ORIGINAL ARTICLE

Relevance of individual bronchial symptoms for asthma diagnosis and control in patients with rhinitis: A MASK-air study

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Abstract

Rationale: It is unclear how each individual asthma symptom is associated with asthma diagnosis or control.

Bernardo Sousa-Pinto and Gilles Louis, the first two authors, contributed equally to the paper.

Benoit Pétré, Renaud Louis and Jean Bousquet, the last three authors, contributed equally to the paper.

A complete list of the think tank members appears in the "MASK-air think tank" section.

Objectives: To assess the performance of individual asthma symptoms in the identification of patients with asthma and their association with asthma control.

Methods: In this cross-sectional study, we assessed real-world data using the MASK-air® app. We compared the frequency of occurrence of five asthma symptoms (dyspnea, wheezing, chest tightness, fatigue and night symptoms, as assessed by the Control of Allergic Rhinitis and Asthma Test [CARAT] questionnaire) in patients with probable, possible or no current asthma. We calculated the sensitivity, specificity and predictive values of each symptom, and assessed the association between each symptom and asthma control (measured using the e-DASTHMA score). Results were validated in a sample of patients with a physician-established diagnosis of asthma.

Measurement and Main Results: We included 951 patients (2153 CARAT assessments), with 468 having probable asthma, 166 possible asthma and 317 no evidence of asthma. Wheezing displayed the highest specificity (90.5%) and positive predictive value (90.8%). In patients with probable asthma, dyspnea and chest tightness were more strongly associated with asthma control than other symptoms. Dyspnea was the symptom with the highest sensitivity (76.1%) and the one consistently associated with the control of asthma as assessed by e-DASTHMA. Consistent results were observed when assessing patients with a physician-made diagnosis of asthma.

Conclusions: Wheezing and chest tightness were the asthma symptoms with the highest specificity for asthma diagnosis, while dyspnea displayed the highest sensitivity and strongest association with asthma control.

KEYWORDS

asthma, diagnosis, dyspnea, mHealth, wheezing

1 | INTRODUCTION

Asthma guidelines such as the Global INitiative for Asthma (GINA) have improved the knowledge and management of asthma.[1] However, (i) there is a substantial number of asthma misdiagnoses (under/over-diagnosis) in primary care, partly associated with the under-use of spirometers and under-reporting of symptoms by patients,[2] (ii) many patients diagnosed with asthma are insufficiently monitored and (iii) poor adherence to treatment remains frequent.[3] These aspects are all associated with poor control, increased risk of asthma attacks and increased healthcare utilization.[4,5]

The aspects related to asthma treatment adherence, follow-up, and control may be partly addressed by patient-centered digital solutions allowing the collection of real-world data on asthma symptoms and treatment. The EU Commission has proposed digital solutions, such as wearables and mHealth apps, to engage citizens in the self-management of chronic diseases.[6] Thus, the patient perspective, captured by patient-reported outcome measures (PROMs), can strengthen patient-centered care.[7] Although various mHealth apps have been developed to support asthma monitoring and management in recent years[8,9], very few have been developed

for both asthma diagnosis and asthma control using validated PROMs. One of the apps, MASK-air® (Mobile Airways Sentinel networK), is a Good Practice of Directorate-General Health and Food Safety[10] and the only one listed as a Best Practice of Organisation for Economic Cooperation and Development for chronic diseases.[10] It includes several validated asthma PROMs.[11–13] One such PROM is the Control of Allergic Rhinitis and Asthma Test (CARAT) questionnaire, aiming to assess asthma and AR control together. It has 10 questions addressing upper and lower airway symptoms, sleep disturbances, limitation of activities and the need to increase medication over the previous four weeks.[14–16] CARAT includes three questions on key asthma symptoms (wheezing, dyspnea and chest tightness).

mHealth tools and other patient-centered digital solutions can help address the misdiagnosis of asthma not only through the assessment of directly provided daily patient data but also by the use of such data to identify specific variables associated with asthma diagnosis or control. In particular, mHealth-based direct patient data may allow us to assess whether some specific asthma symptoms or symptom patterns are associated with asthma diagnosis or control. This is in line with the recent European Respiratory Society guidelines for the diagnosis of asthma in adults, which, in their conclusion, highlighted

that more research should be undertaken on the value of symptoms to predict accurate asthma diagnoses.[17]

In this study, we used the MASK-air® PROMs data (CARAT) to assess how individual asthma symptoms (wheezing, dyspnea, chest tightness) were associated with both asthma diagnosis and control. Knowing which symptom is the most associated with asthma can help reduce the important number of misdiagnoses in the clinical practice, while knowing which symptom has the greatest impact on asthma control is fundamental in the guidance of strategies aiming to improve daily patient management.

2 | METHODS

2.1 | Study design

A full description of the Methods is available in the Online Data Supplement. In this cross-sectional study, we used MASK-air® data to compare the frequency of occurrence (over the previous month) of five asthma symptoms (dyspnea, wheezing, chest tightness, fatigue and night symptoms) in patients with probable asthma, possible asthma and no evidence of asthma[18] as assessed by CARAT. Results of this study were validated in a sample of patients in whom asthma diagnosis had been assessed by a physician in the context of a transfer of innovation project (Twinning) of the European Innovation Partnership on Active and Healthy Ageing.

2.2 | Settings and participants

MASK-air® is freely available in 27 countries. In this study, we included data from MASK-air® users from May 21, 2015 to December 2021, reporting MASK-air® data in at least three different months. The users (i) had a self-reported diagnosis of allergic rhinitis and (ii) ranged in age from 16 to 90 years (or lower than 16 years in countries with a lower age of digital consent).[19,20] We also included data from participants of the Twinning project who were enrolled during a medical consultation with an asthma specialist.[18] Asthma was diagnosed according to GINA,[1] with patients having a pulmonary function test and answering the CARAT questionnaire. Following that consultation, participants were classified as having "current asthma" or "no evidence of (current or past) asthma".

2.3 | Ethics

An Independent Review Board (Bohn-Köln) approval was obtained for the MASK-air studies.[21] For the Twinning project, additional local review board approvals were obtained, and written consent was provided by the patients. All data were anonymized before the study, and users agreed to the analysis of their data.

2.4 | Data sources and variables

The MASK-air® app comprises a daily monitoring questionnaire assessing (i) the impact of asthma and rhinitis symptoms on a daily basis by means of 0–100 visual analog scales (VASs) (with a higher score corresponding to a higher impact of allergy symptoms) and (ii) the daily use of asthma and rhinitis medication (available from country-specific lists with prescribed and over-the-counter medications).[19] The symptom and medication information provided in the MASK-air® daily monitoring questionnaire allows for the computation of the e-DASTHMA, a 0–100 score assessing the daily control of asthma.[22]

In addition to the daily monitoring questionnaire, MASK-air® also includes (although in a non-daily basis) CARAT, a questionnaire assessing the control of allergic rhinitis and asthma in the previous 4 weeks (Table E1).[15]

2.5 | Data analysis

When responding to the MASK-air® daily monitoring questionnaire, it is not possible to skip any of the questions, and data are saved to the dataset only after the final answer. This precludes any missing data. All analyses were performed using software R (version 4.0.0).

Using a two-step approach, k-means cluster analysis methods were applied to group MASK-air® users on their probability of having asthma. Obtained clusters subsequently enabled the classification of patients as having "probable asthma", "possible asthma" or "no evidence of asthma" (i.e., rhinitis alone).[18]

For patients with "probable asthma", "possible asthma" or "no evidence of asthma", using the CARAT questions,[12,15,16] we assessed the frequency of having at least one day per week of (i) dyspnea, (ii) wheezing, (iii) chest tightness, (iv) tiredness/limitations in doing tasks and (v) night symptoms. For each question, we calculated its sensitivity, specificity and predictive values. We also assessed the performance of each question in the discrimination between "possible asthma" and "probable asthma". As each patient could have answered CARAT more than once, we considered, in our main analysis, the first CARAT reported by each patient. We then performed a sensitivity analysis considering (i) all reported CARAT questionnaires or (ii) pre-COVID-19 data only. To validate the obtained results, the sensitivity, specificity and predictive values of each symptom were assessed in a sample of patients in whom the diagnosis of asthma was established by a physician (Twinning participants).

Finally, in patients with probable asthma, we computed the median e-DASTHMA by each category of each CARAT question to assess the most discriminative symptom regarding asthma control. We considered both the median and maximal e-DASTHMA levels for the 4 weeks before answers to CARAT were provided.

3 | RESULTS

3.1 | Demographic characteristics

We included 951 patients with rhinitis (62.8% women; mean ± SD age = 38.6 ± 13.6), of which 468 (49.2%) had probable asthma, 166 (17.5%) possible asthma and 317 (33.3%) no evidence of asthma. The demographic characteristics of the patients are presented in Tables 1 and 2 and Figure 1). There were 2154 CARAT assessments (Table 3 and Table E2), with 302 indicating controlled CARAT and 1852 indicating uncontrolled complete CARAT. For all these participants, e-DASTHMA information was available.

3.2 | Association between individual asthma symptoms and asthma diagnosis

We compared patients with "no evidence of asthma" versus those with "probable or possible asthma" on the presence of each asthma symptom as assessed by CARAT. We examined the first CARAT questionnaire reported by each patient ($N = 951$, Table 4; Figure 1) and observed that the specificity decreased from wheezing (90.5%) to chest tightness, dyspnea, night symptoms and fatigue (51.1%). On the other hand, the sensitivity was around 50% for wheezing and chest tightness and increased up to 63% for dyspnea. The positive predictive value (PPV) ranged from 71.6% to 90.8% (patients with wheezing had a 90.8% risk of having asthma) and the negative predictive value (NPV) was always lower than 50%.

Similar results were observed when all CARAT assessments were reported (Table 4) or when considering pre-COVID-19 data only (Table 4).

In a subsequent analysis, we compared patients with "probable asthma" versus those with "possible asthma" on the presence of each asthma symptom as assessed by CARAT. When considering only the first CARAT questionnaires reported by the patients (Figure 2; Table 5), the specificity decreased from chest tightness (79.5%) and wheezing (77.7%) to dyspnea (64.5%) and fatigue and night symptoms (50.0%). On the other hand, the sensitivity was around 55% for wheezing and chest tightness and increased up to 76.1% for dyspnea. The PPV ranged from 77.9% to 88.9%. Furthermore, the NPV was always lower than 50%. Similar results were observed when all CARAT assessments were reported (Table 5) or when considering pre-COVID-19 data only (Table 5).

TABLE 1 Demographic characteristics of the patients.

	Main analysis (N = 951)
Females—N (%)	597 (62.8)
Age—mean (SD)	38.6 (13.6)
Self-reported asthma—N (%)	393 (41.3)
Asthma classification—N (%)	
Probable asthma	468 (49.2)
Possible asthma	166 (17.5)
No evidence of asthma	317 (33.3)
Self-reported conjunctivitis—N (%)	625 (65.7)
	Twinning sample (N = 283)
Females—N (%)	162 (57.2)
Age—mean (SD)	43.4 (17.2)
Asthma classification diagnosis—N (%)	
Current asthma	97 (34.3)
No evidence of asthma	186 (65.7)
Conjunctivitis—N (%)	148 (52.3)

Abbreviation: SD, Standard-deviation.

TABLE 2 Frequency of MASK-air® users per country.

| Country	N (%)
Argentina	19 (2.0)
Australia	6 (0.6)
Austria	17 (1.8)
Belgium	9 (0.9)
Brazil	19 (2.0)
Canada	3 (0.3)
Czech Republic	6 (0.6)
Denmark	3 (0.3)
Finland	16 (1.7)
France	102 (10.7)
Germany	94 (9.9)
Great Britain	14 (1.5)
Greece	36 (3.8)
Hungary	12 (1.3)
Italy	65 (6.8)
Japan	29 (3.0)
Lebanon	4 (0.4)
Lithuania	113 (11.9)
Mexico	82 (8.6)
Netherlands	20 (2.1)
Poland	62 (6.5)
Portugal	77 (8.1)
Slovenia	14 (1.5)
Spain	59 (6.2)
Sweden	11 (1.2)
Switzerland	13 (1.4)
Turkey	46 (4.8)
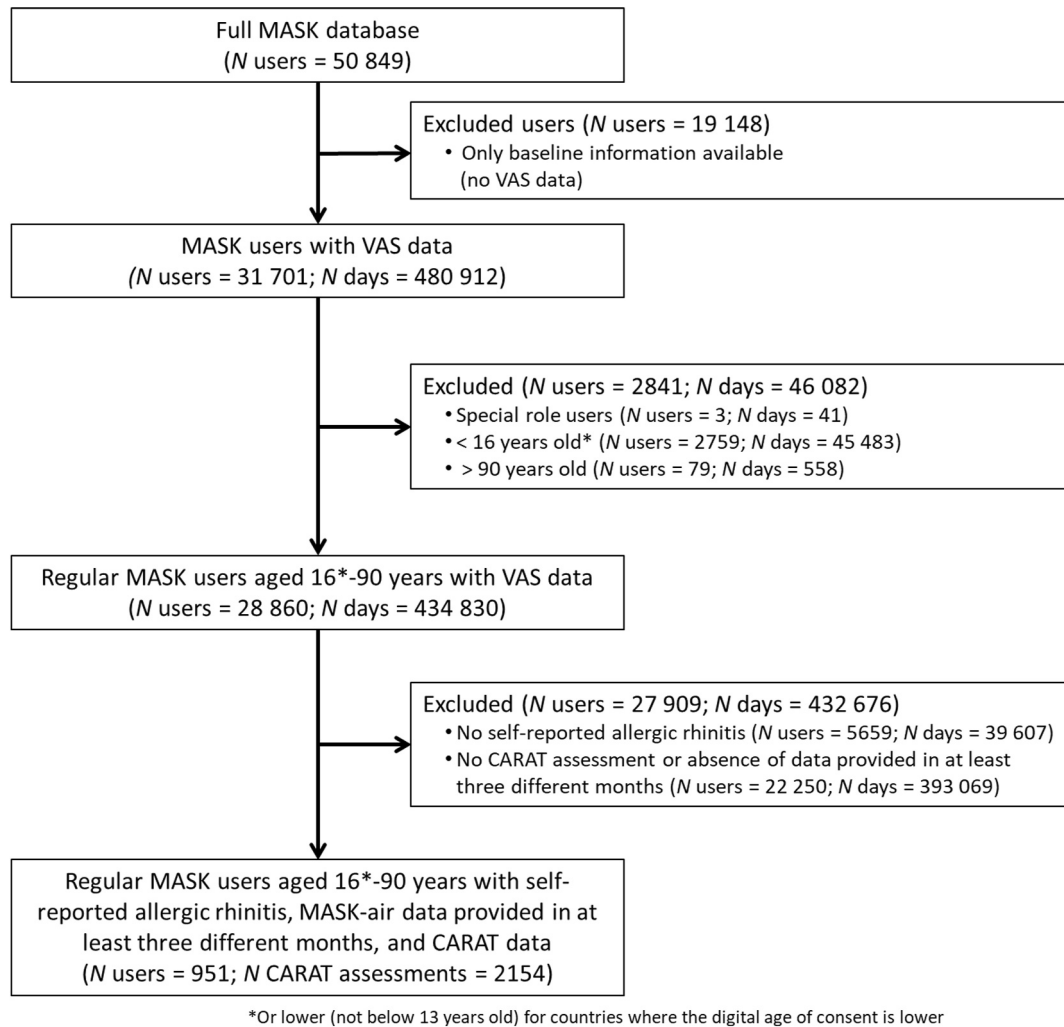
FIGURE 1 Flow chart illustrating participants' selection.

TABLE 3 Number of CARAT questionnaires provided by user.

Number of CARAT questionnaires answered by patient	N patients (%)
1	694 (73.0)
2	154 (16.2)
3	32 (3.4)
4	20 (2.1)
5	13 (1.4)
6–10	22 (2.3)
>10	16 (1.7)

Abbreviation: CARAT, control of allergic rhinitis and asthma test.

When considering the first CARAT questionnaire reported by each patient, only 21% of patients reported having never experienced dyspnea in the previous 4 weeks versus 28% for fatigue, 38% for night symptoms, 42% for chest tightness and 44% for wheezing (Figure 3). On the other hand, 45% of patients reported an average of more than 2 days per week of dyspnea, 41% reported the same for fatigue, 35% for night symptoms, 28% for chest tightness and 21% for wheezing.

3.3 | Validation of results in a sample of patients assessed by a physician

We analyzed 283 Twinning participants, of whom 97 (34.3%) had current asthma and 186 had no evidence of asthma, as assessed by a physician (Table 1). As observed in our main analysis, wheezing was the symptom which displayed the highest specificity (86.0%) and PPV (70.5%), while dyspnea was the symptom associated with the highest

TABLE 4 Comparison between patients classified as having "possible or probable asthma" versus "no evidence of asthma" on the ever occurrence of asthma symptoms as assessed by the Control of Allergic Rhinitis and Asthma Test (CARAT).

CARAT questions	Sensitivity—% (95% CI)	Specificity—% (95% CI)	PPV—% (95% CI)	NPV—% (95% CI)
A. First CARAT assessment by user				
Shortness of breath/dyspnea[a]	65.5 (61.8–69.2)	68.5 (63.3–73.6)	80.6 (77.2–84.0)	49.8 (45.1–54.5)
Wheezing in the chest	46.8 (43.0–50.7)	90.5 (87.3–93.8)	90.8 (87.7–94.0)	46.0 (42.1–49.9)
Chest tightness	48.1 (44.2–52.0)	84.5 (80.6–88.5)	86.2 (82.6–89.8)	44.9 (40.9–48.9)
Tiredness/limitations doing tasks	65.1 (61.4–68.9)	51.1 (45.6–56.7)	72.7 (69.0–76.4)	42.3 (37.3–47.2)
Night symptoms	59.1 (55.3–63.0)	53.0 (47.5–58.5)	71.6 (67.7–75.4)	39.3 (34.7–44.0)
B. All CARAT assessments				
Shortness of breath/dyspnea[b]	75.2 (73.1–77.2)	69.6 (65.2–73.8)	90.4 (88.9–91.9)	42.2 (38.7–45.8)
Wheezing in the chest	49.9 (47.5–52.2)	90.6 (87.9–93.3)	95.3 (93.9–96.7)	32.1 (29.5–34.6)
Chest tightness	58.4 (56.1–60.7)	84.5 (81.2–87.9)	93.5 (92.1–95.0)	34.7 (31.9–37.5)
Tiredness/limitations doing tasks	71.1 (69.0–73.3)	52.2 (47.6–56.9)	85.1 (83.2–86.9)	32.1 (28.7–35.5)
Night symptoms	57.2 (54.9–59.6)	49.8 (45.1–54.4)	81.3 (79.1–83.6)	23.3 (20.6–26.1)

Abbreviations: NPV, negative predictive value; PPV, positive predictive value.
[a]Considering data before 2020 only: Sensitivity = 59.5%; Specificity = 66.7%; PPV = 83.1%; NPV = 37.4%.
[b]Considering data before 2020 only: Sensitivity = 64.8%; Specificity = 68.9%; PPV = 89.5%; NPV = 32.6%.

sensitivity (76.3%) and NPV (85.6%) (Table 6). The sensitivity and specificity for each symptom were similar to those observed in the main analysis. On the other hand, a lower PPV and a higher NPV were observed in the Twinning study, reflecting differences in the proportion of patients with asthma.

3.4 | Association between individual asthma symptoms and asthma control

Dyspnea was the symptom the most strongly associated with e-DASTHMA levels (Table 7). In fact, for dyspnea, the median e-DASTHMA ranged from 5.7 (IQR = 9.5] in patients who had never reported that symptom in the past 4 weeks to 27.8 (IQR = 30.0) in those who reported the symptom every day or almost every day (high effect size = 0.97). Chest tightness and wheezing had a heterogeneous impact on e-DASTHMA (high to no effect). Fatigue and night symptoms had a more consistent effect, with medium effect sizes being observed when comparing median e-DASTHMA levels for patients who had experienced those symptoms every day (or almost) versus those who had never experienced them in the previous 4 weeks. Only dyspnea had a consistent effect on e-DASTHMA levels when all CARAT questionnaires were considered (Table 7).

4 | DISCUSSION

This is the first study to assess individual asthma symptoms for the diagnosis and control of the disease. Although all symptoms were associated with asthma diagnosis, wheezing was the symptom with the highest specificity and PPV for diagnosis. Dyspnea was the most sensitive symptom associated with the control of asthma as assessed by a novel daily electronic symptom-medication score (e-DASTHMA).

4.1 | Strengths and limitations

This study has several limitations. First, the main analysis was not performed in patients with a physician-made diagnosis of asthma. However, we did not solely rely on self-reported asthma (as asthma-self-reporting tends to be associated with the overestimation or the underestimation of asthma) but rather classified patients as having "probable asthma", "possible asthma" and "no evidence of asthma" based on self-reported asthma status, use of asthma medication and daily asthma symptoms. In a previous study assessing a sample of patients with physician-confirmed diagnosis, we observed that patients clustered as having "probable asthma" had a physician diagnosis of current or past asthma in 92.3% of cases, while patients clustered as having "no evidence of asthma" had a physician diagnosis of "no current asthma" in 90.4% of cases.[18] In addition, in this study, we assessed a sample of patients in whom the diagnosis of asthma was established by a physician (Twinning), obtaining consistent results with those of our main analysis. Of note, we did not solely assess patients from asthma clinics with a confirmed diagnosis of asthma since we wanted to have as many patients as possible and in a real-life context.

All assessed patients displayed self-reported rhinitis. Therefore, results are valid only for patients with nasal symptoms, who do however represent a very large proportion of patients with asthma.[23]
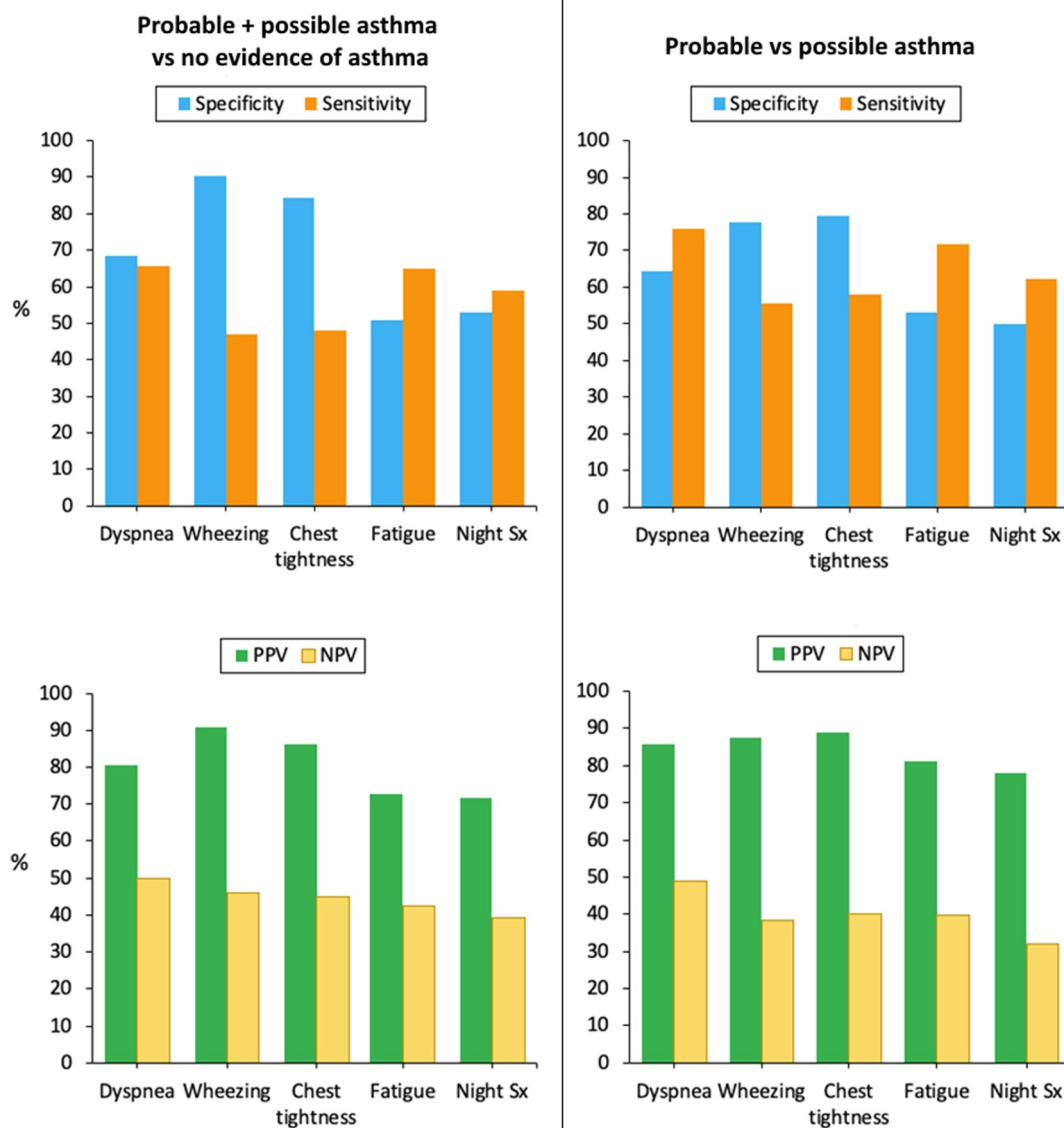
FIGURE 2 Sensitivity, specificity, positive predictive value and negative predictive value of the occurrence of specific asthma symptoms (Sx; as assessed by the control of allergic rhinitis and asthma test) on asthma diagnosis.

We could not test patients with asthma and without rhinitis as they represent less than 10% of patients with asthma in the MASK-air® database.

Some of the symptoms examined (e.g. dyspnea, wheezing, chest tightness, and fatigue) are rather nonspecific and may be the expression of different diseases (e.g. asthma and chronic obstructive pulmonary disease (COPD)), and also of the response to treatment in terms of control.[24]

Additionally, we were not able to assess all asthma symptoms. For example, we did not have information on the presence of cough and secretions. Nevertheless, secretions appear to be less common than wheezing, dyspnea, chest tightness and sleep disturbances. Importantly, treatments received prior to or during the time when the patients used the MASK-air app may have influenced the relationship between diagnosis and symptoms.

The tools used have been validated. CARAT is a validated questionnaire of asthma.[16] CARAT displays (i) an internal consistency similar to that of the Asthma Control Test (ACT), (ii) a reliability similar to that of the Asthma Control Questionnaire (ACQ) and higher than that of ACT, (iii) a correlation with clinical assessment of asthma control higher than that of ACT (despite being lower than that of ACQ) and (iv) an AUC-ROC for asthma higher than that of the ACQ.[16] e-DASTHMA is a daily data-driven asthma control medication score based on MASK-air® data. It is strongly correlated with daily dyspnea symptoms and moderately correlated with work- and quality-of-life-related comparators. It had high test-retest reliability and displayed moderate-to-high responsiveness. e-DASTHMA was validated in an external cohort of asthma patients enrolled by physicians[25] and was associated with the GINA classification of asthma control.[1]

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TABLE 5 Comparison between patients classified as having "probable asthma" versus "possible asthma" on the ever occurrence of asthma symptoms, as assessed by the control of allergic rhinitis and asthma test.

CARAT questions	Sensitivity—% (95% CI)	Specificity—% (95% CI)	PPV—% (95% CI)	NPV—% (95% CI)
A. First CARAT assessment by user				
Shortness of breath/dyspnea[a]	76.1 (72.2–79.9)	64.5 (57.2–71.7)	85.8 (82.4–89.1)	48.9 (42.2–55.5)
Wheezing in the chest	55.6 (51.1–60.1)	77.7 (71.4–84.0)	87.5 (83.8–91.3)	38.3 (33.1–43.5)
Chest tightness	57.9 (53.4–62.4)	79.5 (73.4–85.7)	88.9 (85.3–92.4)	40.1 (34.8–45.4)
Tiredness/limitations doing tasks	71.6 (67.5–75.7)	53.0 (45.4–60.6)	81.1 (77.3–84.9)	39.8 (33.7–46.3)
Night symptoms	62.4 (58.0–66.8)	50.0 (42.4–57.6)	77.9 (73.7–82.1)	32.0 (26.4–37.7)
B. All CARAT assessments				
Shortness of breath/dyspnea[b]	80.9 (78.9–82.9)	57.6 (51.6–63.7)	91.6 (90.1–93.1)	34.7 (30.1–39.2)
Wheezing in the chest	54.0 (51.4–56.6)	73.7 (68.3–79.1)	92.1 (90.3–93.9)	22.0 (19.2–24.7)
Chest tightness	63.2 (60.7–65.7)	69.0 (63.3–74.7)	92.1 (90.4–93.8)	24.8 (21.6–28.0)
Tiredness/limitations doing tasks	76.4 (74.2–78.7)	58.8 (52.8–64.9)	91.4 (89.8–92.9)	30.4 (26.4–34.5)
Night symptoms	59.4 (56.9–62.0)	55.3 (49.1–61.4)	88.3 (86.3–90.3)	19.3 (16.5–22.2)

Abbreviations: NPV, negative predictive value; PPV, positive predictive value.

[a]Considering data before 2020 only: Sensitivity = 69.3%; Specificity = 70.8%; PPV = 87.8%; NPV = 43.2%.

[b]Considering data before 2020 only: Sensitivity = 71.0%; Specificity = 67.6%; PPV = 92.0%; NPV = 30.9%.

We performed the analysis on the first CARAT questionnaires reported and used all CARAT questionnaires as a sensitivity analysis.

4.2 | Interpretation of the results

The first important message is the significance of wheezing in the diagnosis of asthma. Although PPV is influenced by disease prevalence, the high PPV of wheezing for asthma diagnosis is consistent with a study in a secondary care center with a precise asthma diagnosis.[26] However, wheezing lacks sensitivity and 45% of patients with asthma did not report this symptom. Chest tightness displayed a high PPV and a high specificity. It was the symptom showing the highest specificity in differentiating probable from possible asthma. By contrast, dyspnea was the symptom with the highest sensitivity. Despite this result, sensitivity was close to 80%, indicating that approximately one fifth of asthma patients do not experience dyspnea, which likely reflects the good control of the disease in that subset of patients. According to the SpIN and SnOUT rules,[27] a test with high specificity, when positive, helps to rule in the disease. This is the case of wheezing and chest tightness. On the other hand, a test with high sensitivity, when negative, helps to rule out the disease. This may be the case for dyspnea.

The second important message is the significance of dyspnea in asthma control assessment. Dyspnea is a multidimensional, subjective perception of breathing difficulty, commonly seen in patients affected by respiratory diseases, among others.[28] Although dyspnea perception may differ between subjects[29] (decreasing with the worsening of asthma, in older adults or in patients with depression[30]), the present study suggests that it is a cardinal symptom associated with asthma

control. In a previous MASK-air® study, VAS asthma was correlated with VAS dyspnea in patients with different degrees of severity, with a particularly strong correlation being observed in severe asthmatic patients.[31] Dyspnea was the symptom which mostly affected the asthma-related quality of life in both mild[26] and severe asthma patients. In a large cross-sectional study encompassing the whole disease severity spectrum, asthma-related quality of life was essentially determined by the level of asthma control.[32] These results were also confirmed in a longitudinal study.[33] Fatigue and night symptoms were much less specific, and their relationship with asthma control appeared less solid than that of dyspnea and chest tightness.

Future studies may assess whether individual respiratory symptoms can also be useful in the distinction between asthma and other respiratory diseases (namely, COPD).

4.3 | Generalizability

This study was conducted with patients from 27 countries. Previous MASK-air® studies have pointed to a high similarity of outcomes in different countries.[34]

5 | CONCLUSION

Among MASK-air® users who have rhinitis, wheezing and chest tightness are the symptoms that best predict the presence of asthma (displaying high specificity and PPV), while dyspnea is the symptom that showed the highest sensitivity and the strongest relationship
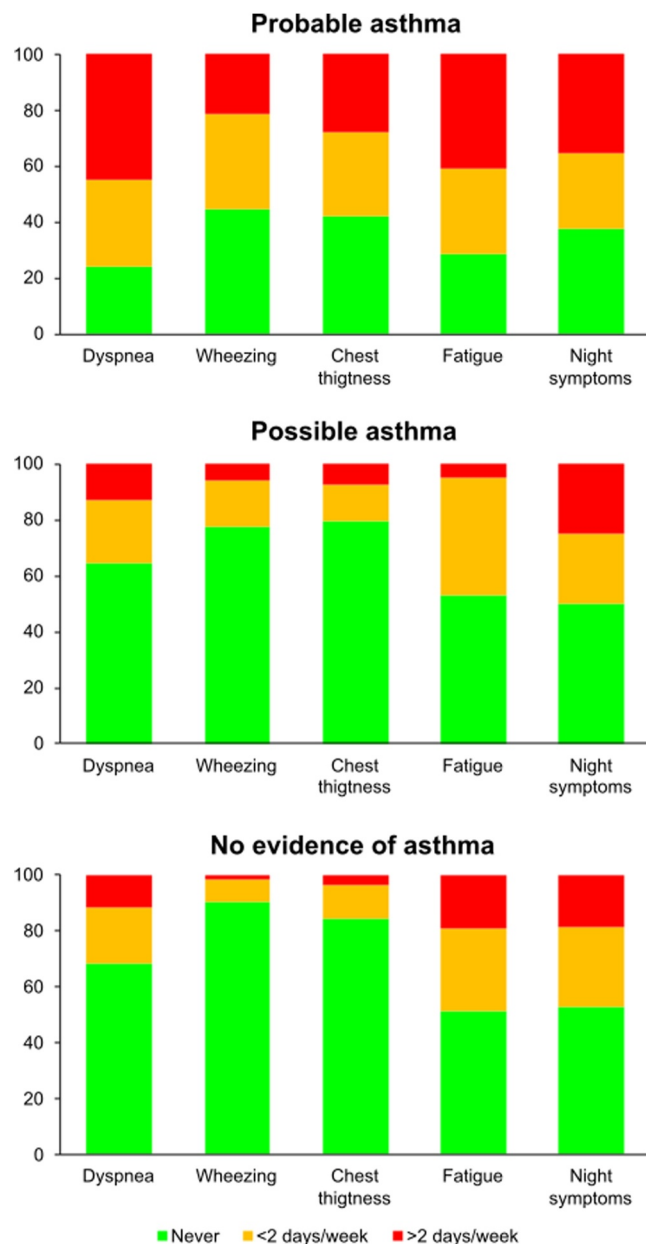
FIGURE 3 Frequency of symptoms reported by the control of allergic rhinitis and asthma test in patients with probable asthma.

with the level of asthma control. These results highlight the importance of thoroughly assessing individual symptoms during both the initial and follow-up assessments of patients with suspected or confirmed asthma.

THE MASK-AIR THINK TANK

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TABLE 6 Comparison between participants of the Twinning project classified as having "current asthma" versus "no evidence of asthma" on the ever occurrence of asthma symptoms, as assessed by the control of allergic rhinitis and asthma test.

CARAT questions	Sensitivity—% (95% CI)	Specificity—% (95% CI)	PPV—% (95% CI)	NPV—% (95% CI)
Shortness of breath/dyspnea	76.3 (67.8–84.8)	73.7 (67.3–80.0)	60.2 (51.5–68.8)	85.6 (80.2–91.1)
Wheezing in the chest	63.9 (54.4–73.5)	86.0 (81.0–91.0)	70.5 (60.9–80.0)	82.1 (76.7–87.4)
Chest tightness	55.7 (45.8–65.6)	80.1 (74.4–85.8)	59.3 (49.2–69.4)	77.6 (71.7–83.5)
Tiredness/limitations doing tasks	64.9 (55.5–74.4)	65.1 (58.2–71.9)	49.2 (40.6–57.9)	78.1 (71.5–84.6)
Night symptoms	47.4 (37.5–57.4)	68.8 (62.2–75.5)	44.2 (34.7–53.8)	71.5 (64.9–78.1)

Abbreviations: NPV, negative predictive value; PPV, positive predictive value.

TABLE 7 Median e-DASTHMA in patients with probable asthma according to the frequency of specific asthma symptoms as assessed by the control of allergic rhinitis and asthma test.

CARAT questions	Never—median (IQR) [N]	Up to 2 days per week—median (IQR) [N]	More than 2 days per week—median (IQR) [N]	Everyday or almost—median (IQR) [N]	Effect size			
					Never versus < 2 days/ week	<2 days versus +2 days/ week	+2 days/week versus every day or almost	Never versus any day/ week
A. First CARAT assessment by user (N = 468)								
Shortness of breath/dyspnea	5.7 (9.5) [112]	12.3 (18.8) [145]	20.9 (24.6) [107]	27.8 (30.0) [104]	0.64	0.56	0.34	0.97
Wheezing in the chest	10.2 (18.0) [208]	15.2 (20.5) [161]	27.1 (34.3) [55]	29.1 (23.7) [44]	0.37	0.55	0.08	0.53
Chest tightness	8.3 (18.0) [197]	14.7 (18.0) [141]	31.0 (28.9) [60]	26.5 (26.9) [70]	0.55	0.85	−0.20	0.77
Tiredness/task limitations	8.1 (11.8) [133]	13.7 (22.4) [144]	19.6 (24.3) [93]	24.5 (26.8) [98]	0.45	0.37	0.27	0.74
Night symptoms	9.1 (17.8) [176]	13.4 (19.7) [126]	20.1 (29.4) [94]	24.4 (25.6) [72]	0.36	0.37	0.21	0.63
B. All CARAT assessments (N = 1452)								
Shortness of breath/dyspnea	6.8 (11.3) [277]	12.3 (14.9) [436]	31.2 (31.4) [350]	35.6 (21.7) [389]	0.57	1.07	0.22	0.93
Wheezing in the chest	13.3 (19.1) [668]	18.8 (31.7) [488]	39.8 (32.0) [190]	38.1 (26.3) [106]	0.34	1.20	−0.10	0.65
Chest tightness	13.5 (21.5) [534]	15.0 (17.7) [370]	38.3 (29.6) [296]	33.5 (36.2) [252]	0.11	1.61	−0.22	0.45
Tiredness/task limitations	10.1 (12.3) [343]	12.7 (18.5) [510]	33.4 (26.1) [267]	33.4 (23.4) [332]	0.24	1.25	0	0.73
Night symptoms	17.8 (36.5) [589]	20.6 (23.0) [324]	22.5 (29.5) [153]	13.2 (25.0) [386]	0.14	0.10	−0.56	0.03

Note: IQR, interquartile range; Effect size: small: 0.20–0.49, medium: 0.50–0.79, high ≥0.80.

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DATA AVAILABILITY STATEMENT

Individual participant data underlying the results reported in this Article can be made available (after de-identification) between 12 and 36 months after Article publication. These data can be supplied to researchers who provide a methodologically sound proposal. Proposals should be directed to the corresponding author (jean.bousquet@orange.fr). We made every effort to follow the EU General Data Protection Regulation; therefore, we can transfer data only if there is a protocol and an agreement between the owner of the data and the person (or institution) requesting the data. To gain access, data requestors will need to sign a data access agreement.

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REFERENCES

1. Reddel HK, Bacharier LB, Bateman ED, et al. Global Initiative for Asthma Strategy 2021: executive summary and rationale for key changes. *Eur Resp J*. 2021;59(1):2102730.
2. Aaron SD, Boulet LP, Reddel HK, Gershon AS. Underdiagnosis and overdiagnosis of asthma. *Am J Respir Crit Care Med*. 2018;198(8):1012-1020. https://doi.org/10.1164/rccm.201804-0682ci
3. Monteiro C, Maricoto T, Prazeres F, Augusto Simoes P, Augusto Simoes J. Determining factors associated with inhaled therapy adherence on asthma and COPD: a systematic review and meta-analysis of the global literature. *Respir Med*. 2022;191:106724. https://doi.org/10.1016/j.rmed.2021.106724
4. Pike KC, Levy ML, Moreiras J, Fleming L. Managing problematic severe asthma: beyond the guidelines. *Arch Dis Child*. 2018;103(4):392-397. https://doi.org/10.1136/archdischild-2016-311368

5. Jansen EM, van de Hei SJ, Dierick BJH, Kerstjens HAM, Kocks JWH, van Boven JFM. Global burden of medication non-adherence in chronic obstructive pulmonary disease (COPD) and asthma: a narrative review of the clinical and economic case for smart inhalers. *J Thorac Dis*. 2021;13(6):3846-3864. https://doi.org/10.21037/jtd-20-2360

6. Downes E, Horigan A, Teixeira P. The transformation of health care for patients: information and communication technology, digiceuticals, and digitally enabled care. *J Am Assoc Nurse Pract*. 2019;31(3):156-161. https://doi.org/10.1097/jxx.0000000000000109

7. Black N. Patient reported outcome measures could help transform healthcare. *BMJ*. 2013;346(jan28 1):f167. https://doi.org/10.1136/bmj.f167

8. Bosnic-Anticevich S, Bakerly ND, Chrystyn H, Hew M, van der Palen J. Advancing digital solutions to overcome longstanding barriers in asthma and COPD management. *Patient Prefer Adherence*. 2023;17:259-272. https://doi.org/10.2147/ppa.s385857

9. Farzandipour M, Nabovati E, Sharif R, Arani MH, Anvari S. Patient self-management of asthma using mobile health applications: a systematic review of the functionalities and effects. *Appl Clin Inf*. 2017;8(4):1068-1081. https://doi.org/10.4338/aci-2017-07-r-0116

10. Bousquet J, Bedbrook A, Czarlewski W, et al. Guidance to 2018 good practice: ARIA digitally-enabled, integrated, person-centred care for rhinitis and asthma. *Clin Transl Allergy*. 2019;9(1):16. https://doi.org/10.1186/s13601-019-0252-0

11. Sousa-Pinto B, Fonseca JA, Gemicioglu B, et al. Patient-reported outcome measures (PROMs) using the MASK-air(R) app in severe asthma. *Allergy*. 2022;77(5):1600-1602. https://doi.org/10.1111/all.15248

12. Sousa-Pinto B, Sa-Sousa A, Amaral R, et al. Assessment of the control of allergic rhinitis and asthma test (CARAT) using MASK-air. *J Allergy Clin Immunol Pract*. 2022;10(1):343.e2-345.e2. https://doi.org/10.1016/j.jaip.2021.09.012

13. Sousa-Pinto B, Sa-Sousa A, Vieira RJ, et al. Cut-off values of MASK-air(R) patient-reported outcome measures (PROMs). *J Allergy Clin Immunol Pract*. 2023;11(4):1281-1289.e5.

14. Azevedo P, Correia de Sousa J, Bousquet J, et al. Control of allergic rhinitis and asthma test (CARAT): dissemination and applications in primary care. *Prim Care Respir J*. 2013;22(1):112-116. https://doi.org/10.4104/pcrj.2013.00012

15. Fonseca JA, Nogueira-Silva L, Morais-Almeida M, et al. Validation of a questionnaire (CARAT10) to assess rhinitis and asthma in patients with asthma. *Allergy*. 2010;65(8):1042-1048. https://doi.org/10.1111/j.1398-9995.2009.02310.x

16. Vieira R, Sousa-Pinto B, Cardoso-Fernandes A, et al. The Control of Allergic Rhinitis and Asthma Test (CARAT): a systematic review of its measurement properties and COSMIN analysis. *Clin Transl Allergy*. 2022;12(9):e12194. https://doi.org/10.1002/clt2.12194

17. Louis R, Satia I, Ojanguren I, et al. European respiratory society guidelines for the diagnosis of asthma in adults. *Eur Respir J*. 2022;60(3):2101585. https://doi.org/10.1183/13993003.01585-2021

18. Bousquet J, Sousa-Pinto B, Anto J, et al. Identification by cluster analysis of patients with asthma and nasal symptoms using the MASK-air® mHealth app. *Pulmonology*. 2023;29(4):292-305. https://doi.org/10.1016/j.pulmoe.2022.10.005

19. Bousquet J, Arnavielhe S, Bedbrook A, et al. Mask 2017: ARIA digitally-enabled, integrated, person-centred care for rhinitis and asthma multimorbidity using real-world-evidence. *Clin Transl Allergy*. 2018;8(1):45. https://doi.org/10.1186/s13601-018-0227-6

20. Bousquet J, Anto JM, Bachert C, et al. ARIA digital anamorphosis: digital transformation of health and care in airway diseases from research to practice. *Allergy*. 2021;76(1):168-190. https://doi.org/10.1111/all.14422

21. Bousquet J, Agache I, Aliberti MR, et al. Transfer of innovation on allergic rhinitis and asthma multimorbidity in the elderly (MACVIA-ARIA) - reference Site Twinning (EIP on AHA). *Allergy*. 2017;73(1):77-92. https://doi.org/10.1111/all.13218

22. Sousa-Pinto B, Jacome C, Pereira AM, et al. Development and validation of an electronic daily control score for asthma (e-DASTHMA): a real-world direct patient data study. *Lancet Digit Health*. 2023;5(4):e227-e238. https://doi.org/10.1016/s2589-7500(23)00020-1

23. Leynaert B, Neukirch C, Kony S, et al. Association between asthma and rhinitis according to atopic sensitization in a population-based study. *J Allergy Clin Immunol*. 2004;113(1):86-93. https://doi.org/10.1016/j.jaci.2003.10.010

24. Agusti A, Rapsomaniki E, Beasley R, et al. Treatable traits in the NOVELTY study. *Respirology*. 2022;27(11):929-940. https://doi.org/10.1111/resp.14325

25. Amaral R, Jacome C, Almeida R, et al. Profiling persistent asthma phenotypes in adolescents: a longitudinal diagnostic evaluation from the INSPIRERS studies. *Int J Environ Res Publ Health*. 2021;18(3):1015. https://doi.org/10.3390/ijerph18031015

26. Louis G, Schleich F, Louis R, et al. How respiratory symptoms impact asthma-related quality of life in mild asthmatics. *Respir Med*. 2023;207:107098. https://doi.org/10.1016/j.rmed.2022.107098

27. Baeyens JP, Serrien B, Goossens M, Clijsen R. Questioning the "SPIN and SNOUT" rule in clinical testing. *Arch Physiother*. 2019;9(1):4. https://doi.org/10.1186/s40945-019-0056-5

28. Parshall MB, Schwartzstein RM, Adams L, et al. An official American Thoracic Society statement: update on the mechanisms, assessment, and management of dyspnea. *Am J Respir Crit Care Med*. 2012;185(4):435-452. https://doi.org/10.1164/rccm.201111-2042st

29. Barbaro MP, Lacedonia D, Palladino GP, et al. Dyspnea perception in asthma: role of airways inflammation, age and emotional status. *Respir Med*. 2011;105(2):195-203. https://doi.org/10.1016/j.rmed.2010.09.013

30. Ohwada A, Sato K. Heterogeneity of perception of symptoms in patients with asthma. *J Thorac Dis*. 2019;11(12):5218-5227. https://doi.org/10.21037/jtd.2019.11.72

31. Benfante A, Sousa-Pinto B, Pillitteri G, et al. Applicability of the MASK-air® app to severe asthma: a pilot longitudinal study. *Int J Mol Sci*. 2022;23(19):11470. https://doi.org/10.3390/ijms231911470

32. Louis G, Petre B, Schleich F, et al. Predictors of asthma-related quality of life in a large cohort of asthmatics: a cross-sectional study in a secondary care center. *Clin Transl Allergy*. 2021;11(7):e12054. https://doi.org/10.1002/clt2.12054

33. Louis G, Petre B, Schleich F, et al. Predictors of change in asthma-related quality of life: a longitudinal real-life study in adult asthmatics. *Qual Life Res*. 2023;32(5):1507-1520. https://doi.org/10.1007/s11136-022-03339-0

34. Bousquet J, Anto JM, Sousa-Pinto B, et al. Digitally-enabled, patient-centred care in rhinitis and asthma multimorbidity: the ARIA-MASK-air((R)) approach. *Clin Transl Allergy*. 2023;13(1):e12215. https://doi.org/10.1002/clt2.12215

SUPPORTING INFORMATION

Additional supporting information can be found online in the Supporting Information section at the end of this article.