

ORIGINAL RESEARCH

# Prevalence of chronic obstructive pulmonary disease among patients with systemic arterial hypertension without respiratory symptoms

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**Background:** The diagnosis of chronic obstructive pulmonary disease (COPD) is often delayed until later stages of the disease. The purpose of the present study was to determine the prevalence of COPD among adults on treatment for systemic arterial hypertension independently of the presence of respiratory symptoms.

**Methods:** This cross-sectional study included adults aged  $\geq$ 40 years with tobacco/occupational exposure and systemic arterial hypertension diagnosed at three Primary Health Care facilities in Goiania, Brazil. Patients were evaluated using a standardized respiratory questionnaire and spirometry. COPD prevalence was measured considering the value of forced vital capacity and/or forced expiratory volume in 1 second <0.70.

**Results:** Of a total of 570 subjects, 316 (55%) met inclusion criteria and were invited to participate. Two hundred and thirty-three (73.7%) patients with arterial hypertension reported at least one respiratory symptom, while 83 (26.3%) reported no respiratory symptoms; 41 (17.6%) patients with arterial hypertension and at least one respiratory symptom, and 10 (12%) patients with arterial hypertension but no respiratory symptoms were diagnosed with COPD (P=0.24). The prevalence of COPD in people with no previous COPD diagnosis was greater among those with no respiratory symptoms (100%) than among those with respiratory symptoms (56.1%) (P=0.01).

**Conclusion:** Our findings suggest that regardless of the presence of respiratory symptoms, individuals aged ≥40 years with tobacco/occupational exposure and arterial hypertension may benefit from spirometric evaluation.

Keywords: COPD, hypertension, public health

# **Background**

The World Health Organization (WHO) reports that chronic obstructive pulmonary disease (COPD) was responsible for more than three million fatalities in 2011, accounting for 5.8% of all deaths. Although 30% of adults ≥35 years of age with tobacco/occupational exposure and at least one respiratory symptom exhibited airway obstruction on spirometry, this test is often delayed until later stages of the disease because early symptoms of COPD are not frequently recognized. However, it has already been demonstrated that airflow obstruction may be observed on spirometry in COPD patients in the initial stages of the disease, even in the absence of respiratory symptoms. It is known that arterial hypertension occurs in more than 50% of COPD patients diagnosed at a primary health care level. However, the prevalence of COPD among subjects with arterial hypertension, a disease often diagnosed and followed by primary health care providers, is unknown. The purpose of the present study was to determine the prevalence of COPD among adults with tobacco/occupational exposure and arterial

Correspondence: Marcelo Fouad Rabahi Av.B, número 483, Setor Oeste, Goiania, Goiás CEP 74110-030, Brazil Tel +55 62 3521 3333 Fax +55 62 3521 3333 Email mfrabahi@gmail.com hypertension with and without respiratory symptoms in three Primary Health Care facilities in Goiania, Brazil.

#### **Methods**

# Design

Cross-sectional study.

## Subject selection

Adults aged ≥40 years with tobacco/occupational exposure and systemic arterial hypertension diagnosed according to the Brazilian Society of Cardiology criteria and who were undergoing treatment at a Primary Health Care facility were eligible for the study.7 Signed informed consent was necessary for enrollment. Exclusion criteria were reported to be asthma, presence of reversibility of the airway obstruction demonstrated on spirometry test; pneumothorax; angina; recent (within 3 months) myocardial infarction or pulmonary embolism or hemoptysis or brain, eye, ear, thoracic, or abdominal surgery; presence of known thoracic, abdominal, or cerebral aneurysms; severe hypertension (mean arterial pressure >130 mmHg) at the time of spirometry; acute illness or symptoms that might interfere with test performance (nausea, vomiting, coughing, or acute diarrhea); and patients with mental confusion or dementia.8 Subjects were also excluded if they had incomplete interviews or could not complete the spirometric test. Patients were consecutively enrolled from January 1, 2011 to May 31, 2012.

# Setting and subject selection

Brazil's health care system is organized into primary, secondary, and tertiary levels. The primary health service is named Family Health Program (FHP). Each FHP facility is responsible for a community of about 1,000 families, and has a family health team of one doctor, one nurse, one auxiliary nurse, and four to six community health agents. Community health agents visit 150 families every month on average, and the FHP provides promotive, preventive, curative, and rehabilitative services. This study was conducted at CIAMS (Center for Integrated Health Care Health) Novo Horizonte, UABSF (Primary Care Unit of the Family Health Program) Caravelas, and UABSF Santa Rita facilities in the same geographic area of Goiania, a city in Brazil's Midwest region.

## Data collection

After signing the informed consent, participants completed a pretested, standardized respiratory questionnaire including the following: demographics; smoking habits; exposure to biomass smoke; education; occupational exposure to chemicals and particulate matter; respiratory symptoms; use of medications; previous spirometry testing; previous medical diagnosis of tuberculosis, asthma, chronic bronchitis, emphysema, or COPD; self-reported exacerbation and hospitalization due to COPD and comorbid conditions (heart disease, stroke, diabetes, and asthma). Anthropometrics, blood pressure, spirometry, and oxygen saturation measurements were performed in all enrolled subjects. Spirometry was performed using the portable, battery-operated ultrasound Easy One spirometer (NDD Medical Technologies, Zurich, Switzerland). Spirometry was performed at baseline and 15 minutes after the administration of 400 µg of salbutamol, according to the criteria of the American Thoracic Society and European Respiratory Society for acceptability and reproducibility.9 Positive bronchodilator responsiveness was defined using the following criteria: forced vital capacity (FVC) and/or forced expiratory volume in 1 second (FEV<sub>1</sub>) ≥12% plus ≥200 mL improvement. 10 The presence of COPD was based on the postbronchodilator FEV<sub>1</sub>/FVC <0.70, a fixed value proposed by the Global Initiative for Obstructive Lung Disease (GOLD) criteria, among subjects 40 years old or more with tobacco/occupational exposure and no asthma diagnosis.11 Reversibility of the airway obstruction diagnosis was defined as the presence of positive bronchodilator responsiveness as defined by the American Thoracic Society.<sup>12</sup> A significant biomass exposure was defined as a subject with more than 80 hours/year with a minimum exposure of 10 years.<sup>13</sup> A subject was included in the group with respiratory symptoms in the presence of any of the following symptoms: dyspnea, cough, wheezing, or sputum production, and was included in the group without respiratory symptoms if all were absent.

# Statistical analysis

The results were analyzed using Stata version 13.0 (Stata-Corp, College Station, TX, USA). For all statistical tests, P<0.05 was considered statistically significant. t-tests were used to compare two means. Chi-square or Fisher's exact test (when applicable) was used for dichotomous variables. Pairwise correlation analysis was performed between pulmonary function parameters and the blood pressure values, and the Bonferroni method was used for adjusting significance levels.

# Ethical aspects

This study was conducted according to good clinical practices, and was approved by the Ethics Committee of the Hospital of the Federal University of Goiás under number 195/2011.

#### Results

Over a 17-month period, 1,325 patients were admitted to all three Primary Health Care facilities, of which 570 (40%) reported treatment for arterial hypertension in one of three facilities (Figure 1). Among these 570 subjects, 316 (55%) met the inclusion criteria and were invited to participate in the study. Two hundred and five were nonsmokers without significant biomass exposure, 10 were <40 years of age, 17 patients refused to sign the consent form, and 22 patients could not perform spirometry; thus, 316 subjects were enrolled in the study. Two hundred and thirty-three patients (73.7%) with arterial hypertension reported at least one respiratory symptom, while 83 (26.3%) reported no respiratory symptoms. Characteristics of the sample are presented in Table 1. Forty-one patients (17.6%) with arterial hypertension and a respiratory symptom, and 10 patients (12%) with arterial hypertension and no respiratory symptoms were diagnosed with COPD (P=0.24). However, the prevalence of COPD cases with no previous diagnosis was greater among patients with no respiratory symptoms than among those with respiratory symptoms (100%; 56.1%) (P=0.01). There were no additional differences related to other variables such as age, smoking history, and others as shown in Table 1. We found no association between pulmonary function parameters and the blood pressure values.

## **Discussion**

This study shows that there was no difference in the prevalence of COPD among patients with arterial hypertension and tobacco/occupational exposure; they either had respiratory symptoms or not. More than half of the patients with respiratory symptoms were unaware of their diagnosis, whereas all the patients without symptoms did not know they had COPD. In fact, systemic arterial hypertension has been described as a major comorbidity associated with COPD, as demonstrated by a population-based case-control study conducted

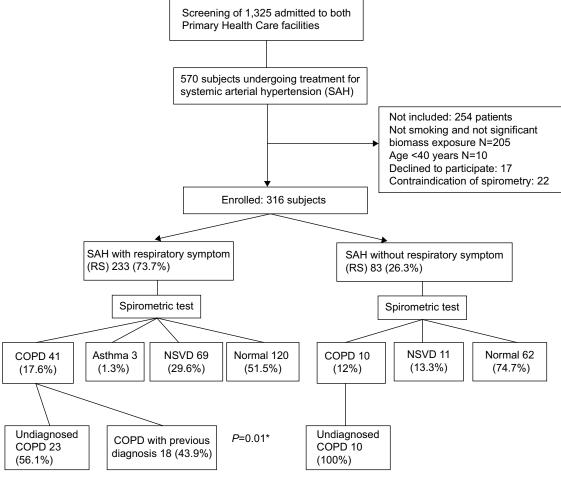


Figure I Study design and subjects.

Note: \*Fisher exact test.

Abbreviations: COPD, chronic obstructive pulmonary disease; NSVD, nonspecific ventilator disturbance.

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Table I Characterization of COPD with and without respiratory symptoms in hypertensive patients

	Group with respiratory symptoms, n=233 (73.7%) COPD, n=41 (17.6)	Group without respiratory symptoms, n=83 (26.3%) COPD, n=10 (12)	Р
50–59	6 (14.6)	0 (0)	0.33
≥60	35 (85.4)	10 (100)	
Male (race)	22 (53.7)	7 (70)	0.48
White	14 (34.2)	4 (40)	
Mestizo	20 (48.8)	6 (60)	0.69
Black	6 (14.6)	0 (0)	
Asian	I (2.4)	0 (0)	
Education (years)	3.3±3.0	4.4±2.8	0.30
BMI (m/kg <sup>2</sup> )	27.1±5.6	24.0±5.8	0.12
Systolic arterial pressure	145.1±23.3	136.7±29.0	0.33
Diastolic arterial pressure	80.7±12.4	77.7±12.2	0.47
Oxygen saturation (%)	93.8±2.8	95.7±2.1	0.06
Smoking (pack-years)	32.8±26.8	36.5±18.2	0.68
Beginning smoking (age)	13.7±7.1	12.7±4.0	0.67
Exposure to biomass (hour/year)	65.8±75.4	46.6±60.6	0.46
Occupational exposure to dust (years)	20.9±20.9	15.5±19.5	0.47
Years living in rural area	25.6±19.3	23.2±14.1	0.72
Smoking status			
Current	12 (29.3)	5 (50)	0.44
Former	24 (58.5)	5 (50)	
Never	5 (12.2)	0 (0)	
Antihypertensive medication			
β-blocker (selective)	3 (7.3)	I (I0)	0.13
β-blocker (n-selective)	17 (41.5)	I (I0)	
Other	21 (51.2)	8 (80)	
Heart disease	11 (26.8)	2 (20)	1.00
Diabetes	4 (9.7)	2 (20)	0.58
Vascular disease	3 (7.3)	2 (20)	0.25
Previous diagnosis of TB	2 (4.9)	0 (0)	1.00

Note: Data presented as mean  $\pm$  SD or n (%).

Abbreviations: COPD, chronic obstructive pulmonary disease; BMI, body mass index; TB, tuberculosis; SD, standard deviation.

in Switzerland with 340,948 COPD hospitalizations. The authors found that among 82 comorbidities studied, arterial hypertension was the comorbidity with the highest prevalence among COPD patients (24%) versus 16% in patients without COPD (95% confidence interval 1.70 [1.6–1.7], P<0.001). <sup>14</sup> Similar results were observed in the KNHANES V cohort (Korean National Health and Nutrition Examination Survey): among 15 comorbidities, only arterial hypertension and history of pulmonary tuberculosis were independently associated with COPD after adjustment for confounders. <sup>15</sup> Furthermore, a Japanese study demonstrated that 25% of patients with cardiovascular disease have COPD and were not previously diagnosed. These findings emphasize the need for active case finding strategies among higher-risk groups in order to diagnose COPD in the earlier stages of the disease. <sup>16</sup>

Our findings suggest that regardless of the presence of respiratory symptoms, people aged 40 or above with

tobacco/occupational exposure and arterial hypertension may benefit from a spirometry to screen a potential case of COPD.

The current study has an important limitation. The diagnosis of COPD is based on the association between risk factor, airway obstruction in spirometry, and the presence of symptoms among subjects aged older than 40 years (GOLD).<sup>17</sup> Our sample included individuals without symptoms. However, it should be pointed out that there are no data to prove or refute the efficacy of screening COPD cases among asymptomatic subjects using the spirometry test (GOLD).<sup>17</sup> Considering that all individuals who reported asthma and those who had reversibility of the spirometry test were excluded from our study, we believe that the positive predictive value of the airway obstruction as demonstrated by the spirometry in this particular sample (subjects with the most common COPD comorbidity sharing COPD risk factors), even without respiratory symptoms, is probably very high for COPD diagnosis.

Furthermore, respiratory symptoms cannot be detected in the earliest stages of COPD.<sup>18</sup> It is therefore necessary to perform studies assessing the value of the spirometry test for screening and diagnosis (among subjects without and with respiratory symptoms, respectively) of COPD among high-risk populations in different epidemiologic scenarios in order to confirm these current findings.<sup>18</sup>

#### **Conclusion**

Our findings suggest that regardless of the presence of respiratory symptoms, individuals aged ≥40 years with tobacco/ occupational exposure and arterial hypertension may benefit from spirometric evaluation for diagnosis and screening of COPD cases in the earliest stages of the disease.

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#### **Author contributions**

MFR contributed to conceiving this study and participated in the design, data collection, data interpretation, and drafting of the manuscript. SAP, APR, JLRJ, and ACC contributed to data collection, data interpretation, and drafting of the manuscript. KSC participated in the study design and coordination, and helped draft the manuscript. MBC participated in conceiving this study and participated in the study design, coordination, and statistical analysis and review of the manuscript. All authors read and approved the final manuscript.

#### **Disclosure**

This study was financially supported by Novartis. The authors did not receive any reimbursement or financial benefits, and declare no competing interests. Novartis had no role in the design, methods, data management, analysis, or the decision

to publish. The authors report no conflicts of interest in this work.

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