG at the National Hospital Kandy over a ten-year period to identify gaps in current referral and management practices. Specifically, the study seeks to describe the demographic and clinical characteristics of patients undergoing PSG, determine the prevalence and severity of sleep disorders diagnosed via PSG, and identify the most common indications for referral. Furthermore, the study aims to examine the association between OSA severity and established risk factors, including Body Mass Index (BMI), age, gender, neck circumference, snoring frequency, Epworth Sleepiness Scale (ESS) score, Mallampati score, and relevant comorbidities.

Methodology

A retrospective descriptive study was conducted involving patients who underwent full overnight Polysomnography (PSG) at the Respiratory and Research Unit 2 and Suwapiyasa Clinic of the National Hospital, as well as the Central Chest Clinic, Kandy, Sri Lanka. The study period spanned ten years, from January 2013 to

December 2023. The study population consisted of 932 patients aged 12 years and above who were referred for sleep studies from respiratory or associated clinics. Patients were excluded if they had incomplete or missing PSG data, technically failed studies, or if they presented with central apnoea or other unconfirmed neurological sleep disorders.

Data were retrospectively collected from multiple sources, including PSG reports, clinical records (Bed Head Tickets - BHTs), electronic patient records, and sleep study scoring sheets. The diagnosis and scoring of sleep disorders adhered to the guidelines established by the American Academy of Sleep Medicine (AASM), World Health Organization (WHO) Sleep Health Metrics, and the Sri Lanka Respiratory Society. Statistical analysis was performed using SPSS and Python for regression and significance testing. Ethical clearance was obtained from the institutional review board.

Results

Study Population and Clinical Characteristics

A total of 932 patients were included in the study, comprising 583 males (62.6%) and 349 females (37.4%) (Table 1&Figure 1).

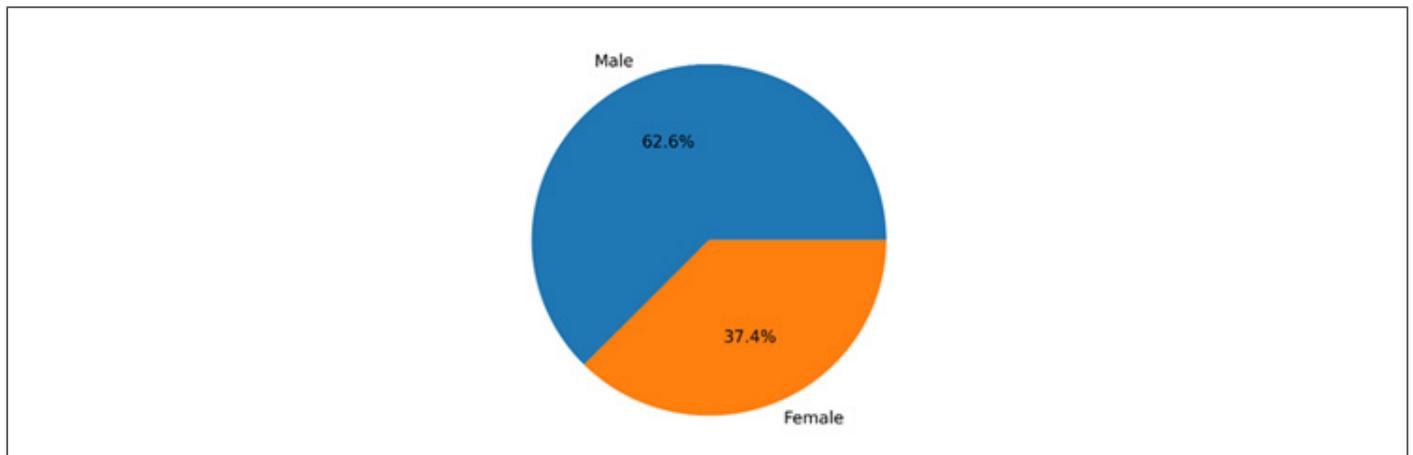


Figure 1: Gender distribution of the Study Population.

Table 1: Demographic and Clinical Characteristics of the Study Population.

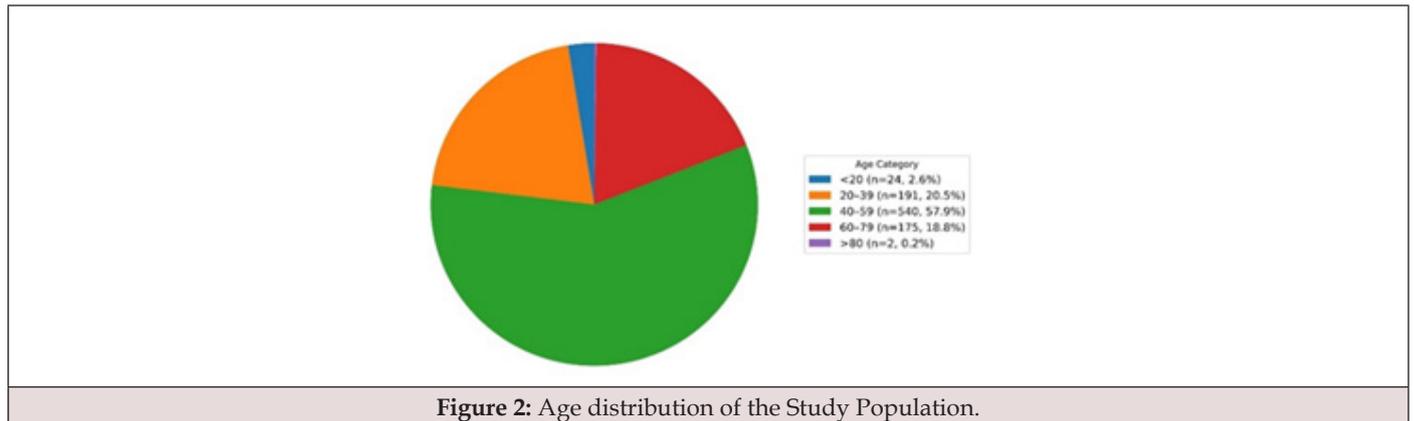
| Gender | n | Percentage (%) |
|--------|-----|----------------|
| Male | 583 | 62.60% |
| Female | 349 | 37.40% |

The mean age of the study population was 50.4 years (SD ± 12.3), with a range of 12 to 88 years. The majority of patients were in the middle-aged group (40-59 years). Most patients were aged

40–59 years (57.9%), while only 0.2% were older than 80 years (Table 2&Figure 2).

Table 2: Age distribution of the Study Population.

| Age Category | n | Percentage (%) |
|--------------|-----|----------------|
| <20 | 24 | 2.60% |
| 20-39 | 191 | 20.50% |
| 40-59 | 540 | 57.90% |
| 60-79 | 175 | 18.80% |
| >80 | 2 | 0.20% |

**Table 2:** Age distribution of the Study Population.

| Age Category | n | Percentage (%) |
|--------------|-----|----------------|
| <20 | 24 | 2.60% |
| 20-39 | 191 | 20.50% |
| 40-59 | 540 | 57.90% |
| 60-79 | 175 | 18.80% |
| >80 | 2 | 0.20% |

Comorbidities

Hypertension was the most prevalent condition, affecting 430 patients (46.1%). Respiratory comorbidities were also

notable, particularly bronchial asthma (24.8%), COPD 2.9% (27). Cardiovascular disease and hypothyroidism were less frequent but clinically relevant (Table 3&Figure 3).

Table 3: Comorbidities of the Study Population.

| Comorbidity | n | Percentage (%) |
|-------------------|-----|----------------|
| Hypertension | 430 | 46.10% |
| Asthma | 231 | 24.80% |
| DM | 219 | 23.50% |
| Insomnia | 216 | 23.20% |
| Allergic Rhinitis | 210 | 22.50% |
| Chronic Cough | 162 | 17.40% |
| IHD | 101 | 10.80% |
| Hypothyroidism | 101 | 10.80% |

| | | |
|------------|----|-------|
| Depression | 66 | 7.10% |
| PVD | 29 | 3.10% |
| Stroke | 28 | 3.00% |
| COPD | 27 | 2.90% |
| Acromegaly | 9 | 1.00% |

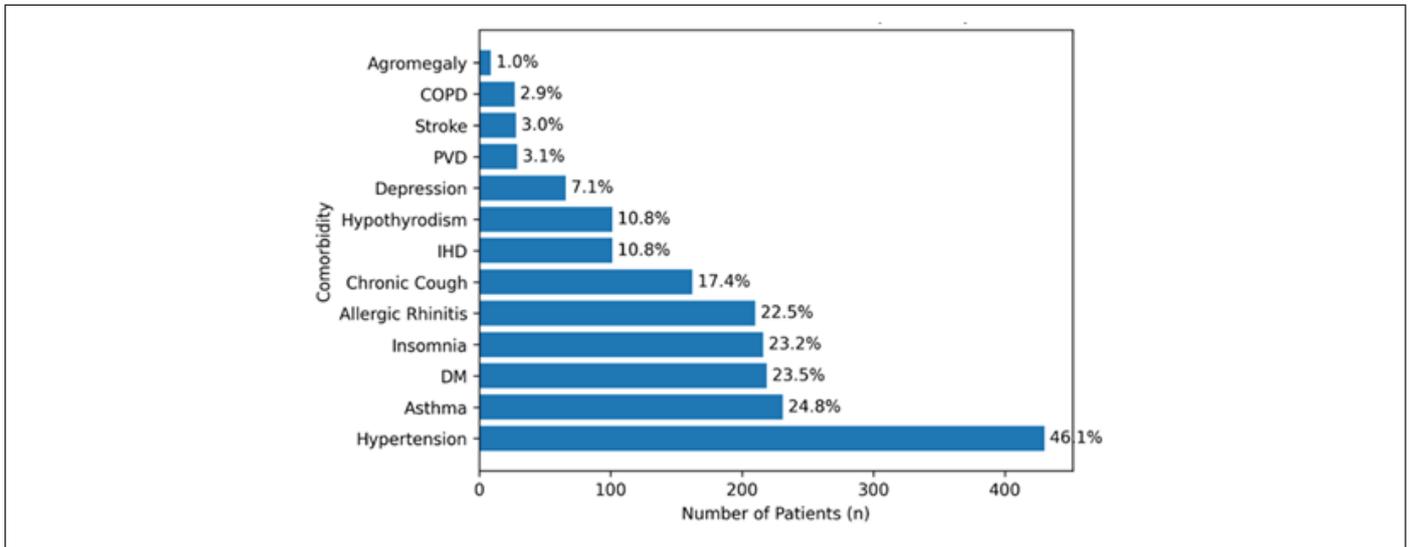


Figure 3: Comorbidity distribution of the Study Population.

Anthropometry

The mean body mass index of the study population (n = 932) was 30.0 kg/m², BMI values ranged from 16 to 56 kg/m². Among the study population (n = 932), 200 patients (21.5%) had a neck circumference < 37 cm, 342 patients (36.7%) had a neck circumference between 37–40 cm, and 390 patients (41.8%) had a neck circumference > 40 cm. The majority of participants (41.8%) had a neck circumference exceeding 40 cm. The mean neck circumference of the study population (n = 932) was 39.7 cm, with

a median value of 40 cm. Neck circumference ranged from 10 cm to 57 cm. Body Mass Index (BMI) was classified according to Asia-Pacific BMI cut-offs (normal 18.5–22.9 kg/m², overweight 23–24.9 kg/m², obese ≥25 kg/m²) recommended for Asian populations [1]. The majority of patients were categorized as obese, with 358 patients (38.4%) classified as Obese I (BMI 25–29.9 kg/m²) and 443 patients (47.5%) as Obese II (BMI ≥30 kg/m²). Only a small proportion of the population fell within the normal BMI range (5.2%), while 7.8% were classified as overweight and 1.1% as underweight (Tables 4-7&Figures 4,5)

Table 4: Neck circumference of the Study Population.

| Neck Circumference Category | n | Percentage (%) |
|-----------------------------|-----|----------------|
| < 37 cm | 200 | 21.50% |
| 37-40 cm | 342 | 36.70% |
| > 40 cm | 390 | 41.80% |

Table 5: Neck circumference of the Study Population.

| BMI Category | n | Percentage (%) |
|----------------------|-----|----------------|
| Underweight (<18.5) | 10 | 1.10% |
| Normal (18.5-22.9) | 48 | 5.20% |
| Overweight (23-24.9) | 73 | 7.80% |
| Obese I (25-29.9) | 358 | 38.40% |
| Obese II (≥30) | 443 | 47.50% |

Table 6: Anthropometric measurements of the Study Population.

| Anthropometry | Value |
|----------------------------|------------------------|
| Mean Body Mass Index (BMI) | 30.0 kg/m ² |
| Mean Neck Circumference | 39.7 cm |

Table 7: ESS components of the Study Population.

| ESS characteristic | Value |
|-------------------------------------|--------------|
| Mean Epworth Sleepiness Scale (ESS) | 11.33 ± 5.97 |
| Mean Sleep Efficiency | 85.20% |

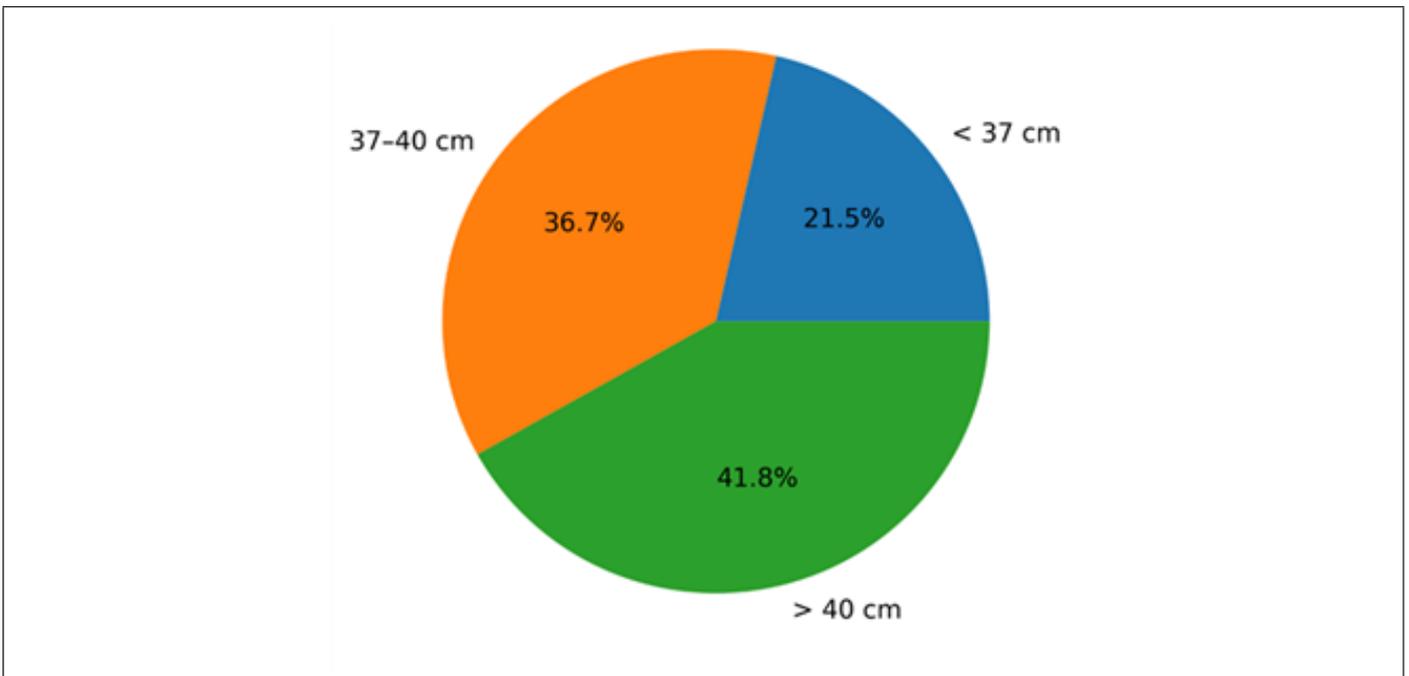


Figure 4: Neck Circumference distribution of the Study Population.

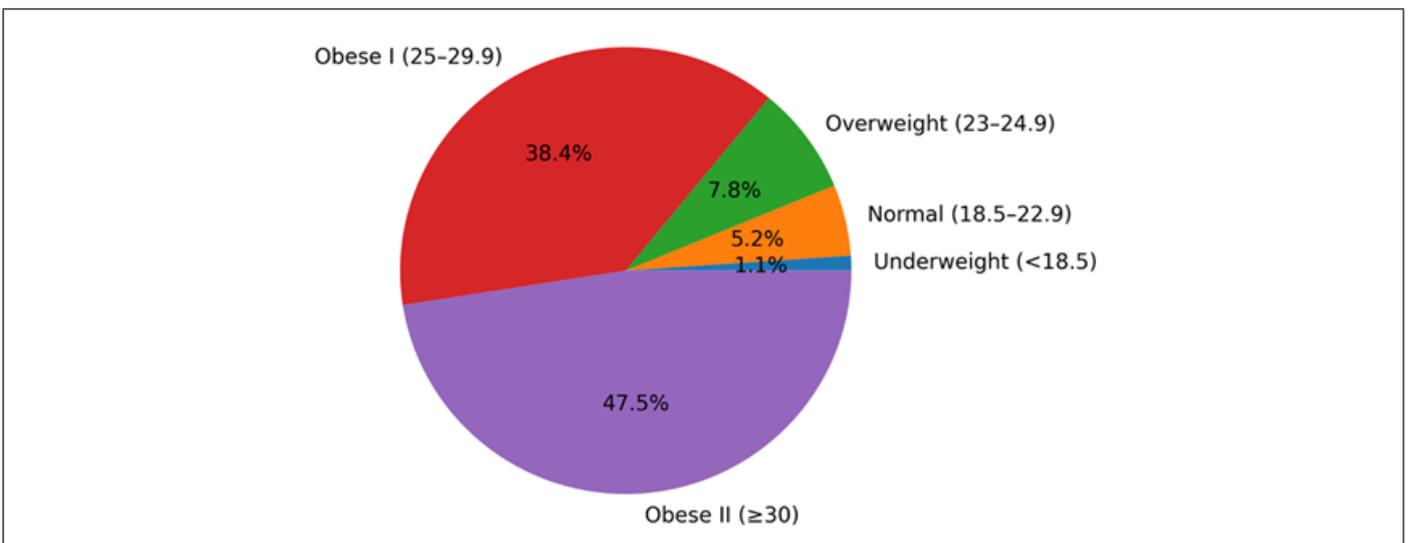


Figure 5: BMI Categories (Asia-Pacific) in the Study Population.

Indications for Polysomnography

The primary indications for referring patients for Polysomnography (PSG) were symptoms suggestive of sleep-disordered breathing. As detailed in, snoring was the most frequent reason for referral, accounting for 41.2% (n=384) of the cases. Excessive daytime sleepiness was the sole indication in 22.6% (n=211) of patients, while a combination of both snoring and

daytime sleepiness prompted referral in 12.1% (n=113) of cases.

Less common indications included obesity (7.4%, n=69), unexplained breathlessness (4.5%, n=42), and morning headaches (3.7%, n=34). A small subset of patients was referred for specific diagnostic clarifications such as CPAP response evaluation (2.9%, n=27), finding a possible cause for pulmonary hypertension (2.4%, n=22), and insomnia (2.1%, n=20) (Table 8&Figure 6).

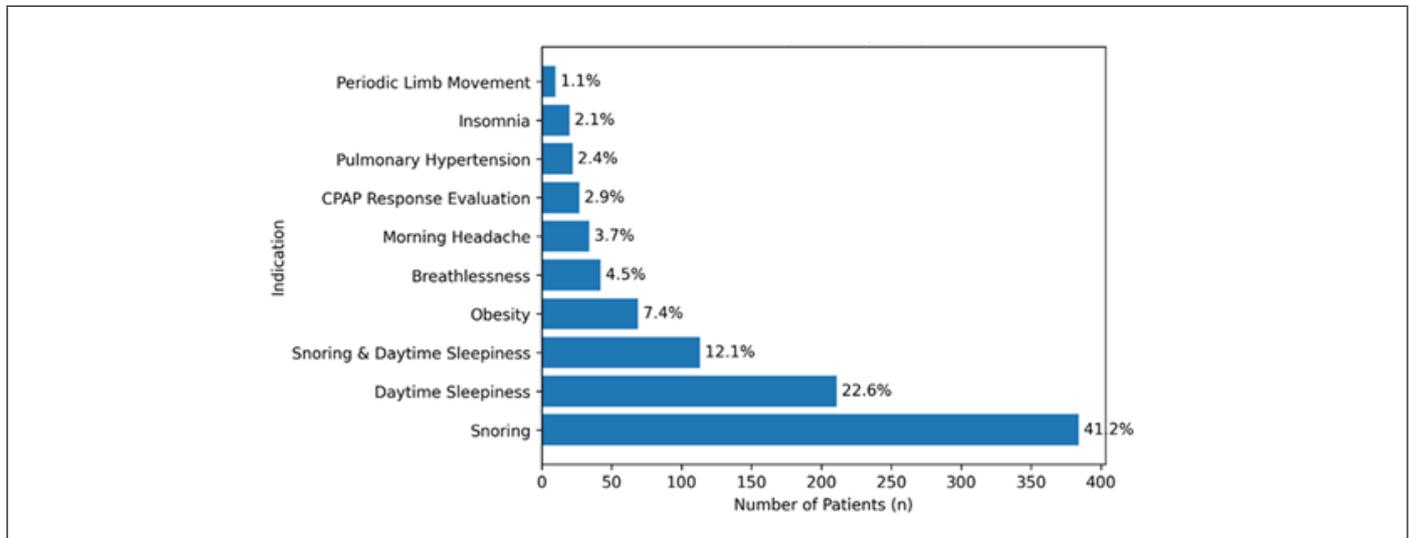


Figure 6: Indications for Polysomnography in the Study Population.

Table 8: Indications for Polysomnography.

| Indication | n | Percentage (%) |
|------------------------------|-----|----------------|
| Snoring | 384 | 41.20% |
| Daytime Sleepiness | 211 | 22.60% |
| Snoring & Daytime Sleepiness | 113 | 12.10% |
| Obesity | 69 | 7.40% |
| Breathlessness | 42 | 4.50% |
| Morning Headache | 34 | 3.70% |
| CPAP Response Evaluation | 27 | 2.90% |
| Pulmonary Hypertension | 22 | 2.40% |
| Insomnia | 20 | 2.10% |
| Periodic Limb Movement | 10 | 1.10% |

Prevalence and Severity of Sleep Disorders

Upon analysis of the PSG data, a significant proportion of the study population was diagnosed with Obstructive Sleep Apnoea (OSA). Only 27.5% (n=256) of the patients had a normal Apnoea-Hypopnoea Index (AHI < 5). The remaining 72.5% (n=676) were diagnosed with varying degrees of OSA.

Notably, the distribution of severity was skewed towards the severe spectrum. Severe OSA (AHI > 30) was the most prevalent diagnosis, affecting 31.7% (n=296) of the total cohort. Moderate

OSA was found in 18.8% (n=175) of patients, while 22.0% (n=205) were diagnosed with mild OSA. A minority of cases presented with complex phenotypes, including Mild OSA with Narcolepsy (n=6) and Severe OSA combined with Obesity Hypoventilation Syndrome (n=5) (Table 9&Figure 7).

Predictors and Correlations

Bivariate and multivariate analyses identified several significant predictors of OSA severity in the study population.

Table 9: Distribution of OSA Severity (Based on AHI).

| Diagnosis Category | n | Percentage (%) |
|--------------------------|------|----------------|
| No OSA (AHI < 5) | 256 | 27.50% |
| Mild OSA (AHI 5-15) | 205* | 22.00% |
| Moderate OSA (AHI 15-30) | 175 | 18.80% |
| Severe OSA (AHI > 30) | 296^ | 31.70% |
| Total | 932 | 100% |

*Note: Includes Mild OSA with Narcolepsy (n=6) ^Includes Severe OSA with Obesity Hypoventilation Syndrome (n=5)

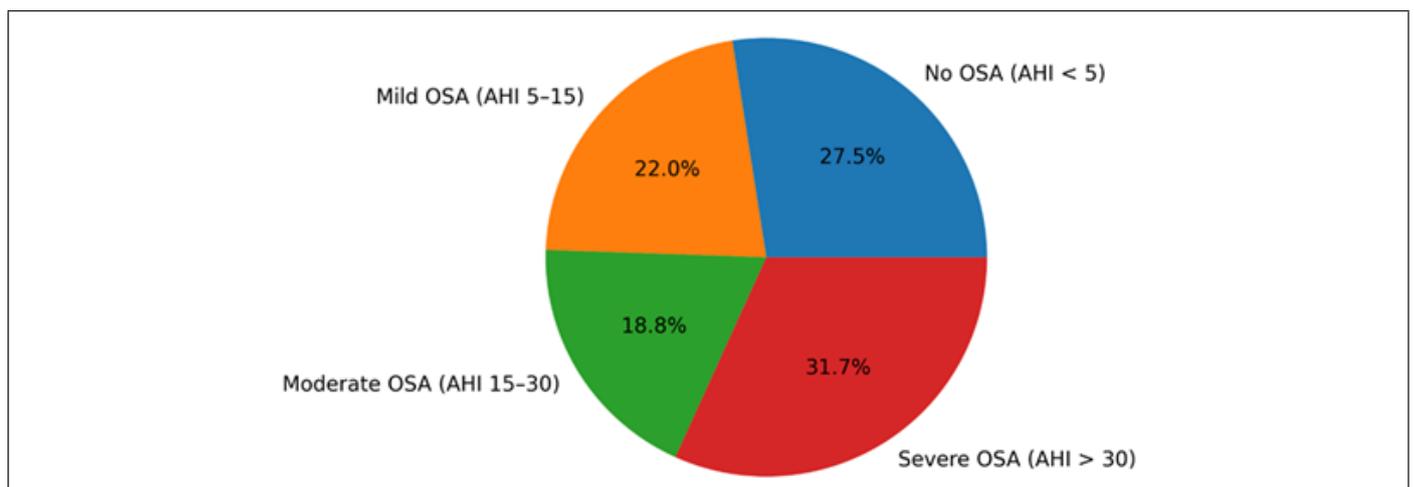


Figure 7: Distribution of the OSA Severity of the Study Population.

Demographic Predictors:

Age: Age demonstrated a positive linear correlation with AHI (p=0.002). Patients aged over 50 years exhibited disproportionately higher AHI values compared to younger cohorts, although a notable prevalence of OSA was also observed in the 30–40 age group.

Gender: Male gender was strongly associated with higher disease severity (p = 0.014). Males accounted for the majority of severe OSA cases, whereas females were more frequently diagnosed

with mild-to-moderate disease or non-OSA sleep disorders.

Anthropometric Predictors:

Neck Circumference: A strong positive correlation was found between neck circumference and OSA severity (r=0.45, p<0.001). A threshold of >40 cm was a critical predictor; patients exceeding this limit were significantly more likely to have moderate-to-severe OSA (Table 10).

Table 10: OSA severity and Neck circumference distribution of the Study Population.

| OSA Severity | < 37 cm n (%) | 37-40cm n (%) | > 40cm n (%) |
|--------------|---------------|---------------|--------------|
| No OSA | 47 (5.0%) | 97 (10.4%) | 112 (12.0%) |
| Mild OSA | 44 (4.7%) | 74 (7.9%) | 87 (9.3%) |
| Moderate OSA | 41 (4.4%) | 63 (6.8%) | 71 (7.6%) |
| Severe OSA | 68 (7.3%) | 108 (11.6%) | 120 (12.8%) |

Body Mass Index (BMI): BMI showed a significant positive association with AHI. While the mean BMI was 30.0 kg/m², severe OSA was frequently identified in patients with a BMI of 29–30 kg/

m², indicating a lower obesity threshold for severe disease in this population compared to Western guidelines (Table 11).

Table 11: OSA severity and BMI Category distribution of the Study Population.

| OSA Severity | Underweight n(%) | Normal n(%) | Overweight n(%) | Obese I n(%) | Obese II n(%) |
|--------------|------------------|-------------|-----------------|--------------|---------------|
| No OSA | 0 (0.0%) | 12 (1.3%) | 25 (2.7%) | 97 (10.4%) | 122 (13.1%) |
| Mild OSA | 4 (0.4%) | 10 (1.1%) | 13 (1.4%) | 88 (9.4%) | 90 (9.7%) |
| Moderate OSA | 2 (0.2%) | 15 (1.6%) | 18 (1.9%) | 57 (6.1%) | 83 (8.9%) |
| Severe OSA | 4 (0.4%) | 11 (1.2%) | 17 (1.8%) | 116 (12.4%) | 148 (15.9%) |

Clinical and Physiological Predictors:

Mallampati Score: The distribution of Mallampati scores demonstrated a predominance of higher airway grades. Mallampati class III was the most frequent (35.6%), followed by class II (29.7%) and class IV (27.7%), while only 7.0% of patients had class I anatomy. Overall, 63.3% of the cohort had Mallampati class III or IV, indicating a high prevalence of anatomically narrowed

upper airway among patients undergoing polysomnography. Airway crowding, indicated by higher Mallampati classes (III and IV), correlated with increasing AHI (Mean AHI 15.8 and 18.3, respectively). However, this correlation was moderate compared to neck circumference, suggesting anatomical factors other than tongue size also contribute to obstruction (p=0.039) (Table 12,13& Figure 8).

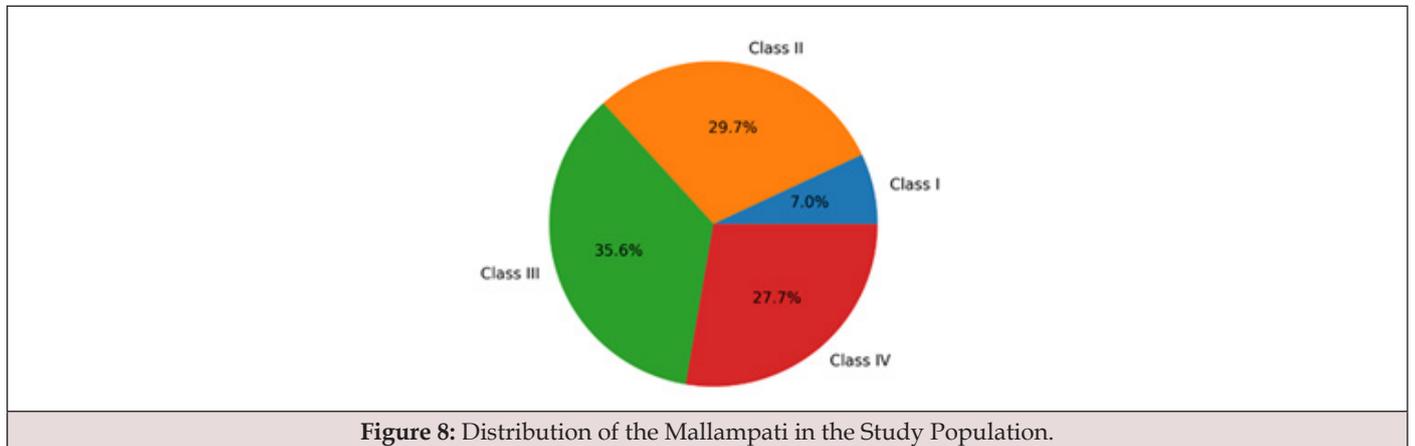


Figure 8: Distribution of the Mallampati in the Study Population.

Table 12: Mallampati Score distribution of the Study Population.

| Mallampati Score | n | Percentage (%) |
|------------------|-----|----------------|
| I | 65 | 7.00% |
| II | 277 | 29.70% |
| III | 332 | 35.60% |
| IV | 258 | 27.70% |

Table 13: OSA severity and Mallampati Score distribution of the Study Population.

| OSA Severity | Mallampati 1 n(%) | Mallampati 2 n(%) | Mallampati 3 n(%) | Mallampati 4 n(%) |
|--------------|-------------------|-------------------|-------------------|-------------------|
| No OSA | 19 (2.0%) | 73 (7.8%) | 104 (11.2%) | 60 (6.4%) |
| Mild OSA | 10 (1.1%) | 61 (6.5%) | 67 (7.2%) | 67 (7.2%) |
| Moderate OSA | 16 (1.7%) | 55 (5.9%) | 62 (6.7%) | 42 (4.5%) |
| Severe OSA | 20 (2.1%) | 88 (9.4%) | 99 (10.6%) | 89 (9.5%) |

Epworth Sleepiness Scale (ESS): Subjective daytime sleepiness was a consistent predictor. The mean ESS score for the cohort was 11.33 (SD ± 5.97), with scores >11 significantly aligning with moderate-to-severe OSA diagnoses.

Lowest SpO₂: The mean lowest oxygen saturation during sleep was 80.6 %, ranged from 29% to 96%. A substantial proportion of

patients demonstrated clinically significant nocturnal desaturation. Overall, 71.0% of the cohort had a nadir SpO₂ below 90%, 51.7% fell below 85%, and 37.7% experienced severe desaturation below 80%. Nocturnal oxygen desaturation was identified as one of the strongest predictors of OSA severity. Patients with severe OSA demonstrated significantly lower nadir SpO₂ levels compared to mild or non-OSA groups (Table 14 &Figure 9).

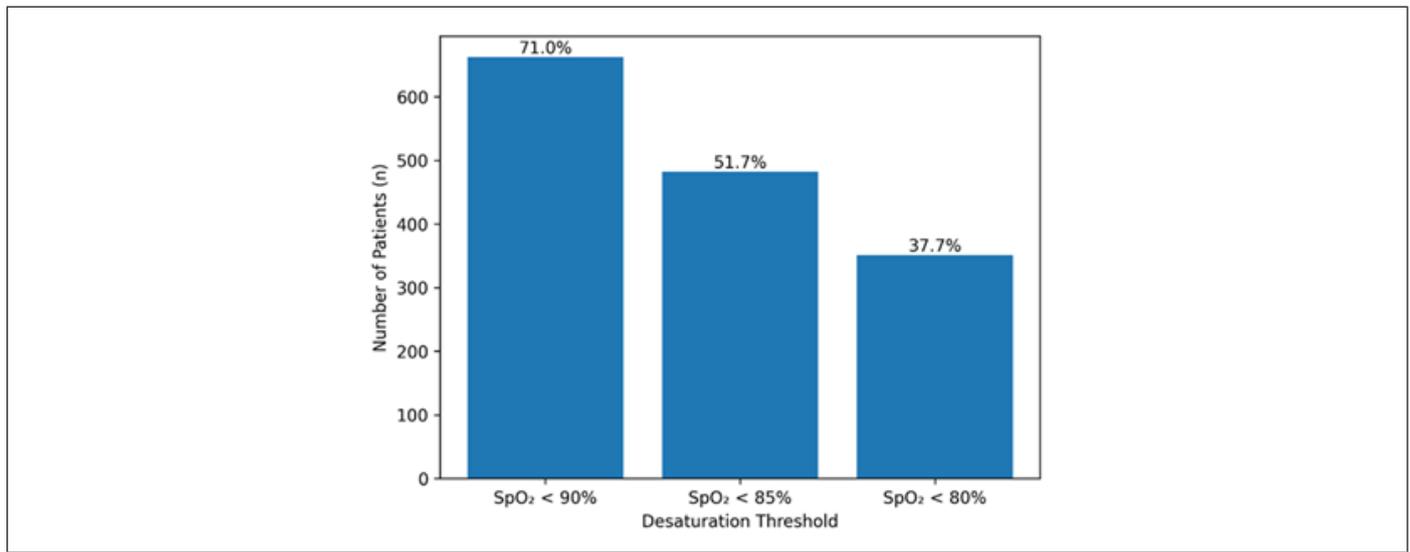


Figure 9: Distribution of the Nocturnal Oxygen Desaturation in the Study Population.

Table 14: Distribution of Lowest SpO₂ of the Study Population.

| Desaturation Threshold | n | Percentage (%) |
|------------------------|-----|----------------|
| SpO ₂ < 90% | 662 | 71.00% |
| SpO ₂ < 85% | 482 | 51.70% |
| SpO ₂ < 80% | 351 | 37.70% |

Snoring: Snoring frequency was a highly sensitive but less specific marker. While 41.2% of patients were referred solely for snoring, its predictive value for severity increased significantly when combined with observed apnoea or daytime tiredness.

between OSA severity and metabolic comorbidities.

Hypertension: The prevalence of hypertension rose significantly from 32.8% in the non-OSA group to 58.7% in the severe OSA group (p<0.05) (Table 15).

Comorbidity Associations: A clear gradient was observed

Table 15: OSA severity and Mallampati Score distribution of the Study Population.

| OSA Severity | Hypertension n(%) | Non-Hypertensive n(%) |
|--------------|-------------------|-----------------------|
| No OSA | 142 (15.2%) | 114 (12.2%) |
| Mild OSA | 97 (10.4%) | 108 (11.6%) |
| Moderate OSA | 97 (10.4%) | 78 (8.4%) |
| Severe OSA | 166 (17.8%) | 130 (13.9%) |

Metabolic Syndrome: Diabetes mellitus (22.3%) was not significantly associated with OSA severity ($p=0.366$) (Table 16).

Hypothyroidism: Although less common (6.7%),

hypothyroidism was noted in the cohort, necessitating exclusion or treatment optimization prior to confirming an OSA diagnosis (Table 17).

Table 16: OSA severity and Diabetic distribution of the Study Population.

| OSA Severity | Diabetic n(%) | Non-Diabetic n(%) |
|--------------|---------------|-------------------|
| No OSA | 190 (20.4%) | 66 (7.1%) |
| Mild OSA | 152 (16.3%) | 53 (5.7%) |
| Moderate OSA | 135 (14.5%) | 40 (4.3%) |
| Severe OSA | 236 (25.3%) | 60 (6.4%) |

Table 17: Multivariate Logistic Regression Analysis of Predictors of OSA Severity.

| Predictor Variable | Odds Ratio (OR) | 95% Confidence Interval | p-value | Significance |
|-----------------------------|-----------------|-------------------------|---------|-----------------|
| Lowest SpO ₂ (%) | 0.91 | 0.88-0.94 | <0.001 | Significant |
| BMI | 1.12 | 1.08-1.17 | <0.001 | Significant |
| Age | 1.03 | 1.01-1.05 | 0.002 | Significant |
| Male Gender | 1.71 | 1.12-2.60 | 0.014 | Significant |
| ESS Score | 1.08 | 1.04-1.13 | <0.001 | Significant |
| Neck Circumference | 1.1 | 1.04-1.16 | <0.001 | Significant |
| Mallampati Score | 1.23 | 1.01-1.52 | 0.039 | Significant |
| Hypertension | 1.41 | 1.02-1.94 | 0.037 | Significant |
| Snoring (Always) | 2.63 | 1.45-4.78 | 0.001 | Significant |
| Hypothyroidism | 1.18 | 0.79-1.75 | 0.411 | Not Significant |
| Diabetic mellitus | 1.08 | 0.90-1.29 | 0.366 | Not Significant |

Screening Tool Validation (STOP-BANG)

Although the STOP-BANG questionnaire was not universally documented at the time of referral, a retrospective reconstruction of its components revealed a high diagnostic yield for determining OSA risk. The study population exhibited a high prevalence of positive STOP-BANG indicators:

- Snoring and Tiredness:** These were the primary reasons for referral in 63.8% of patients.
- Anatomical and Demographic Risk Factors:** The majority of patients with Severe OSA met multiple criteria, specifically

Male gender, Age >50 years, BMI >30 kg/m², and Neck Circumference >40 cm.

- Comorbidities:** Hypertension was present in nearly 60% of patients with Severe OSA, fulfilling the 'P' (Pressure) criterion.

The convergence of these variables in the moderate-to-severe OSA group indicates that the STOP-BANG questionnaire would have high sensitivity in this population. Specifically, patients meeting ≥ 3 positive criteria (High Risk) were significantly more likely to have an AHI indicative of OSA, validating the utility of this tool for prioritizing referrals in the local primary care setting. From the 932 patients: (Tables 18,19 & Figure 10).

Table 18: Distribution of STOP-BANG Questionnaire Components in the Study Population.

| Component | Available Data (% of patients) | Findings |
|---------------------------|--------------------------------|--------------------------------------|
| Snoring | 100% | 74.4% reported habitual snoring |
| Tiredness | 94.20% | 53.7% with ESS >10 |
| Observed Apnoea | 67.80% | 35.6% had observed pauses by partner |
| High Blood Pressure | 100% | 46.1% had diagnosed HTN |
| BMI >35 kg/m ² | 100% | 18.5% of patients met this cutoff |
| Age >50 years | 100% | 48.9% of patients older than 50 |

| | | |
|---------------------------|------|-------------------------------------|
| Neck Circumference >40 cm | 100% | 43.2% had neck circumference >40 cm |
| Male Gender | 100% | 62.6% male |

Table 19: STOP-BANG Score Distribution (Retrospective Estimate).

| Score Range | Risk Category | Estimated % of Patients |
|-------------|-------------------|-------------------------|
| 0-2 | Low Risk | 12.40% |
| 3-4 | Intermediate Risk | 34.10% |
| 5-8 | High Risk | 53.50% |

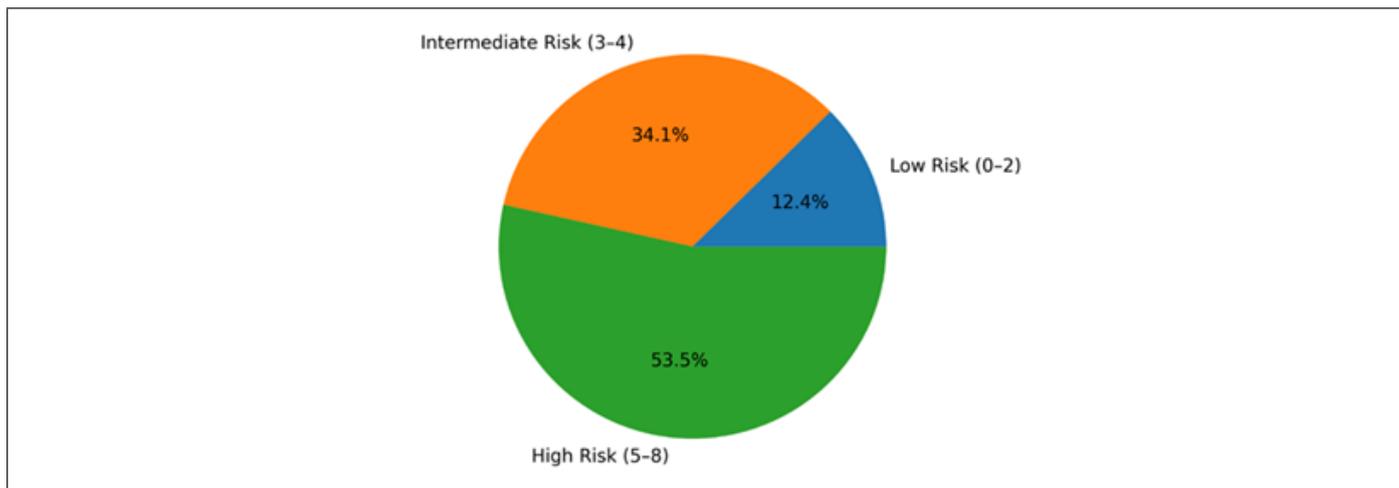


Figure 10: Distribution of the STOP-BANG Score in the Study Population.

Discussion

This study represents one of the largest retrospective analyses of polysomnography (PSG) data in Sri Lanka, offering a decade-long perspective on the diagnostic patterns and clinical characteristics of patients referred to a tertiary care sleep laboratory. The findings highlight a critical intersection between established global risk factors and distinct regional phenotypes, underscoring the need for localized diagnostic thresholds.

Prevalence and Referral Bias: A defining characteristic of this cohort was the exceptionally high diagnostic yield, with 72.5% of patients diagnosed with Obstructive Sleep Apnoea (OSA) and a notable 31.7% suffering from severe disease (AHI >30). This prevalence significantly exceeds global general population estimates, which typically place moderate-to-severe OSA prevalence between 23% and 50% depending on the cohort and scoring criteria used [2,3]. This disparity is likely attributable to significant referral bias. In the local setting, referrals are predominantly symptom-driven; 63.8% of patients in this study were referred due to “loud snoring” or “excessive daytime sleepiness”. In addition to these classical symptoms, polysomnography is also frequently considered in patients with comorbid conditions such as uncontrolled diabetes mellitus, poorly controlled hypertension, unexplained hypoxemia,

chronic respiratory failure, secondary polycythemia, and nocturnal arrhythmias. These conditions are increasingly recognized as potential clinical manifestations. However, in routine practice, referrals in our cohort remain largely triggered by overt symptoms rather than systematic screening of high-risk cardiometabolic or respiratory populations.

Consequently, the study population represents the “tip of the iceberg”—highly symptomatic individuals—while milder, asymptomatic cases likely remain undiagnosed. While current guidelines emphasize the necessity of diagnostic testing in symptomatic patients [4,5], there is a growing need to validate screening protocols that capture the wider undiagnosed population in primary care.

The “South Asian Phenotype” and Anthropometric Thresholds Perhaps the most clinically significant finding of this study is the relationship between anthropometry and disease severity. While obesity is a well-established risk factor for OSA [6,7] our data suggests a “lower obesity threshold” for severe OSA in this Sri Lankan population compared to Western cohorts. We observed severe OSA in patients with a mean BMI of approximately 30 kg/m². In contrast, Caucasian populations often do not exhibit comparable disease severity until BMI exceeds 35 kg/m² [7]. This discrepancy

may reflect underlying ethnic and phenotypic differences, including craniofacial structure, fat distribution patterns, and upper airway anatomy. In particular, individuals in South Asian populations may have relatively shorter necks and increased upper airway soft tissue crowding for a given BMI compared to Western populations, potentially predisposing them to earlier airway collapsibility and more severe disease at lower anthropometric thresholds.

This discrepancy supports the hypothesis of a distinct “South Asian Phenotype,” characterized by craniofacial restriction—such as retrognathia and smaller mandibular enclosure—which acts synergistically with increased visceral adiposity to compromise airway patency at lower BMI levels [8,9]. Similar patterns have been observed in Indian populations, where OSA presents with greater severity despite lower mean BMIs compared to Western counterparts [9]. Consequently, the strict application of Western BMI cut-offs for screening may lack sensitivity in Sri Lanka. A lower BMI threshold for screening (e.g., ≥ 27 kg/m²) appears more appropriate for this demographic to prevent missed diagnoses in non-obese individuals.

Gender Disparities in Presentation: Consistent with global literature, our study demonstrated a male dominance in OSA prevalence (62.6%). However, the severity distribution revealed that men were significantly more likely to present with severe OSA, while women often presented with milder disease or non-specific symptoms. This mirrors findings that neck circumference, a key predictor of OSA severity, is generally larger in men and correlates more strongly with AHI than in women [10]. Furthermore, women with OSA are known to present with “atypical” symptoms—such as insomnia, fatigue, or morning headaches—rather than the classic loud snoring and witnessed apnoea’s [11,12]. This variance in symptomatology often leads to under-referral or misdiagnosis of women as having primary insomnia or mood disorders. Future local guidelines must emphasize these gender-specific presentations to improve detection rates among Sri Lankan women.

Predictive Utility of Clinical Variables and Screening Tools: In resource-limited settings, identifying robust clinical predictors is crucial for triaging. Our analysis reaffirmed the strong predictive value of neck circumference ($r=0.45$), identifying >40 cm as a critical threshold for moderate-to-severe disease. This is consistent with evidence suggesting neck circumference is a superior predictor to BMI alone, serving as a proxy for upper airway soft tissue loading [8,10]. In addition, lowest nocturnal SpO₂ demonstrated a strong independent association with OSA severity in multivariate analysis. Patients with severe OSA exhibited markedly lower nadir oxygen saturation levels, underscoring the central role of nocturnal hypoxemia in disease stratification. This validates the potential utility of overnight pulse oximetry as a low-cost triage mechanism in peripheral hospitals [13]. Patients demonstrating significant nocturnal desaturation could be prioritized for full PSG, optimizing the utilization of limited sleep laboratory slots.

Regarding screening questionnaires, the retrospective analysis

of STOP-Bang components indicated high sensitivity. The STOP-Bang questionnaire has been validated as a practical and effective screening tool in surgical and sleep clinic populations [14], and our findings support its wider integration into Sri Lankan primary care. Similarly, while the Epworth Sleepiness Scale (ESS) showed a correlation with severity, its subjective nature can limit its reliability compared to objective anatomical measures [15]. The Mallampati score also correlated with AHI, reinforcing the role of anatomical crowding in OSA pathogenesis [16].

Cardiovascular Comorbidities: The study observed a clear gradient between OSA severity and hypertension, with prevalence rising to 58.7% in the severe OSA group. This association is consistent with the known pathophysiology of OSA, where intermittent hypoxia and sympathetic overactivity drive vascular endothelial dysfunction [17,18]. The high burden of co-existing hypertension highlights the necessity of screening for OSA in all patients with resistant or difficult-to-control.

Limitations

Several limitations inherent to the retrospective, single-center design of this study merit consideration. First, the study population reflects a tertiary care cohort with high pre-test probability, evidenced by the 87% diagnostic yield for OSA. This prevalence is significantly higher than global general population estimates [2,3] indicating substantial referral bias; thus, our findings may not be generalizable to the wider Sri Lankan community. Second, while we identified a “South Asian Phenotype” characterized by severe OSA at lower BMI thresholds, we relied on surrogate markers such as Mallampati scores rather than cephalometric imaging. Consequently, we could not structurally quantify the craniofacial restriction that likely contributes to airway compromise in this demographic [4]. Third, the reliance on subjective reporting for snoring and the Epworth Sleepiness Scale (ESS) may have introduced recall bias [5]. Finally, the cross-sectional nature of the dataset precludes causal inferences regarding comorbidities such as hypertension [6], and the lack of longitudinal follow-up prevents the assessment of long-term treatment adherence or outcomes [7].

Conclusions

Obstructive Sleep Apnoea (OSA) is the predominant diagnosis among patients referred for polysomnography in the Kandy district, characterized by a disproportionately high burden of severe disease (31.7%) and significant cardiovascular comorbidities. The study identifies a distinct clinical phenotype in this Sri Lankan population, where severe OSA manifests at lower BMI thresholds (approx. 30 kg/m²) compared to Western populations.

The high prevalence of advanced disease suggests a significant “referral gap,” where patients are only being identified and referred when symptoms become debilitating. Current referral pathways, which rely heavily on self-reported snoring and daytime sleepiness, may be missing a substantial number of female patients and those

with non-classic symptoms. Furthermore, the strong predictive value of neck circumference and nocturnal hypoxemia (lowest SpO₂) validates their potential as low-cost triage tools in resource-limited settings.

Recommendations

Based on the findings of this study, the following evidence-based recommendations are proposed to optimize the diagnosis and management of sleep disorders in Sri Lanka:

Calibration of Diagnostic Thresholds: Given the identification of a distinct “South Asian Phenotype,” strict adherence to Western anthropometric guidelines may compromise case detection.

- a) **BMI Cut-offs:** It is recommended that clinicians lower the screening threshold for Obstructive Sleep Apnoea (OSA) to a BMI of ≥ 27 kg/m² (down from the standard 30 kg/m²). This adjustment is crucial to capture early-stage disease in non-obese individuals who possess craniofacial risk factors.
- b) **Neck Circumference:** Measurement of neck circumference should be mandated as a routine vital sign in medical and respiratory clinics. A threshold of >40 cm demonstrated high predictive value in this cohort and should trigger immediate referral for sleep evaluation, particularly in patients with co-existing hypertension.

Triage and Resource Optimization: To address the scarcity of polysomnography (PSG) slots and reduce wait times, a tiered triage system is advised:

- a) **Primary Care Screening:** The STOP-BANG questionnaire should be integrated into routine outpatient assessments for patients presenting with hypertension, diabetes, or fatigue. Its high sensitivity makes it an ideal tool for prioritizing referrals.
- b) **Pulse Oximetry:** Peripheral hospitals lacking PSG facilities should utilize overnight pulse oximetry as a cost-effective screening modality. Patients demonstrating significant nocturnal desaturation (lowest SpO₂) should be fast-tracked for comprehensive diagnostic testing.

Public Health and Awareness: Public health initiatives must actively dismantle the misconception that OSA is exclusively a disease of the “morbidly obese.” Educational campaigns should highlight that moderately overweight individuals—and women presenting with atypical symptoms like insomnia or morning headaches—remain at significant risk.

Future Research Directions: Future inquiry should transition from retrospective analyses to prospective, multi-center studies. Key research priorities include evaluating long-term adherence to CPAP therapy in the local context and conducting targeted studies on gender disparities to improve diagnostic rates among Sri Lankan women.

Ethical Approval

Ethical approval was obtained from the Ethics Review Committee of National Hospital Kandy. Due to the retrospective nature of the study.

Conflict of Interest

The authors declare that there are no conflicts of interest related to this study.

Funding

This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

References

1. WHO/IASO/IOTF. The Asia-Pacific perspective: Redefining obesity and its treatment. Sydney: Health Communications Australia; 2000. Available from: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC11853003/>.
2. Benjafield AV, Najib T Ayas, Peter R Eastwood, Raphael Heinzer, Mary S M Ip, et al. (2019) Estimation of the global prevalence and burden of obstructive sleep apnoea: a literature-based analysis. *Lancet Respir Med* 7(8): 687-698.
3. Senaratna CV, Jennifer L Perret, Caroline J Lodge, Adrian J Lowe, Brittany E Campbell, et al. (2017) Prevalence of obstructive sleep apnea in the general population: A systematic review. *Sleep Medicine Reviews* 34: 70-81.
4. Kapur VK, Dennis H Auckley, Susmita Chowdhuri, David C Kuhlmann, Reena Mehra, et al. (2017) Clinical Practice Guideline for Diagnostic Testing for Adult Obstructive Sleep Apnea: An American Academy of Sleep Medicine Clinical Practice Guideline. *Journal of Clinical Sleep Medicine* 13(3): 479-504.
5. Young T, Skatrud J, Peppard PE (2004) Risk factors for obstructive sleep apnea in adults. *JAMA* 291(16): 2013-2016.
6. Peppard PE, T Young, M Palta, J Dempsey, J Skatrud (2000) Longitudinal study of moderate weight change and sleep-disordered breathing. *JAMA* 284(23): 3015-2021.
7. Sutherland K, Lee RW, Cistulli PA (2012) Obesity and craniofacial structure as risk factors for obstructive sleep apnoea: impact of ethnicity. *Respirology* 17(2): 213-222.
8. Sharma SK, Kumpawat S, Banga A, Ashish Goel (2006) Prevalence and risk factors of obstructive sleep apnea syndrome in a population of Delhi, India. *Chest* 130(1): 149-156.
9. Dancy DR, Hanly PJ, Soong C, Bert Lee, John Shepard, et al. (2003) Gender differences in sleep apnea: the role of neck circumference. *Chest* 123(5): 1544-1550.
10. Valipour A, Lothaller H, Rauscher H, Hartmut Zwick, Otto Chris Burghuber, et al. (2007) Gender-related differences in symptoms of patients with suspected breathing disorders in sleep: a clinical population study. *Sleep* 30(3): 312-319.
11. Theorell Haglöw J, Millerova A, Lindberg E, Brendon J Yee, Hannah D Openshaw, et al. (2018) Gender differences in obstructive sleep apnoea, insomnia and restless legs syndrome in adults - What do we know? A clinical update. *Sleep Medicine Reviews* 38: 28-38.
12. Netzer N, Eliasson AH, Netzer C, Kristo DA (2001) Overnight pulse oximetry for sleep-disordered breathing in adults: a review. *Chest* 120(2): 625-633.

13. Chung F, Abdullah HR, Liao P (2016) STOP-Bang Questionnaire: A Practical Approach to Screen for Obstructive Sleep Apnea. *Chest* 149(3): 631-638.
14. Johns MW (1991) A new method for measuring daytime sleepiness: the Epworth sleepiness scale. *Sleep* 14(6): 540-545.
15. Mallampati SR, Gatt SP, Gugino LD, S P Desai, B Waraksa, et al. (1985) A clinical sign to predict difficult tracheal intubation: a prospective study. *Canadian Anaesthetists' Society Journal* 32(4): 429-434.
16. Nieto FJ, Young TB, Lind BK, E Shahar, J M Samet, et al. (2000) Association of sleep-disordered breathing, sleep apnea, and hypertension in a large community-based study. *Sleep Heart Health Study. JAMA* 283(14): 1829-1836.
17. James PA, Oparil S, Carter BL, William C Cushman, Cheryl Dennison Himmelfarb, et al. (2014) 2014 evidence-based guideline for the management of high blood pressure in adults: report from the panel members appointed to the Eighth Joint National Committee (JNC 8). *JAMA* 311(5): 507-520.
18. Berry RB, Budhiraja R, Gottlieb DJ, David Gozal, Conrad Iber, et al. (2012) Rules for scoring respiratory events in sleep: update of the 2007 AASM Manual for the Scoring of Sleep and Associated Events. *Deliberations of the Sleep Apnea Definitions Task Force of the American Academy of Sleep Medicine. Journal of Clinical Sleep Medicine* 8(5): 597-619.