Clinical Predictors of Sleep Apnea-Hypopnea Syndrome Susceptible to Treatment With Continuous Positive Airway Pressure

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OBJECTIVE: To analyze the predictive value of clinical data for identifying patients suspected of sleep apnea-hypopnea syndrome with an apnea-hypopnea index (AHI) ≥30.

MATERIAL AND METHODS: Patient characteristics, cardiorespiratory medical history, and clinical signs and symptoms were recorded for all patients. Exclusion criteria were daytime respiratory insufficiency or heart failure. All patients underwent polysomnographic testing (AutoSet Portable Plus II, ResMed Corp, Sydney, Australia) for automatic AHI calculation and manual determination of central and obstructive apneas. A logistic regression model was constructed to calculate the likelihood of an individual’s presenting an AHI ≥30 as well as the predictive value of each variable and of the final model.

RESULTS: Three hundred twenty-nine patients with a mean (SD) age of 58 (13.45) years were studied; 76.4% were men. Data for 207 patients were used to construct the logistic regression model: logit (P) = 2.5 hypertension + 1.5 Epworth test + body mass index + 0.6 repeated observed episodes of apnea – 2.1. Logit(P) = log((1-P)/P) and variables were dichotomized with cut points of 11 for the Epworth test and of 30 kg/m² for body mass index. The diagnostic sensitivity of the model was 80.2% (75%-86%); specificity was 93.4% (89%-95%); positive predictive value was 86.9% (84%-93%); and negative predictive value was 89.6% (81%-90%), such that 89.6% of the patients were correctly classified. The variable with the greatest predictive value was high blood pressure. The model was validated prospectively in the remaining 102 patients.

CONCLUSIONS: Prior to diagnostic tests for sleep apnea-hypopnea syndrome, clinical data can be useful for identifying patients suspected to have a AHI ≥30.

Key words: Apnea-hypopnea syndrome. Sleep. Logistic regression. AutoSet®.

Valor predictivo de la clínica para la identificación de los pacientes con síndrome de apneas-hipopneas durante el sueño susceptibles de tratamiento con presión positiva continua de la vía aérea (CPAP)

OBJETIVO: Analizar el valor predictivo de las variables clínicas en la identificación de pacientes con sospecha de síndrome de apneas-hipopneas durante el sueño (SAHS) con un índice de apneas-hipopneas (IAH) superior a 30.

MATERIAL Y MÉTODOS: Se recogieron datos referentes a variables generales, antropométricas, antecedentes personales cardiorespiratorios, clínica y la sensación subjetiva del clínico. Se excluyó a los pacientes con insuficiencia respiratoria diurna o cardíaca. A todos ellos se les realizó un estudio poligrafico (AutoSet) con determinación automática del IAH y manual del índice de apneas obstructivas y centrales. Mediante la construcción de un modelo lógistico se calculó la probabilidad individual de presentar un IAH ≥30 así como el valor predictivo de cada variable estudiada por separado y de la ecuación logística final.

RESULTADOS: Se estudió a 329 pacientes, con una edad media ± desviación estándar de 58 ± 13,45 años; el 76,4% eran varones. Las variables de 207 pacientes se utilizaron para la construcción de la ecuación logística: logit P = 2,5 hipertensión arterial + 1,5 test de Epworth + índice de masa corporal + 0,6 apneas presenciadas y repetidas – 2,1; siendo logit P = log((1-P)/P) y valorando las variables como dicotómicas con puntos de corte para el test de Epworth de 11 y para el índice de masa corporal de 30 kg/m². El valor diagnóstico de dicha ecuación fue: sensibilidad del 80,2% (75%-86%); especificidad del 93,4% (89%-95%); valor predictivo positivo del 86,9% (84%-93%) y valor predictivo negativo del 89,6% (81%-90%), lo que supuso un porcentaje de pacientes correctamente clasificados del 89,6%. La variable que presentó mayor capacidad predictora fue la presencia de hipertensión arterial. La ecuación se validó prospectivamente en los restantes 102 pacientes.

CONCLUSIONES: Los parámetros clínicos podrían ser útiles en la identificación, previa a la realización del estudio diagnóstico de SAHS, de aquellos pacientes con sospecha de SAHS que presentaran un IAH ≥ 30.

Palabras clave: Síndrome de apneas-hipopneas. Sueño. Regresión logística. AutoSet
Introduction

Sleep apnea-hypopnea syndrome (SAHS) constitutes a recognized public health problem both because of its high prevalence in the general population and the morbidity and mortality it causes.¹ If we consider an apnea-hypopnea index (AHI) over 10 together with the presence of excessive daytime sleepiness to indicate a diagnosis of SAHS, then the prevalence of SAHS among the middle-aged in Spain is estimated to be approximately 3% to 3.5%.²

SAHS should be diagnosed by polysomnography (PSG), although a valid diagnosis can be established by respiratory polygraphy that has been properly validated for populations with high or low probability of the diagnosis.³,⁴ Nevertheless, diagnosis is usually delayed significantly because the few sleep laboratories that are available are working at capacity.³ Considering the demonstrated relation between SAHS and a 2- to 7-fold greater likelihood of a patient having a traffic accident,⁵,⁶ an increased risk of cardiovascular disease or related death,⁷,⁸ and the great efficacy of continuous positive airway pressure (CPAP) treatment on the main symptoms,¹²,¹⁵ the search for alternative diagnostic approaches would seem to be a priority, particularly in the effort to identify the most severe forms of the disease and initiate early treatment until a sleep study can confirm the diagnosis.

Therefore, various suggestions—from subjective clinical assessment¹⁴ to the application of clinical,¹⁵–²¹ functional,²² or anthropometric²³ parameters—have been put forth for identifying a priori the likelihood that a patient has SAHS or a certain AHI. Among the range of options, the ones most often studied have been clinical parameters. Several studies have evaluated their role as diagnostic tools through the creation of predictive models using multivariate analysis.¹⁴,¹⁵,¹⁸,²¹ Results have varied, although the models generally have high sensitivity (between 78% and 95%) and low specificity (between 41% and 63%) for AHI cut points between 5 and 20 and different prevalences of SAHS in the studied population.²⁴

For Spain, the Spanish Society of Pulmonology and Thoracic Surgery (SEPAR) has issued a series of recommendations for treating SAHS patients, establishing an arbitrary AHI threshold of 30 to distinguish patients who, depending on their symptoms and cardiovascular history, will be most likely to respond to CPAP treatment.¹² Accordingly, we believe that predicting which patients are likely to have an AHI ≥30 would have useful therapeutic application and allow CPAP treatment to reach the most severe cases early in the disease (provided clinical criteria and medical history are sufficient to warrant prescription), while waiting for tests to confirm the diagnosis. At least such an approach would allow such patients to have priority when scheduling tests. We have not found any studies in the Spanish literature on the diagnostic value of clinical parameters for predicting an AHI ≥30 in patients referred for specialist consultation. Therefore, the present study was designed to analyze the predictive value of such parameters relative to an AHI cut point of ≥30.

Material and Methods

All patients referred to our service with a suspected diagnosis of SAHS from January 2001 through August 2002 were studied. Our respiratory medicine department is part of a first-referral regional hospital that provides specialist care to a population of 60 000. SAHS was suspected if 1 of 3 cardinal symptoms was reported: chronic snoring, excessive daytime sleepiness, or observed apneas. Patients with daytime respiratory insufficiency or congestive heart failure were excluded. All patients were given a polygraph test using the AutoSet® (AS) Portable Plus II (ResMed Corp, Sydney, Australia). When the AS auto-CPAP device is set in diagnostic mode, various respiratory variables and heart rate can be recorded. Nasal airflow is measured by a cannula with a pressure transducer and oxygen saturation by a digital pulse oximeter, apneas are counted according to the patient’s position by a body position sensor and thoracoabdominal movements are recorded by way of signals from an elastic band with a piezoelectric sensor. Automatically, using appropriate software (Autoview 98, version 2.0), the AS calculates the AHI as well as the apnea index and the hypopnea index by subtraction of each of the previous 2 variables. Although the AS does not permit the total apnea index to be changed, each apnea can be classified manually as obstructive, mixed, or central with information from recordings of respiratory effort provided by the thoracoabdominal band. A respiratory event was defined as apnea when nasal airflow fell more than 75% and as hypopnea when it fell between 50% and 75%, for longer than 10 seconds in each case. The AHI was defined as the number of respiratory events (apneas or hypopneas) per recording hour. All data were calculated in function of total recording time. All tests were performed in dedicated hospital rooms prepared by trained personnel. Patient characteristics (age and sex), anthropometric data (body mass index [BMI] in kg/m² and neck circumference in centimeters), medical history (mainly cardiorespiratory signs such as hypertension, cardiac or cerebrovascular events, bronchial asthma, and chronic obstructive pulmonary disease), signs and symptoms (daytime sleepiness by a validated Spanish language version of the Epworth test,²⁵ the existence of observed apneas and their frequency, and the occurrence of asphyxia), and the referring caregiver’s subjective feeling (dichotomized) as to each patient’s probability of having an AHI ≥30. A diagnosis of hypertension was established according to the recommendations of the World Health Organization.²⁶ The morning after the polygraph test, the patient filled in a form about his or her subjective feeling about the amount (in hours) and quality (good–average–bad) of sleep. Tests were considered valid if the patient reported having had at least 3 hours with a minimum sleep quality estimated as average. Tests were considered invalid if there was a technical failure or if the patient had disconnected the device and recording had not lasted at least 3 hours. In both cases, the polygraph was repeated. SAHS was diagnosed if the AHI was ≥10.

Statistical Analysis

The commercial statistics software packet SPSS 9.0 (SPSS Inc., Chicago, Illinois, USA) was used. Quantitative variables were reported as means (SD) and qualitative variables as
Results

The number of patients initially enrolled was 329. Patients were excluded if they had daytime respiratory insufficiency (n=10), congestive heart failure (n=3), declined to participate (n=5), or died before the study took place (n=2). Therefore, 309 patients (76.4% men) entered into analysis. Their mean (SD) age was 58(13.45) years (range 24-83 years). Seventy-three percent were referred from primary care, 15% came from an otorhinolaryngologist, and 12% from a variety of internal medicine specialists. Data from 207 patients were analyzed retrospectively to construct a logistic regression model and the resulting equation was validated prospectively with data from the remaining 102 patients. No significant differences were found between the patient characteristics for the two groups, as shown in Table 1.

Bivariate analysis identified variables that were candidates for inclusion in the model from data available for the set of 207 patients initially analyzed (Table 2). BMI, the presence of observed and repeated apneas, the presence of hypertension, subjective clinical suspicion, Epworth test score, and the occurrence of asphyxia were significantly more frequent or higher in group 1 patients (AHI ≥30). To convert quantitative to qualitative variables, the cut points that best distinguished between groups 1 and 2 were a BMI ≥30 and an Epworth test score ≥11. The diagnostic values of individual variables entered into the model are shown in Table 3.

The best regression equation (n=207) was as follows:

\[
\logit P = 2.5 \times HT + 1.5 \times Epw + \text{BMI} + 0.6 \times \text{Apr} - 2.1
\]

where \(\logit P = \log(1-p)/p\), HT is the presence (1) or absence (0) of hypertension, Epw is an Epworth test

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Patient Characteristics for the Group Used To Create the Predictive Model and for the Group Used to Validate the Model Prospectively*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Variable</td>
<td>Regression Model Group (n=207)</td>
</tr>
<tr>
<td>Age, years</td>
<td>58(10.1)</td>
</tr>
<tr>
<td>Sex, male</td>
<td>85%</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>33(4.65)</td>
</tr>
<tr>
<td>Neck circumference, centimeters</td>
<td>42.2(3.7)</td>
</tr>
<tr>
<td>Asthma</td>
<td>9%</td>
</tr>
<tr>
<td>COPD</td>
<td>20(23%)</td>
</tr>
<tr>
<td>Hypertension</td>
<td>61(68.5%)</td>
</tr>
<tr>
<td>Stroke</td>
<td>9(9%)</td>
</tr>
<tr>
<td>Ischemic heart disease</td>
<td>6(6.7%)</td>
</tr>
<tr>
<td>Chronic snoring</td>
<td>18(96.4%)</td>
</tr>
<tr>
<td>Apneas†</td>
<td>60(60%)</td>
</tr>
<tr>
<td>Epworth test</td>
<td>12(3.6)</td>
</tr>
<tr>
<td>Asphyxia</td>
<td>30(34%)</td>
</tr>
</tbody>
</table>

*Quantitative values are means (SD). Qualitative variables are absolute numbers (percentages). AHI indicates apnea-hypopnea index; BMI, body mass index; COPD, chronic obstructive pulmonary disease; HT, presence (1) or absence (0) of hypertension; Epw, Epworth test.
assigning priority to such patients when scheduling tests. Waiting for PSG to confirm the diagnosis or at least for useful for making early treatment decisions while identifying those with an AHI diagnoses of SAHS had high predictive value for respiratory medicine specialists with suspected

Discussion

the group used to validate the model.

observed in the results for the group from whose data (87.1%, and gain in correctly classified patients 48.9% (95% CI 76%-91%), percentage correctly classified (95% CI 79%-91%), specificity 93.4% (95% CI, 89%-95%), PPV 89.6% (95% CI, 84%-93%), and NPV 86.9% (95% CI, 81%-90%). The percentage of correctly classified patients was 87.9%, meaning there were 11 false positives and 14 false negatives. The false positives had significantly higher Epworth scores than the rest of the patients [15](3) vs 8(3), P<.001, whereas there were significantly more hypertensive patients among the false negatives (79% vs 44.6%; P<.008). Therefore, if the pretest probability of correctly classifying an individual (prevalence of patients with an AHI ≥30 on the polysomnographic study) was 43%, the posttest probability (after applying the logistic regression model) was 89.6%, indicating a 46.6% gain in correctly classified patients (P>.0001).

The following results were obtained when the model was applied prospectively (n=102): sensitivity 83.1% (95% CI 79%-91%), specificity 91.1% (95% CI 85%-96%), PPV 87.1% (95% CI 84%-95%), NPV 84.5% (95% CI 76%-91%), percentage correctly classified 87.3%, pretest probability 38.2%, posttest probability 87.1%, and gain in correctly classified patients 48.9% (P>.0001). There were no significant differences observed in the results for the group from whose data the logistic regression model was derived and those for the group used to validate the model.

Discussion

Clinical parameters for patients referred to the respiratory medicine specialists with suspected diagnoses of SAHS had high predictive value for identifying those with an AHI ≥30. This finding may be useful for making early treatment decisions while waiting for PSG to confirm the diagnosis or at least for assigning priority to such patients when scheduling tests.

Several studies have sought to find a diagnostic procedure for identifying patients with SAHS or for predicting various AHIs before PSG, as part of an effort to avoid more expensive and less readily available diagnostic tests as well as to initiate early CPAP treatment under the assumption that patients usually face fairly long waiting lists.14-23

Among such studies have been those using unusual lung function parameters,22 measures of upper airway structures,23 or calculations performed with complex neural network computer programs.19 All have been shown to have considerable diagnostic value for identifying SAHS patients but little practical clinical utility given their complexity or lack of availability.

The most often studied parameters have been the clinical signs and symptoms that are easiest to see and measure. Studied individually, such clinical variables have not had acceptable predictive value for diagnosing SAHS.18 Only neck circumference has demonstrated a certain degree of predictive value in some studies,27 although some authors conclude that that measurement may combine linearly with other variables such as age, sex, or BMI and, therefore, would provide redundant information.15-18 We found no significant differences, however, between neck circumference in group 1 (AHI ≥30) and group 2 (AHI <30) patients. The reported predictive value of this variable may only appear when lower cutoff points are used (<20) and may lose its power to discriminate when disease is more severe. Other clinical variables such as the presence of

### Table 3

Diagnostic Value of Each Variable (Assessed as Dichotomies) Initially Considered for Constructing the Logistic Regression Model (Minimum Significance for Between-Group Comparison of .10)*

<table>
<thead>
<tr>
<th>Variable</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
<th>CorrCl</th>
<th>OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypertension</td>
<td>66.7</td>
<td>68.4</td>
<td>54</td>
<td>81.6</td>
<td>67.9</td>
<td>4.33 (2.17-8.64)</td>
</tr>
<tr>
<td>Apneas†</td>
<td>58.6</td>
<td>71.4</td>
<td>57.1</td>
<td>69.8</td>
<td>64.8</td>
<td>3.27 (1.63-6.56)</td>
</tr>
<tr>
<td>SubOb</td>
<td>42.9</td>
<td>57.1</td>
<td>75</td>
<td>25</td>
<td>53.6</td>
<td>1.25 (0.48-2.01)</td>
</tr>
<tr>
<td>Asphyxia</td>
<td>60</td>
<td>56.3</td>
<td>46.2</td>
<td>60</td>
<td>53.6</td>
<td>1.28 (0.70-2.37)</td>
</tr>
<tr>
<td>BMI ≥30 kg/m²</td>
<td>61.1</td>
<td>68.8</td>
<td>59.5</td>
<td>70.2</td>
<td>65.5</td>
<td>3.46 (1.82-6.56)</td>
</tr>
<tr>
<td>Test de Epworth ≥11†</td>
<td>63.6</td>
<td>70.6</td>
<td>58.3</td>
<td>75</td>
<td>67.90</td>
<td>4.20 (2.17-8.11)</td>
</tr>
</tbody>
</table>

*Values are expressed as percentages with the exception of the OR. PPV indicates positive predictive value; NPV, negative predictive value; SubOb, subjective clinical observation to classify patients in either group 1 or group 2; BMI, body mass index.

†Apneas observed and repeated.

‡Variables entering the final equation.

### Table 4

Logistic Regression Model for Prediction Using an AHI Cutoff of 30. Adjusted Odds Ratio and 95% Confidence Interval for Each Variable in the Model

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR</th>
<th>95% CI of the OR</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>HT</td>
<td>11.9</td>
<td>3.9-36.8</td>
<td>&lt;.0001</td>
</tr>
<tr>
<td>Epw ≥11</td>
<td>4.47</td>
<td>2.2-11.4</td>
<td>.002</td>
</tr>
<tr>
<td>BMI ≥30 kg/m²</td>
<td>2.92</td>
<td>1.3-7.1</td>
<td>.01</td>
</tr>
<tr>
<td>Apr</td>
<td>1.73</td>
<td>1.1-4.2</td>
<td>.045</td>
</tr>
</tbody>
</table>

OR indicates odds ratio; CI, confidence interval; HT, presence or absence of hypertension; Epw, Epworth score ≥11; BMI, body mass index; Apr, presence or absence of apneas that are observed and repeated.
hypertension, observed apneas, BMI or excessive
daytime sleepiness have been reported to have modest
diagnostic value when studied individually using an
AHI cutoff between 5 and 20, usually because those
variables have low NPVs. Deegan et al. however,
found that although clinical variables studied
individually have low NPVs and sensitivities, but very
high PPVs and specificities, at low AHI cutoffs (≥10),
NPV and specificity increase considerably while PPV
and specificity decrease only moderately as higher
cutoffs are chosen (≥20), with a consequent increase in
overall diagnostic value. Our results suggest that the
aforementioned variables have better-than-average
value for distinguishing patients with an AHI ≥30, with
correct diagnoses exceeding 65% in most cases.
Nevertheless, because this improvement in diagnostic
capability for high AHI cut points is still modest, the
clinical application for individual variables is still
scarce. Finally, the clinician’s subjective guess about
the diagnosis did not have predictive value; in other
studies, as in our study, the percentage of correct
diagnoses exceeding 65% in most cases. As a result,
various combinations of clinical variables have been
used in regression models to try to predict the
presence of SAHS for different AHI cut points (usually
between 5 and 20) in patients referred to sleep
clinics. Results have varied depending mainly
on the probability of having SAHS based on symptoms
and on the AHI cut point used for diagnosis, although
sensitivity has usually been high (>85%) and specificity
low (<55%) for AHI cutoffs between 5 and 20. As
such results, these equations may have value for ruling
out the diagnosis but not for confirming it or for
supporting early treatment.

The logistic regression model from our study showed
excellent ability to predict which patients would have
an AHI ≥30. The equation includes 4 variables typical
of predictive models published to date: the presence of
hypertension, the presence of observed and repeated
apneas, the Epworth test score, and BMI. All of them
were dichotomized, and the last two are relevant to
cutoffs of 11 and 30, respectively. The OR for each
variable seems to indicate that using higher than usual
AHI cutoffs leads to a significant change in the relative
weight of each variable’s predictive value, the greatest
changes occurring for hypertension (OR=11.9) and a
high Epworth test score (OR=4.47) as opposed to age,
sex, presence of apneas, or anthropometric variables
(neck circumference or BMI), although there is no
change in which variables finally enter the model. It
is important to point out that the presence of apneas only
had predictive value when the sleeping partner indicated
that they were repeated. It seems logical to think
that most snorers experience apneic events normally
and even that a few are pathological. The sleeping partner
becomes aware of such events and reports them
faithfully, even when apneas are not repeated often
enough to define a high AHI. This situation can lead to
overestimating the existence of isolated nighttime
apneas. Our study would therefore not apply to subjects
without companions who can become aware of the
existence of such apneic events, for example to
individuals who live or sleep alone (12% in our patient
series).

Of the 25 patients (12.1%) who were not correctly
classified by the model, 11 were false positives and 14
were false negatives. A careful look at these patients
indicates that the false positives were different from the
other patients in having very high Epworth test scores
(over 15). All had been referred for PSG because of
excessive daytime sleepiness in spite of having a
negative AS because of the “relatively” low NPV of the
AS (78%) in comparison with that of PSG in patients
similar to those in our series. Three had increased upper
airway resistance syndrome and were finally treated
with CPAP. 4 had SAHS (with AHI findings of 19, 22,
33, and 29), and the remaining 4 had negative PSG
findings and are undergoing tests to investigate the
reason for pathological daytime hypersomnia. The false
negatives were mostly hypertensive individuals.
Hypertension in our study was not actively investigated
but was recognized in the medical history.

The AS polygraphic study used instead of PSG
assessment is logically of limited diagnostic value
according to our study. However, it is important to point
out that this device is widely validated in the literature
for different cutoff points and prevalences of SAHS.

The reasons for our model’s high diagnostic and
predictive ability are complex. The explanation for the
higher overall value of the model may lie in 2 features
of our study: the high AHI selected as the cut point and
the high pretest probability. For none of the variables in
the final model was the sensitivity low; rather they all
had moderate sensitivities, between 50% and 67%. If
each variable is considered an individual diagnostic test,
the use of several alongside one another to classify
patients (as occurs in the use of predictive equations)
would increase sensitivity and NPV considerably. The
parallel decrease in specificity and PPV that would
 correspond to the increase in sensitivity might be
compensated for, in the case of specificity, by the high
cut point chosen to classify the patients and, in the case
of PPV, by the high pretest probability for that cut point
in our series. Finally, the high specificity values and
PPV for the individual variables in the equation may
influence the behavior of the model. Therefore, the
diagnostic value of our model may change if it is
applied to different patient populations.

In conclusion, we think that clinical parameters may
have considerable predictive value for distinguishing
patients with an AHI ≥30 among those referred to a
respiratory medicine specialist, allowing the eventual
mention of such parameters in SEPAR recommendations
for the early treatment of SAHS. Such
inclusion may save considerable time in initiating CPAP
for patients who are most ill or may serve
to give priority to severely ill patients when scheduling
diagnostic tests.
REFERENCES