

# ORIGINAL RESEARCH ARTICLE

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# Gastroesophageal reflux disease in our asthma patients: the presence of dysphagia can influence pulmonary function

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# Abstract

**Background:** The prevalence of Gastroesophageal Reflux Disease (GERD) in Turkey is reported as 11.6%. Studies of pulmonary function in asthmatics have demonstrated a correlation between lung resistance and the occurrence of spontaneous gastroesophageal reflux. Few studies have included measures of lung diffusing capacity for carbon monoxide. The aim of this study is to assess whether asthma patients had worse lung function and gas diffusion according to diversity of GERD symptoms they concurrently experienced. The secondary aim of the study is to determine the frequency and different faces of GERD in our asthma patients compared to healthy controls.

**Methods:** Sixty consecutive asthma patients evaluated at the pulmonary specialty outpatient clinic were included in the study. The control group included 60 healthy volunteers who had normal pulmonary function and routine laboratory tests. A modified version of a self-reported questionnaire developed by Locke and associates at the Mayo Clinic was conducted face-to-face with consecutive asthma patients and control subjects. Pulmonary function measurements were taken using spirometry. DLCO (mL/dk/mmHg) and DLCO/VA (DLCO adjusted according to alveolar volume) were measured using a single-breath technique. Statistical analyses were performed using the SPSS 17.0 statistical software.

**Results:** DLCO and DLCO/VA were significantly lower in asthma patients who had dysphagia symptoms. Frequent and significant acid regurgitations were seen in 28.33% (n = 17) of patients in the study group and 6.7% (n = 4) of patients in the control group. Severe, troublesome heartburn symptoms were reported by 28.2% (n = 17) of patients in the study group and 16.7% (n = 10) of subjects in the control group. Dysphagia was detected in 38.3% (n = 23) of all asthma cases and in 1.7% (n = 1) of the subjects in the control group.

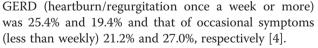
**Conclusions:** There were many faces of gastroesophageal reflux disease in our asthmatic patients. Dysphagia was the only GERD symptom influencing on pulmonary function tests, while gastroesophageal reflux symptoms and nocturnal awakening attacks were common in this study.

Keywords: Asthma, Dysphagia, Gastroesophageal reflux disease, Lung function

# Background

Gastroesophageal reflux disease (GERD) is rather common and its prevalence in Turkey is 11.6%. GERD is frequent in adult patients with asthma and the reported prevalence ranges from 32 to 82% [1-3].

Bor et al. in a study in Turkey reported in patients with asthma and in controls that the prevalence of



Extraesophageal reflux disease is commonly seen at gastroenterology outpatient clinics as well as ear, nose and throat, allergy and asthma clinics. The reflux of acid and pepsin may affect the respiratory tract and cause respiratory problems such as asthma, pneumonia and interstitial fibrosis. Chronic cough, chronic laryngitis and asthma are significantly associated with GERD [5]. There is a cause and effect relationship between asthma



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and gastroesophageal reflux which can turn into a vicious cycle. Reflux may precipitate asthma either via a vagal reflex initiated by gastric fluid in the esophagus or by micro-aspiration of gastric content into the trachea. Inversely, asthma may promote reflux due to the increased gradients pressure between the abdominal cavity and thorax, outgoing the lower esophageal sphincter pressure barrier [6-8]. Experimental evidence in both animals and humans has demonstrated reflex stimulation of bronchospasm and cough as a response to esophageal acidification.

Animal studies also have demonstrated the development of laryngeal ulceration and severe bronchospasm as a result of the direct application of acid into the larynx or acid inhalation into the airways [9-11]. Some studies of pulmonary function in asthmatics have demonstrated a correlation between lung resistance and occurrence of spontaneous gastroesophageal reflux [12]. In a review by Field *at al.* there was no objective improvement in pulmonary function attributable to GERD medical therapy. The discomfort associated with GERD can cause reflux-associated respiratory symptoms even when the pulmonary function is normal [9].

However, few studies have included measures of the lung diffusing capacity for carbon monoxide. The only study on the subject has found that severe GERD is associated with an impairment of gas exchange [13]. The aim of this study is to assess whether asthma patients have worse lung function and gas diffusion according to the diversity of GERD symptoms they concurrently experience. The secondary aim of the study is to determine the frequency of GERD in asthma patients compared to healthy controls.

## **Methods**

Sixty consecutive asthma patients participated in the study, and they had been seen at the pulmonary specialty outpatient clinic at the Taksim Training and Research Hospital and the Yedikule Chest Disease Hospital between January 2008 and January 2009. Several inclusion parameters were used to determine eligibility for the study. Eligible patients had been diagnosed with asthma according to the guidelines of the Global Strategy for Asthma Management (GINA) at least 3 years prior to entry into the study. The asthma severity of the patients without exacerbation was defined in this study as intermittent, mild persistent, moderate persistent, and severe persistent according to GINA [14]. None of the patients were experiencing an asthma exacerbation. All patients were receiving inhaled steroids, beta agonists or leukotriene antagonists at an appropriate level for the severity of their disease. None of the patients had been previously diagnosed with any other obstructive or restrictive chronic pulmonary diseases and patients who had pulmonary abnormalities on chest radiographs were excluded. Additionally, patients with known esophageal disease such as cancer, achalasia, stricture, active peptic ulcer disease, Zollinger-Ellison syndrome, and scleroderma as well as patients who were currently abusing alcohol (more than three alcoholic drinks a day) were excluded. This was a the symptom based study, thus an invasive diagnostic toollike the esophago-gastroduodenoscopy was not used.

## **Control group**

The healthy control group consisted of 60 age-matched voluntary persons who had a normal pulmonary function and routine laboratory tests. These subjects denied having any respiratory symptoms such as dyspnea or chronic sputum production and did not have a previous diagnosis of asthma or COPD or any other respiratory illness. The above-mentioned exclusion causes were also evaluated in the control group.

## Protocol

A modified version of a self-reported questionnaire developed by Locke and associates at the Mayo Clinic [15] was conducted face-to-face with consecutive asthma patients and control subjects. We established the feasibility, reproducibility, reliability and validity of the Turkish version for examining four symptoms in detail: heartburn (which was defined as a burning sensation in the retrosternal area), acid regurgitation (which was defined as the perception of the flow of refluxed gastric content into the mouth or hypopharynx), dysphagia (which was defined as a perceived impairment of the passage of food from the mouth into the stomach) and chronic cough. The first question for each symptom served as a branch point- subjects who indicated "no" proceeded to the next symptom. The next two questions for each symptom addressed the frequency (once in a month, more than once in a month, once in a week, more than once in a week and daily symptoms) and the severity of the symptoms in the last year. Further questions assessed specific attributes of each symptom. Other questions assessed the effect of both heartburn and acid regurgitation on shortness of breath, cough, wheezing, chest pain, globus and dysphonia. The remainder of the questions assessed the patient's demographic data, smoking history and use of over-thecounter antacids and prescription anti-reflux medications. In general, the questionnaires were completed in < 30 minutes by asthma patients and controls. This study was approved by our hospital ethics committee and written informed consent was obtained from all participants in the study.

## Lung function

All FVC (forced vital capacity), FEV1 (forced expiratory capacity) and PEF (peak expiratory flow) measurements

were taken using spirometry (PC using the Ocean Winspiro program, Spirolab, MIR-Medical International research, Italia). DLCO (mL/dk/mmHg) and DLCO/VA (DLCO adjusted according to alveolar volume) were measured using a single-breath technique (model 1070; Medical Graphics; St. Paul, MN). The hemoglobin level was known at the time of testing. The DLCO was routinely adjusted for the hemoglobin if the value was outside of the normal range. When analyzing the pulmonary function data, the asthma patients with gastroesophageal reflux symptoms were compared to patients without symptoms. Each of the GER symptoms was separately evaluated.

#### Statistical analysis

The reliability of the questionnaire was calculated for dichotomous choices using kappa statistics (test-retest). Kappa values were between 0.69–1. The questions were written using a Likert-type scale and were analyzed with the Spearman correlation (rs) test (rs values were between 0.67-0.95). The internal consistency of measures with dichomotous choices was evaluated by the Kuder-Richardson Formula 20 (KR-20) and the value obtained was 0.89. At the same time, the Cronbach  $\alpha$  was used to assess internal consistency for measures with nondichomotous choices and was calculated to be 0.85. The normality of the distribution was assessed using the Kolmogorov Smirnov test and the Shapiro-Wilk test by drawing histograms. The data were presented as mean  $\pm$  SD, median(min-max), frequency and percentage. The variables in each group were compared using a Mann-Whitney U-test. The nominal variable values were evaluated using the Chi-square test, Yates arranged Chi-square test and Fisher exact test. Analyses were performed using the SPSS 17.0 statistical software. The statistical significance value was set at p < 0.05.

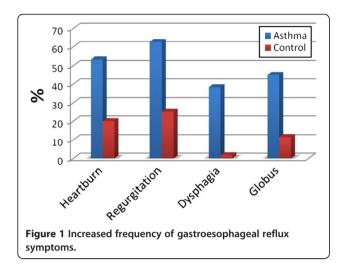
## Results

The male to female ratio of the asthmatic patients was 26/24, while it was 24/26 in the control group. There was no statistically significant difference between these two groups with regard to sex (p = 0.14). The mean age of the study group was  $44.85 \pm 14.25$  years, the mean age of the control group was  $41.48 \pm 13.90$  years(p > 0.05)

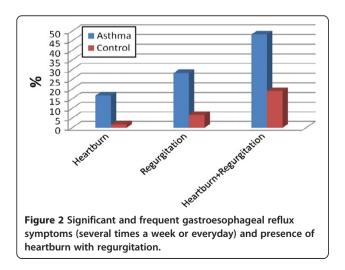
(Table 1). According to the GINA classification, 25 out of 60 patients had intermittent asthma (FEV1 and PEF  $\ge$  80%, predicted and variability <20% – step 1), 8 had mild persistent asthma (nocturnal symptoms, variability 20-30% - step 2), 21 had moderate persistent asthma (daily symptoms FEV1 and PEF 60-80%. variability  $\geq 30\%$  - step 3) and 6 had severe persistent asthma (daily symptoms, FEV1 and PEF  $\leq$  60%, variability > 30% - step 4). None of the asthmatic patients was current smoker, however 10.3% (n = 6) were ex-smokers and 10.0% (n = 6) of the healthy controls were current smokers. No subjects in either group drank alcohol. Patients had not regularly used anti-reflux therapy. However, 8 had used proton-pomp inhibitors several times(13.3%). The frequency of any heartburn in the asthma cases was 53.3% (n = 32) and in the control group it was 20% (n = 12) (p = 0.001) (Figure 1). The incidence of frequent and significant heartburn (occurring daily or at least a few times a week) was 16.7% (n = 10) in the study group and 1.7% (n = 1) in the control group (p = 0.05) (Figure 2). Severe, troublesome heartburn symptoms were reported by 28.2% (n = 17) of patients in the study group and 16.7% (n = 10) of control subjects. The frequency of regurgitation in asthma patients was 62.7% (n = 37), but only 25% (n = 15) in the control group (p = 0.001) (Figure 1). Frequent and significant regurgitation was seen in 28.33% (n = 17) of patients in the study group and 6.7% (n = 4) of control subjects (Figure 2). Severe, troublesome regurgitation (as reported by the patients) was seen in 27% (n = 10) of the asthmatics with regurgitation, and 6.7% (n = 1) of subjects in the control group. Both heartburn and regurgitation were experienced by 48.3% (n = 29) of asthmatics and 19.3% (n = 11) of controls (p = 0.001) (Figure 2). No significant association was seen between heartburn, symptoms of regurgitation and a limitation of daily activities in either the study or the control group (p = 0.07). Dysphagia was detected in 38.3% (n = 23) of asthmatics and in 1.7% (n = 1) of controls (p < 1)0.001) (Figure 1). When patients were questioned in detail, dysphagia was reported intermittently and more often occurred for solid foods than for liquids. The dysphagia symptoms experienced by the patients did not progress. Some patients had a hard time defining the difference between dysphagia and difficulty swallowing. Frequent

 Table 1 Demographics of patients with asthma and control subjects

Variables		Asthma Patients (n = 60)	Control Subjects (n = 60)	р
Gender	F n(%)	34 (56.7)	26 (43.3)	0.14
	M n(%)	26 (43.3)	34 (56.7)	
Age, yr				
mean $\pm$ SD, median (minmax.)		44.85 ± 14.25, 44(17.80)	41.48 ± 13.90 (18-72)	p > 0.05
BMI, kg/m <sup>2</sup>				
mean $\pm$ SD, median (minmax.)		25.93 ± 5.07, 25.9(13-42)	24.93 ± 4.40, 24.8(17.7-33.6)	p > 0.05



and significant dysphagia was found in 16.7% (n = 10) of asthmatics. Dysphonia was seen in 78.3% (n = 47), globus in 45% (n = 27) of asthmatics and 10% (n = 6) and 11.6% (n = 7) of controls, respectively (p < 0.01) (Figure 1). Chest pain was seen in 52.5% (n = 31) of asthma patients and in only 15% (n = 9) of the control subjects (p = 0.001). After a detailed history was obtained from patients with chest pain and the cardiologic workup was done, 15% (n = 9) of asthma patients had findings of ischemia (8.3% (n = 5) and hypertension (6.7% (n = 4)).These findings were detected in 1.7% (n = 1) of controls (p = 0.01). Seventy percent (n = 42) of asthmatics initially presented to the hospital with chronic cough and dyspnea. Gastroesophageal reflux symptoms were the initial presentation in only 6.8% (n = 4) of the asthmatics patients. None of the patients had undiagnosed or previously known hiatal hernia or esophageal disease. There was no significant correlation between the presence of regurgitation symptoms and the clinical severity of asthma (p = 0.79 and p = 0.73, respectively). Similarly,



there was no correlation between heartburn and regurgitation and severe cough (p = 0.163 and p = 0.14, respectively). Nevertheless, there was a significant correlation between the presence of regurgitation and waking up at night with dyspnea in asthmatic patients (p = 0.005). When asthma patients with and without heartburn symptoms were compared, there was no difference in pulmonary function test parameters such as FEV1,FVC, FEV1/FVC, PEF and pulmonary diffusion tests parameters such as DLCO, DLCO/VA (Table 2). The presence of heartburn and regurgitation did not affect pulmonary function. However, when patients with and without dysphagia symptoms were compared, the pulmonary diffusion test parameters such as DLCO, DLCO/VA were significantly lower in patients who had dysphagia (p = 0.008, p = 0.02, respectively). Similarly, pulmonary function test parameters such as FEV1, FVC, PEF were lower in asthma patients with dysphagia (p = 0.044, p = 0.049, p = 0.019) (Table 3).

#### Discussion

The main findings of this study are the changes of pulmonary function in patients with dysphagia. In fact, in this study pulmonary diffusion test parameters such as  $DL_{CO}$ ,  $DL_{CO}$ /VA were significantly lower in patients with dysphagia compared to those without dysphagia. Pulmonary function test parameters such as FEV<sub>1</sub>, FVC, PEF, were lower in asthma patients with dysphagia. Previous studies in the literature have already compared pulmonary function test results in patients with and without GERD, while no study has compared the pulmonary function test results according to diversity of GERD symptoms.

To the best of our knowledge there are in literature no studies other than that done by Schachteret al. in which the pulmonary diffusion capacity was measured in patients with GERD [13]. That study was performed in a group composed of obese people with severe GERD symptoms and a low diffusion capacity was found in those patients. There is no difference in the prevalence of asthma and/or airway obstruction in patients with and without GERD. In our study we did not find any changes in the diffusion capacity of patients who had heartburn/regurgitation. In Tan et al. study, GERD has not been shown to worsen lung function or bronchial reactivity consistently [16]. Field et al. determined increased minute ventilation without changing lung function in non-asthmatic patients exposed to experimental acid perfusion [17]. However, patients with dysphagia, had changes in pulmonary diffusion parameters in this study. The carbon monoxide diffusion test has been previously found to be either normal or even high in asthmatic patients [18]. The spectrum of reflux disease spans from non-erosive reflux disease to esophageal

	Asthma Patients		
	HB (n = 32)	non-HB (n = 28)	р
Age mean ± SD	44.69 ± 12.1	45.04 ± 16.6	0.93
BMI mean ± SD	27.05 ± 5.22	26.8 ± 5	0.85
FEV1 L mean ± SD	$2.20 \pm 0.80$	$2.08 \pm 0.83$	0.561
FEV1% of predicted mean $\pm$ SD	73.25 ± 21.72	$69.32 \pm 20.60$	0.475
FVC.L mean ± SD	3.01 ± 1.11	$2.91 \pm 0.89$	0.703
FVC% of predicted mean $\pm$ SD	82.40 ± 22.11	$82.54 \pm 16.04$	0.977
FEV1 FVC% mean ± SD median (minmax.)	3.02 ± 10.28 73.9(35-88)	70.53 ± 12.32 71.7(48-94)	0.405
DLCO mean ± SD	26.91 ± 8.19	$26.14 \pm 8.06$	0.717
DLCO% mean ± SD	101.31 ± 20.10	96.71 ± 15.37	0.321

Table 2 Spirometry and lung diffusion test measurements comparing patients with and without heartburn

complications such as esophagitis, hemorrhage, stricture, Barrett's esophagus and esophageal carcinoma. Prolonged exposure of the esophagus to gastric content can cause dyphagia and this can suggest that undesirable changes are likely occurring in the lung and esophagus. In a study of patients with idiopathic pulmonary fibrosis, 83% presented with abnormal distal/proximal esophageal acid exposure [19].

GERD can cause permanent histopathological changes in asthma, progression to fibrosis, and decrease in the response rate to treatment through direct irritation, hypersensitivity of the airway, and a neural reflex. An asthma exacerbation results in negative intrathoracic pressure, which may cause reflux, and medications used to treat asthma (theophylline, beta-agonists, steroids) can on turn reduce the lower esophageal sphincter pressure [20-22]. We did not find any study that showed the relationship with pulmonary fibrosis and dysphagia in asthma patients. However, Savarino et al. study explored

 Table 3 Spirometry and diffusion capacity of the lung for

 carbon monoxide in patients with and without dysphagia

PFT Results in Asthma Group						
(mean ± SD)	Dysphagia (n = 23)	non-Dysphagia (n = 37)	p value			
Age, yr	45.3 ± 14.7	44.57 ± 14.16	0.85			
BMI	$26.44 \pm 6.28$	$27.24 \pm 4.27$	0.56			
FEV1.L	$1.89 \pm 0.68$	$2.30 \pm 0.86$	0.044			
FEV1% of predicted	68.69 ± 23.17	73.10 ± 19.88	0.454			
FVC.L	$2.66 \pm 0.82$	3.15 ± 1.07	0.049			
FVC% of predicted	$81.69 \pm 20.74$	83.08 ± 18.71	0.766			
FEV1.FVC.%	70.63 ± 13.48	$72.55 \pm 9.85$	0.564			
DLCO	$23.38 \pm 6.19$	$28.54 \pm 8.54$	0.009			
DLCO%	$92.00 \pm 13.59$	103.62 ± 19.17	0.008			
DLCO/VA	4.96 ± 0.81	$6.30 \pm 4.78$	0.0103			
DLCO/VA	98.66 ± 16.19	114.67 ± 19.77	0.002			

the association of GERD and pulmonary fibrosis in patients with scleroderma with and without pulmonary fibrosis. Their findings suggested that patients with cutaneous systemic sclerosis and pulmonary fibrosis have more extensive esophageal involvement, leading to more severe GERD. Low LES (lower esophageal sphincter) pressure and smooth muscle dysfunction (i.e., uncoordinated pressure waves and reduced or absent peristalsis) predisposed to GERD in these patients [23,24]. Their experiments can be model to explain the coincidence of dysphagia and pulmonary fibrosis [23-25]. The result of Masato et al. study supported that lung inflation modulated the swallowing reflex [26]. The observed impaired breathing and swallowing patterns in the patients with COPD suggests a possible explanation for the development of dysphagia [27-29]. Our study was limited to prove the progression to pulmonary fibrosis or lung inflation with radiologic tools in asthmatic patients with dysphagia. Recurrent airway inflammation and aberrant remodeling is thought to be the basis of the fibrotic response seen in patients with asthma. The numbers of fibrocytes were significantly increased in the lamina propria and in the peripheral blood of patients with severe asthma [29-31]. Both the asthma itself and accompaniment of GERD may predispose to fibrosis. In this study the incidence of dysphagia was similar to that reported by Vakil et al., that is 37% in patients with erosive esophagitis [32]. In another study on asthma patients, we detected dysphagia in 26.7% of patients with esophagitis [33]. When patients were questioned in detail, dysphagia was noted to be intermittent and nonprogressive. Non-troublesome and non-progressive dysphagia is common in GERD. Non-progressive dysphagia can resolve in most patients. In this study, the incidence of frequent and significant dysphagia was 16.7%. However, none of the patients had progressive dysphagia. In patients with troublesome and frequent dysphagia the extent to which the dysphagia progresses should always

be determined. Progressive dysphagia is an alarm symptom that warrants further examination for peptic stricture and upper gastrointestinal malignancy [34].

According to the Montreal definition and classification, heartburn and/or regurgitation are typical gastroesophageal reflux symptoms [12]. In this study the incidence of heartburn and acid regurgitation symptoms in asthma patients were consistent with the literature [1-3]. Frequent acid regurgitation was seen in 28.3% of asthma patients. The incidence of acid regurgitation detected in asthma patients in this study was similar to the findings of Boret al. (25.4% of asthmatic subjects had heartburn/regurgitation once a week or more) [4]. When questioning patients, it is important to make sure that patients understand the meaning of each symptom [5]. While severe heartburn and regurgitation were present in 28.2% and 27% respectively of the studied patients, these percentages reflect the personal perception of the severity of symptoms and may be subject to personal and individual differences. Moreover, none of our patients had reflux symptoms that limited their daily activities. Heartburn/regurgitation symptoms coexisted in 48.3% of asthma patients. In the study by Klauseret al. that examined a patients' population referred for pH monitoring, a sensitivity of 78% and a specificity of 60% for heartburn were reported [35]. The study by Moayyedi P. et al. showed that patients with "dominant heartburn" have a 50% chance of having GERD, as diagnosed by 24-h esophageal pH studies [36]. Furthermore, nonacid esophageal reflux (pH of > 4) has also been recognized [37,38]. A varying prevalence of non-cardiac chest pain has been reported in studies from a variety of countries. A number of studies have reported a population prevalence of chest pain of up to 25% [39]. A study in China was similar to our study with a population prevalence of chest pain of 20.6%. The authors found that GERD was present in 51% of subjects with noncardiac chest pain [40]. GERD can cause episodes of chest pain that resemble ischemic cardiac pain without accompanying heartburn or regurgitation according to the Montreal definition and classification. Esophageal motor disorders can also cause chest pain by a mechanism separate from gastroesophageal reflux [5]. Furthermore, non-respiratory symptoms such as chest pain are common symptoms of an asthma attack prodrome and become more frequent after the onset of the attack [41]. Future studies that investigate chest pain in the setting of GERD and asthma should take into account these contradictions. In the present study there was no significant correlation between the presence of gastroesophageal symptoms and the clinical severity of asthma. On the contrary, in some studies the severity of asthma has been associated with reflux symptoms. In fact, reflux symptoms were present in 30% of patients with mild asthma, 76% of those with moderate asthma and 70% of those with severe asthma [36]. However, other studies do not support these findings. Patients with wellcontrolled asthma with and without pH probedocumented acid reflux were compared in a study by Di Mango et al. and their results did not support the conclusion that asymptomatic reflux is associated with decreased lung function, worsen asthma control, increased airway hyperresponsiveness or increased asthma symptoms [42]. Many more studies are needed to investigate the association between reflux symptoms and the severity of asthma. However, the presence of acid regurgitation was significantly correlated with nocturnal awakening with dyspnea in this study. Compared to non-asthmatics, asthmatic patients have significantly more frequent and severe day and night GERD symptoms and significantly more pulmonary symptoms (nocturnal suffocation, cough or wheezing) often associated with GERD [43]. In a study done in Brazil, awakening from sleep was the most frequent symptom found at the onset of GERD, and preceded 38% of nocturnal GERD. Sleep stage 2 was also prevalent and preceded 36% of GER events [44].

#### Conclusions

In conclusion, dysphagia was the only influential GERD symptom on pulmonary function test while gastroesophageal reflux symptoms and nocturnal awakening attacks were common in this study. Dysphagia should be an alarm sign for the pulmonologist in the same manner that it is for the gastroenterologist. A pulmonary carbon monoxide diffusion test should be used in the follow up of patients who have coexisting asthma and GERD. However, more studies that investigate this topic are needed.

#### Competing interest

The authors declare that they have no competing interests.

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