Gastroesophageal reflux episodes in asthmatic patients and their temporal relation with sleep architecture

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Gastroesophageal reflux (GER) is common in asthma patients and can contribute to sleep disruption. The aim of the present study was to determine the time-related distribution of GER events together with their impact on sleep in asthmatic subjects with GER disease symptoms. The inclusion criteria were: 18–65 years, controlled moderate to severe asthma and GER-compatible clinical evidence. The exclusion criteria were: chronic obstructive lung disease, smoking, infections of the upper airways, use of oral corticosteroids, other co-morbidities, pregnancy, sleep-related disorders, night-time shift work, and the use of substances with impact on sleep. Asthmatic patients with nocturnal symptoms were excluded. All-night polysomnography and esophageal pH monitoring were recorded simultaneously. Of the 147 subjects selected, 31 patients and 31 controls were included. Seventeen patients were classified as DeMeester positive and 14 as DeMeester negative. Both groups displayed similar outcomes when general variables were considered. Sleep stage modification one minute prior to GER was observed in the DeMeester-positive group. Awakening was the most frequent occurrence at GER onset and during the 1-min period preceding 38% of the nocturnal GER. Sleep stage 2 was also prevalent and preceded 36% of GER events. In the DeMeester-negative group, awakening was the most frequent response before and during GER. Modifications in sleep stages, arousals or awakenings were associated with 75% of the total GER events analyzed during the period of one minute before and after the fall of esophageal pH below 4 in the DeMeester-positive group. These data provide evidence that sleep modifications precede the GER events in asthmatic patients.

Key words: Asthma; Gastroesophageal reflux; Polysomnography; Sleep

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Introduction

Occasional regurgitation and heartburn due to gastro-esophageal reflux (GER) are frequent in the general population. GER disease refers to the presence of symptoms which are secondary to the reflux of gastric content through the esophagus with or without signs of esophageal mucosa lesions (1). In more severe manifestations, GER seems to be related to recurrent and long-lasting reflux episodes which occur even during sleep (2). The increase in the transdiaphragmatic pressure, which occurs in the initial phases of sleep, may predispose patients to GER (3). The literature also highlights a greater frequency of GER occurring during arousals and the awakening period, but being markedly absent during rapid eye movement (REM) sleep (4,5).

Approximately 50 to 70% of asthmatic subjects regularly display GER symptoms (6-9). Esophageal mobility alterations occur in 82% of cases due to longer periods of contact between the esophageal mucosa and gastric acid...
Sleep abnormalities frequently detected in these subjects may not only result from asthma aggravation during the night (11,12) but may also stem from GER events (10,13) as reported by Goodall et al. (6) in a double-blind crossover study of 20 asthmatic patients. In that particular study, the investigators reported the beneficial effect of cimetidine on the night symptoms of asthma and GER. Additionally, reproduction of the alterations in nighttime breathing patterns by infusing acid solution into the distal esophagus of asthmatic patients with esophagitis has been reported (14,15). However, the relationship between asthma, GER and disordered sleep is not understood. In a controlled study of nine patients with asthma, Hughes et al. (4) failed to find a higher incidence of GER or any impact of the latter on respiratory and sleep parameters, since GER episodes occurred mainly during arousals or waking periods. Similarly, in a study based on simultaneous and continuous recording of pH and lower airway resistance, Tan et al. (16) failed to identify any impact of GER on day- or night-time asthma attacks.

Assuming that, before their detection, GER events can contribute to sleep disruption in asthma, the present study was conducted with the aim of determining the time-related distribution of GER events together with their impact on sleep in asthmatic subjects.

Patients and Methods

Patients

From 2000 to 2002, 147 subjects with moderate/severe asthma were recruited from the outpatient Pneumology Unit of São Paulo Federal University (UNIFESP). Patient selection was based on the Global Initiative for Asthma (17). These subjects fulfilled the inclusion criteria, i.e., age 18-65 years, a diagnosis of moderate to severe bronchitic asthma controlled with regular inhalatory corticosteroids at doses equivalent to 1000 µg/day beclomethasone by a metered dose inhaler for at least 60 days, and the presence of GER-compatible clinical evidence (18), such as heartburn symptoms or regurgitation up to two weeks before the initial interview. The exclusion criteria were: a diagnosis of chronic obstructive lung disease; a history of smoking during the previous year (even if the individual had already stopped smoking); infectious diseases of the upper airways during the previous 6-month period; use of oral corticosteroids; presence of co-morbidities, particularly gastrointestinal, neurologic or metabolic diseases; pregnancy; illiteracy; alcoholism; sleep-related disorders; night-time shift working, and the use of medications with an impact on sleep. The exclusion of asthmatic subjects who complained of night-time or morning aggravation over 20 days prior to the initiation of the project was based on the decision to minimize the influence of any other respiratory condition on our data.

The study also included anthropometric and polysomnography (PSG) data from 31 healthy controls with sleep disturbances but without respiratory or gastroesophageal complaints, matched for sex and age, taken from the database of the Sleep Medicine and Biology Discipline, UNIFESP.

The study was approved by the Research Ethics Committee of UNIFESP. The participants were provided with all the relevant information and gave written informed consent to participate in the research on a voluntary basis.

Sleep study

All 31 patients included were invited to the Sleep Laboratory of UNIFESP for an initial clinical sleep evaluation by first-night basal full PSG followed by placement of a pH catheter and a subsequent second night of full PSG.

The computerized Oxford System™ (UK) and Sleep Analyzer Computer (version 9.3) were used for sleep recordings, carried out for at least 7 h. Of the 16 channels of the system, three were used for the electroencephalogram (C3-A2, C4-A1, O1-A2), two for the electrooculogram (right and left), two for the electromyogram (chin and the anterior tibialis muscle), one for the electrocardiogram (V2 modified), and the remaining ones for body position sensor, nasal thermistor, microphone, and abdominal and thoracic belts. Pulse oximetry (Ohmeda™, USA) was also carried out. After the recordings, the sleep stages were analyzed visually and blindly by an experienced investigator at a sampling frequency of 256 Hz/channel, in consecutive epochs of 30 s according to standardized criteria (19). The percents of sleep stages were measured as percent total sleep time and the percents of wake after sleep onset as percent total recording time. Respiratory events (20) and arousals (21) were also scored.

Gastroesophageal study

Esophageal pH monitoring (Zinetics, Meditronic Medizinelektronik GmbH, Hohen, Neuendorf, Germany) was initiated immediately after the basal PSG, at 8:00 am with subjects in the fasting condition. The two-channel pH catheter was introduced nasally and placed 5 cm from the lower esophagus, tracked by esophageal manometry. During the study period, catheterized subjects recorded eating times on a daily basis (beginning and end of food ingestion period), time spent lying down, the taking of regular medication, along with all symptomatic periods. Patients were instructed to maintain their usual level of physical activity and meal characteristics, with the exception of acidic foods. Patients were not allowed to lie down...
during the day, when pH monitoring occurred. During the night, pHmetry recordings were performed concomitantly with the second PSG recording. All data obtained were stored in a database. The recording and analysis of GER events were performed by the Esophogram software (Gastrossoft, Irving, TX, USA).

Acid GER was defined as a drop of pH below 4, with a minimum duration of 15 s (22). Patients were classified according to the criteria proposed by DeMeester et al. (22) based on the following parameters measured on the distal electrode: total number of reflux episodes, number of reflux episodes with pH below 4 lasting 5 min or more, duration in minutes of the longest reflux episode, percent of total time with pH below 4, percent of time with pH below 4 in the orostatic position period, and percent of time with pH below 4 in the supine position period. Subjects were classified as DeMeester positive for scores of 14.72 or more and as DeMeester negative for scores below 14.72. According to the position during reflux, GER was classified as orostatic or supine. GER episodes during sleep were analyzed according to time of reflux, duration in seconds, sleep stage at the time of reflux and during the period of 1 min that preceded it. Sleep fragmentation was defined in terms of changes in sleep stage, arousals and awakenings at the time of GER and during the 1-min period that preceded it. Asthma exacerbation related to GER episodes was clinically evaluated for each episode, and again in the morning, upon completion of both sleep recordings.

### Statistical analysis

The normality of data distribution was submitted to the Kolmogorov-Smirnov test. Descriptive data are presented as mean ± SD for parametric data and median (interquartile range: 25th and 75th percentiles) for non-parametric ones. Comparisons between DeMeester-positive and -negative patients regarding the quantitative variables were performed by the Student t-test or by the Mann-Whitney U-test when the distribution of the data was not normal. Qualitative variables were analyzed by the chi-square or Fisher test as appropriate. The level of significance adopted was 0.05. The ‘Statistica’ Statistical package version 5 was used for data processing.

### Results

Of the 147 subjects selected, only 31 completed the trial, while the others dropped out because they presented one or more of the following features during the study: infectious diseases of the upper airways (8 patients), asthma aggravation (30 patients), and being absent during one or more phases of the study (78 patients).

The 31 patients included in the study (age: 47 ± 10 years; range: 18-65 years, 28 females, 3 males) exhibited basal forced expiratory volume in the first second (FEV₁) of 1.6 ± 0.6 L/min, and a DeMeester index of 15 ± 38. In comparison to the 31 healthy controls matched for sex and age (age: 47 ± 9 years; range: 18-65 years, 28 females, 3 males), the body mass index (BMI) of the asthmatic subjects was higher (30 ± 8 vs 21 ± 2 kg/m², respectively, P < 0.001). Of the 31 asthmatic subjects, 9 (29%) had a BMI >25 kg/m² and 13 (42%) a BMI >30 kg/m², whereas only 1 of the control group volunteers displayed a BMI higher than 25 kg/m² (27 kg/m²).

In comparison to the control group, asthmatic patients presented significantly lower total sleep time (434.2 ± 44.3 vs 360.2 ± 38.6 min, P < 0.001), sleep efficiency (92.8 ± 4.6 vs 81.9 ± 8.44%, P < 0.001), percent REM sleep (23.1 ± 3.2 vs 19.5 ± 6.2%, P = 0.006), and also higher mean values for arousals (4.8 ± 5.7 vs 18 ± 13.7/h, P < 0.001), and percent total sleep time with blood oxygen saturation below 90% (0.2 ± 0.2 vs 0.8 ± 2.2%, P < 0.001).

Among the 31 asthmatic subjects, the median number of GER episodes, detected by esophageal pH recording, was greater during the day in the orostatic position than during the night in the supine position (35, range 1-262, vs 5, range 0-48, respectively, P < 0.001).

Seventeen of the 31 asthmatic subjects were DeMeester positive while 14 were DeMeester negative, with no significant difference between groups in terms of sex, age, BMI, or FEV₁ values. Additionally, there was no significant difference between the two basal FEV₁ values obtained after sleep recording for each individual.

Over the 24-h period encompassing GER recordings, the DeMeester-positive group exhibited a total of 1596 GER episodes, 1262 (79%) of which occurred in the orostatic position. In the DeMeester-negative group, a significantly lower number of GER episodes was detected, i.e., 254 events (χ² = 17.22, P < 0.001), 229 (90%) of which occurred in the orostatic position.

No significant difference in PSG parameters was detected between the DeMeester-positive and -negative groups (Table 1). Sleep oscillations and variation in esophageal pH were detected in 10 of the 17 DeMeester-positive subjects, and in 7 of the 14 DeMeester-negative ones, corresponding to a total of 167 GER episodes in the DeMeester-positive group and 12 in the DeMeester-negative group. The data for the remaining patients were not studied due to artifacts in sleep recordings. Modifications in sleep pattern were evaluated at the time when the esophageal pH level fell below 4.0, along with readings during the minute preceding the onset of GER (Table 2). In the DeMeester-negative group, only one GER was fol-
lowed by arousal. Awakening was the most prevalent condition at the time of a GER event and during the minute that preceded it (P = 0.01 in both groups). Although awakening was also the most prevalent condition in the DeMeester-positive group, stage 2 was also a prevalent sleep stage during the 1 min that preceded GER (Table 2). In this group, in 72 of the 103 GER that occurred during sleep, sleep stage modification or arousal events were detected throughout the period from 1 min before to 1 min after the GER event: of 18 GER in stage 1 non-REM sleep, 94% were associated with changes to a superficial pattern of sleep (P < 0.001, Fisher test), and this was the case for 78.3% of 60 GER events in stage 2 non-REM sleep (χ² = 39.14, P < 0.001) and for 85.7% of 7 GER events during REM sleep (P = 0.04, Fisher test; Figure 1). No significant influence of GER on sleep was observed in stages 3 and 4. Arousals preceded the detection of 22 GER episodes, and followed 28 episodes (Table 2). Modifications in sleep showed no significant relationship with asthma exacerbation, since the subjects did not exhibit cough, chest wheezing or dyspnea symptoms, nor did they need to use asthma relief medication during the study period.

**Discussion**

This study was designed to evaluate the role of GER in sleep abnormalities in patients with asthma and GER-related symptoms, using all-night PSG recordings along with a 24-h monitoring of esophageal pH. In comparison to a matched control group without respiratory or gastrointes- tinal symptoms, mean BMI value was higher in the subjects with asthma and GER symptoms. These subjects presented evidence of sleep fragmentation, reduction in blood oxygen saturation, but a normal apnea-hypopnea index.

Although higher BMI values were observed in the group of asthmatics than in controls, the findings of this study are applicable to asthmatics in general, since the prevalence of obesity in asthma has been described (24,25). Considering the role of obesity as a risk factor both for GER disease (26) and sleep-related breathing disorders (27),

| Table 1. Sleep characteristics of asthmatic subjects according to the DeMeester index. |
|-----------------------------|-----------------------------|-----------------------------|
| Variables                  | DeMeester                    | DeMeester                    |
|                            | negative (N = 14)            | positive (N = 17)            |
| TST (min)                  | 362.1 ± 37.8                 | 358.7 ± 40.4                 |
| min-max                    | 288–404                      | 294–437                      |
| Efficiency (%)             | 84.2 ± 7.7                   | 80.1 ± 8.8                   |
| min-max                    | 70.2–93.7                    | 68.6–95.9                    |
| Stages 1 and 2 (%TST)      | 56.5 ± 8.9                   | 60.9 ± 11.8                  |
| min-max                    | 44.5–74.4                    | 47.6–90.4                    |
| Stages 3 and 4 (%TST)      | 23.5 ± 6.4                   | 19.7 ± 8.4                   |
| min-max                    | 7.4–30.5                     | 5.3–36.8                     |
| REM (%TST)                 | 19.96 ± 5.5                  | 19.2 ± 6.9                   |
| min-max                    | 8.3–27.2                     | 4.3–29.6                     |
| Brief arousal index events (N/h) | 20.9 ± 11.9               | 18.9 ± 9.9                   |
| min-max                    | 5.3–50.8                     | 8.5–40.5                     |
| SAT O₂ <90 mmHg (%TST)     | 4.7 (9.8)                    | 0.8 (1.3)                    |
| min-max                    | 0–16                         | 0–4                          |
| AHI (events/h)             | 2.4 (3)                      | 1.6 (5.1)                    |
| min-max                    | 0–23.4                       | 0–26                         |

Data are reported as means ± SD, minimum and maximum, or median interquartile range (25th and 75th percentiles). TST = total sleep time; REM = rapid eye movement; SAT = saturation; AHI = apnea-hypopnea index. There were no significant differences between groups (Student t-test for independent samples or Mann-Whitney U-test).

<table>
<thead>
<tr>
<th>Table 2. Influence of gastroesophageal reflux on sleep.</th>
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<tr>
<td>DeMeester negative (N = 7) 12 GER episodes</td>
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<tr>
<td>1 min before GER episodes</td>
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<tr>
<td>Stage 0</td>
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<td>Stage 1</td>
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GER = gastroesophageal reflux; SWS = slow wave sleep; REM = rapid eye movement.

*The DeMeester-negative group presented more stage 0 sleep 1 min before and during GER episodes than other sleep stages (P < 0.01; Fisher and χ² tests). *The DeMeester-positive group presented more stage 0 sleep 1 min before and during GER episodes than other sleep stages (P = 0.0007; χ² test).
we propose that this aspect may have influenced some of the results of the present study, including the reduction in blood oxygen saturation possibly caused by a certain degree of hypoventilation. Therefore, obesity, associated with the likelihood of subclinical bronchial hyper-reactivity, could justify the increased number of arousals in the group of patients. The influence of upper airways resistance syndrome cannot be excluded in spite of the normality of the apnea-hypopnea index and the absence of daytime sleepiness complaints among patients, in view of the fact that a specific device was not used during sleep recording (28).

The use of DeMeester criteria to analyze esophageal pH variability is justified given that this method considers the duration of events with pH <4 as a key factor. In our view, this corresponds to the parameter that best discriminates physiological from pathological GER events (29,30).

Regarding GER, 17 of the 31 (50%) asthmatic patients were classified as DeMeester positive. The higher prevalence of orthostatic GER in both DeMeester-positive and negative groups highlights the greater competence of the esophagogastric junction in the supine rather than in the orthostatic position (22,31). Furthermore, physiological GER events, which are less severe and rarely symptomatic, mainly occur in the orthostatic position, particularly after meals.

GER episodes occurred more frequently during wakefulness. During sleep, GER events are less common, and may reflect a dysfunction of the lower esophageal sphincter, leading to inflammation and erosion of the esophageal mucosa due to the continuous and prolonged contact with the gastric content (23) or to frequent modifications in esophageal pH (32). Even though the contact time between the esophageal mucosa and acid is a more important factor than the number of GER events, the total number of reflux episodes was taken into consideration in the present study in order to assess sleep alterations related to GER.

Despite the higher prevalence of wakefulness at the time of GER detection, 62% of the events were preceded by sleep, particularly stage 2 of non-REM sleep, showing that sleep alterations occurred at least one minute prior to the fall in esophageal pH to values of 4 or below. A further increase in arousals with modifications of sleep stages preceding GER events during stages 1 and 2 of non-REM sleep was a notable feature. As previously described, GER events were less frequent during REM sleep, a period during which secondary contractions of the esophagus might act as a protecting factor for the esophageal mucosa in contact with the acid refluxate (2,33). Sleep alterations, at least one minute prior to the drop in esophageal pH to values of 4 or below, were independent of the duration of the GER event. Modifications in sleep to a light sleep architecture, characterized by arousals, lighter sleep stages or awakening after sleep onset, can be interpreted as a protection of the esophageal mucosa by permitting swallowing and heightened peristalsis, thereby avoiding prolonged local acidity and lesions to the esophageal mucosa (33).

The reports of sleep disruption in asthmatic patients could be attributed to asthma (26,34), GER (31,33), or to aggravation of asthma by GER events (9,10,13,14,35). Even though the time-related association of GER and blood desaturation episodes was not detected in the present study, nor was exacerbation of asthma observed after GER events, the role of asthma in sleep fragmentation cannot be excluded (26,34).

The impact of sleep (12,34,36) or GER (9,10,13,14,35) on day- or night-time asthma attacks remains unresolved. In a controlled study of 9 patients with asthma, Hughes et al. (4) failed to find a relationship between GER episodes and modification of respiratory sleep parameters, since GER episodes occurred mainly during arousals or awakening periods. In a study based on simultaneous and continuous recordings of pH and lower airway resistance, Tan et al. (16) was also unsuccessful in establishing any impact of GER on day- or night-time asthma attacks.

To the best of our knowledge, this is the first study highlighting sleep fragmentation preceding GER events in patients with asthma. Even though a cause and effect relationship between GER and sleep disruption cannot be fully established, it can be suggested that decreases in pH which precede GER detection by pHmetry (32) could have an impact on sleep architecture. The assessment of any

![Figure 1](https://www.bjournal.com.br)
swallowing activity, the analysis of a longer sleep period preceding GER detection, and the inclusion of a sample of non-asthmatic individuals with GER symptoms would also be required to clarify the relationship between acid GER and sleep disruption.

A limitation of this study is related to the detection of GER events by monitoring esophageal pH, since this detection can be biased by esophageal alkalization due to the presence of saliva, sub-mucous secretion, and the presence of food particles, oral infection or esophageal obstruction (33,37). The combination of intraluminal impedance measurement (38) and esophageal pH monitoring has been considered to be the most appropriate method for the detection of GER (39), despite the fact that the classification of GER as acid and non-acid is theoretical, in view of the deleterious effects of both on the esophageal mucosa (40).

In conclusion, no difference in sleep architecture was detected between DeMeester-positive and -negative subgroups of asthmatic patients, and GER events were more frequent during the daytime and in wakefulness. When GER occurs during sleep, it can be related to sleep instability, preceding the detection of reflux. Further studies are needed to elucidate these findings.

References

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