

Suitability of Forced Expiratory Volume in 1 Second/Forced Vital Capacity vs Percentage of Predicted Forced Expiratory Volume in 1 Second for the Classification of Asthma Severity in Adolescents

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Objective: To determine whether lung function alters asthma severity based on symptom history in asthmatic adolescents.

Design: Data on asthma symptoms and lung function were collected from adolescents randomly selected from the general population.

Setting: Five schools from the central Wellington, New Zealand, area during 2003 to 2005.

Participants: Two hundred twenty-four secondary school students aged 13 to 17 years (asthmatic, 118; non-asthmatic, 106).

Main Exposures: Asthma questionnaire and lung function testing.

Main Outcome Measures: Distribution of asthmatic adolescents in each severity class based on symptoms, lung function, or a combination of both.

Results: Median values for all spirometric parameters for asthmatic adolescents, apart from forced expiratory vol-

ume in the first second of expiration (FEV₁)/forced vital capacity (FVC), were in the normal range. Distribution of severity (based on symptoms and β_2 -agonist use with adjustment for regular inhaled corticosteroid use) was 48.3%, mild; 28.8%, moderate; and 22.9%, severe asthma. For severity based on percentages of predicted FEV₁ and predicted forced expiratory flow, midexpiratory phase (FEF_{25%-75%}) and FEV₁/FVC, the percentages were 89.8%, 86.4%, and 63.5%, mild; 9.3%, 10.2%, and 18.6%, moderate; and 0.9%, 3.4%, and 17.8%, severe asthma, respectively. When percentages of predicted FEV₁ or predicted FEF_{25%-75%} or FEV₁/FVC were added to symptom severity, 6.8%, 5.1%, and 16.9% of asthmatic adolescents were reclassified into another severity group, respectively.

Conclusions: The majority of asthmatic adolescents have normal lung function despite experiencing significant asthma symptoms. Adding FEV₁/FVC to symptom history changes the distribution of severity; however, both percentages of predicted FEV₁ and FEF_{25%-75%} have little added effect in assessing asthma severity in adolescents.

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ASTHMA GUIDELINES HAVE been developed to aid in the evaluation of asthma severity and control and assist in determining optimum treatment regimens. Invariably these guidelines include a measure of lung function, commonly forced expiratory volume in the first second of expiration (FEV₁) or peak expiratory flow rate (PEF),^{1,2} in addition to asthma symptoms. Because patient symptom recall is generally considered unreliable, with approximately 25% of patients either underestimating or overestimating the severity of their symptoms,³ an objective measure of lung function is thought to strengthen the subjective assessment of asthma severity provided by the clinical his-

tory. Several studies of asthma severity in children have shown that lung function, and in particular FEV₁, is in the normal range in the majority of children, including in children with severe asthma.⁴⁻⁶ In addition, asthma symptom frequency correlates poorly with FEV₁ and PEF. It has been suggested that it may be more appropriate to use FEV₁/forced vital capacity (FVC) or forced expiratory flow, midexpiratory phase (FEF_{25%-75%}), which measures airflow in the smaller airways, for objective assessment of asthma severity in children and adolescents.^{4,6} The National Asthma Education and Prevention Program Expert Panel Report 3 (NAEPP EPR-3)⁷ has responded to these findings and no longer recommends PEF in the assessment of asthma severity.

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Table 1. Classification of Asthma Severity Symptom and β_2 -Agonist Use

Symptom/Medication Use	Severity Class	Frequency, No. (%)
No. of wheezing attacks in the last 12 mo		
0-3	Mild	84 (71.2)
4-12	Moderate	23 (19.5)
>12	Severe	11 (9.3)
Sleep disturbance due to wheeze in the last 12 mo, nights/wk		
<1	Mild	112 (94.9)
≥ 1	Moderate	6 (5.1)
Daily symptoms in the last 2 wk	Moderate	13 (11.0)
Daily β_2 -agonist use in the last 2 wk	Moderate	32 (27.1)

FEV₁/FVC has been included together with percentage of predicted FEV₁. It remains unclear as to whether an assessment of lung function aids in determining asthma severity, particularly in the child and adolescent age groups. Including spirometry adds significant extra work for the already busy practitioner, and indeed, it has been shown that PEF is more commonly used by general practitioners rather than spirometry to assess asthma severity.⁸ We have conducted a study investigating asthma disease severity in a random general population sample of asthmatic adolescents to determine whether adding spirometric parameters (percentage of predicted FEV₁, FEV₁/FVC, and percentage of predicted FEF_{25%-75%}) to a disease severity classification system based solely on symptom frequency and β_2 -agonist use alters the assessment of disease severity sufficiently to justify its use.

METHODS

SUBJECT RECRUITMENT

Subjects were recruited from 5 schools that had participated in the Wellington International Study of Asthma and Allergies in Childhood (ISAAC) Phase III survey of asthma symptom prevalence.⁹ These schools were chosen because of their proximity to the study center. Potential asthmatic adolescents (n=224) and nonasthmatic adolescents (n=595) were identified from the ISAAC Phase III study from their response to the question "Have you had wheezing or whistling in the chest in the past 12 months?" From this group, 136 asthmatic adolescents and 134 nonasthmatic adolescents consented to participate in a further study. At the study visit, a further questionnaire based on the ISAAC Phase III questionnaire and including additional questions on medication use for the 12-month and 2-week period prior to the clinic visit was completed. Final assignment to asthmatic or nonasthmatic status was determined from this questionnaire. Asthmatic adolescents had a history of wheezing or whistling in the chest in the last 12 months and/or any asthma medication use in the last 12 months. Nonasthmatic adolescents had no current wheeze, no history of asthma, no nocturnal cough in the last 12 months apart from that associated with a cold or chest infection, and no asthma medication use in the past 12 months. Sixteen subjects were excluded because they did not meet the definitions for asthma or nonasthma either because of a history of asthma but no current symptoms or an ambiguous response to the relevant questions. A further 8 subjects were excluded because they were unable to be contacted after indicating a willingness to participate in the study, 11 were unable to perform adequate spirometry, and

11 were excluded because of medication use, such as β_2 -agonist, within 6 hours prior to spirometry.

The study visit was deferred if subjects had had an acute exacerbation of asthma or respiratory tract infection in the previous 4 weeks. Subjects with chronic illnesses other than asthma were ineligible. School decile rating was used as a proxy for socioeconomic status, ranging from 1-10, with a rating of 10 equating to the highest socioeconomic group.

Ethical approval for the study was obtained from the Wellington Ethics Committee (00/03/010). Written consent was obtained from school principals and from each child in the study and their parents.

SPIROMETRY

Lung function was measured as the best of 3 reproducible forced expiratory maneuvers (EasyOne Spirometer; ndd Medizintechnik AG, Zurich, Switzerland). The spirometer was calibrated daily with a 3-L calibration syringe (Hans Rudolph, Inc, Kansas City, Missouri). Predicted values for lung function parameters were determined using the National Health and Nutrition Examination Survey III prediction equations.¹⁰

SYMPTOM AND LUNG FUNCTION CLASSIFICATION OF SEVERITY

Asthma symptoms were classified as mild, moderate, or severe based on the NAEPP EPR-3 guidelines.⁷ This included frequency of wheezing attacks, nocturnal symptoms, and frequency of β_2 -agonist use and daily symptoms in the 2 weeks preceding the study visit (unadjusted symptom severity) (**Table 1**). Adjustment was made for regular inhaled corticosteroid (ICS) use by moving the severity level up one step (ie, mild to moderate and moderate to severe [ICS-adjusted symptom severity]). For asthma severity using percentage of predicted FEV₁, FVC, and PEF, mild asthma was defined as 80% or more; moderate asthma, as 60% or more and less than 80%; and severe asthma, less than 60%; for percentage of predicted FEF_{25%-75%}, mild, moderate, or severe asthma was defined as 65% or more, 50% or more and less than 65%, and less than 50%, respectively^{6,10}; and for FEV₁/FVC, mild, moderate, or severe asthma was defined as 80% or more, 75% or more and less than 80%, and less than 75%, respectively, using the NAEPP ERP-3 guidelines cutoff points.⁷

DATA ANALYSIS

All analyses were carried out using SAS (SAS Institute Inc, Cary, North Carolina). Lung function was expressed as median values and interquartile range. Comparisons between groups were made using the χ^2 test and Fisher exact test.¹¹ The Wilcoxon rank sum test was used to determine significant differences between nonparametric data. A κ value was calculated to assess the agreement between tests. A *P* value of <.05 was considered significant.

RESULTS

SUBJECT DEMOGRAPHICS

There was no significant difference in the distributions of the asthmatic adolescents and nonasthmatic adolescents by age (mean, 14.9 and 15.1 years, respectively), sex (26% and 32%, female, respectively), or ethnicity (81% and 83%, European, respectively). This study included

Table 2. Spirometry Results for Asthma Symptom Severity Groups, Asthmatic and Nonasthmatic

	Median (Interquartile Range)				
	Mild (n = 57)	Moderate (n = 34)	Severe (n = 27)	Asthma (n = 118)	Nonasthma (n = 106)
FEV ₁ , % predicted	96.9 (18.8)	97.8 (14.5)	95.8 (18.3)	96.9 ^a (17.1)	101.2 (15.1)
FVC, % predicted	98.7 (18.8)	100.2 (18.7)	101.6 (13.0)	99.3 (16.5)	100.4 (17.9)
FEV ₁ /FVC, %	84.3 ^b (9.3)	83.1 (8.7)	78.5 (10.5)	83.2 ^c (10.5)	86.0 (9.1)
FEF _{25%-75%} , % predicted	90.4 ^b (26.8)	87.5 (32.2)	75.8 (41.3)	86.5 ^c (32.0)	100.9 (26.8)
PEF, % predicted	100.6 (19.1)	106.1 (23.3)	96.0 (19.9)	100.6 (22.4)	101.1 (20.0)

Abbreviations: FEF_{25%-75%}, forced expiratory flow, midexpiratory phase; FEV₁, forced expiratory volume in the first second of expiration; FVC, forced vital capacity; PEF, peak expiratory flow rate.

^a $P < .01$ asthma vs nonasthma.

^b $P < .05$ mild asthma vs severe asthma.

^c $P < .001$ asthma vs nonasthma.

subjects attending schools with decile ratings ranging from 5 to 10 (ie, lower socioeconomic groups [school decile 1-4] were not included). There was no difference in the distribution of asthmatic adolescents and nonasthmatic adolescents in school decile rating. Five percent of asthmatic adolescents were smokers compared with 1% of nonasthmatic adolescents. Just more than half (52.5%) of asthmatic adolescents reported ICS use at some time; however, only 24% reported regular ICS use. Of the 118 asthmatic adolescents included in the study, 89% responded positively to the question "Have you ever had asthma?"

BASELINE SPIROMETRY

For the nonasthmatic adolescents, the percentages of predicted FEV₁ and FVC values were all very close to 100% (**Table 2**), suggesting that the reference values from the National Health and Nutrition Examination Survey were appropriate for use in New Zealand adolescents.¹⁰ Asthmatic adolescents had significantly lower percentages of predicted values for FEV₁ and FEF_{25%-75%} and lower values for FEV₁/FVC than nonasthmatic adolescents but not for percentages of predicted FVC and PEF (Table 2). The median values for asthmatic adolescents for all spirometric parameters, apart from FEV₁/FVC, were in the normal range (ie, for percentages of predicted FEV₁, FVC, and PEF, $\geq 80\%$; for percentage of predicted FEF_{25%-75%}, $\geq 65\%$) (Table 2). For the complete asthmatic group and all levels of asthma severity, FEV₁/FVC was less than 85%, the value given as normal for FEV₁/FVC in the NAEPP EPR-3 guidelines for this age group.

SEVERITY AND LUNG FUNCTION

The median values for FEV₁/FVC and for percentage of predicted FEF_{25%-75%} were significantly lower in the severe asthmatic adolescents compared with the mild asthmatic adolescents (Table 2). There were no significant differences among the 3 symptom severity groups for percentage of predicted FEV₁, percentage of predicted FVC, and percentage of predicted PEF and no significant differences between moderate and severe asthmatic adolescents for all spirometric parameters.

ASTHMA SEVERITY DISTRIBUTION

Using the adjusted symptom severity definition, 48.3% of asthmatic adolescents had mild, 28.8% had moderate, and 22.9% had severe symptoms (**Table 3**). Without adjustment for regular ICS use, the distribution was 54.2%, mild, 36.4%, moderate, and 9.3%, severe. The distribution of severity based on percentage of predicted FEV₁ alone was mild, 89.8%; moderate, 9.3%; and severe, 0.9%; on FEV₁/FVC was mild, 63.5%; moderate, 18.6%; and severe, 17.8%; and on percentage of predicted FEF_{25%-75%} was mild, 86.4%; moderate, 10.2%; and severe, 3.4% (Table 3). When lung function was added to the adjusted symptom severity distribution, percentage of predicted FEV₁ or FEF_{25%-75%} caused little change in distribution, with only 6.8% and 5.1% of asthmatic adolescents changing severity, respectively. Adding percentage of predicted FEV₁ or FEF_{25%-75%} to the unadjusted symptom severity caused a change in severity for 7.6% and 4.1% of asthmatic adolescents, respectively. The change in distribution was much greater for FEV₁/FVC, with 16.9% of subjects changing severity for the adjusted symptom severity and 22.9%, for the unadjusted symptom severity. Combining both percentage of predicted FEV₁ and FEV₁/FVC caused a change in severity for 20.3% of asthmatic adolescents for the adjusted symptom severity and 26.3% for the unadjusted severity. κ Values were calculated to assess the agreement between the severity classifications. There was no agreement with symptom severity and severity based on percentage of predicted FEV₁ ($\kappa = -0.04$ for adjusted ICS severity and -0.08 for unadjusted severity) or on percentage of predicted FEF_{25%-75%} ($\kappa = 0.02$ for adjusted ICS severity and 0.09 for unadjusted severity) and minimal agreement with FEV₁/FVC ($\kappa = 0.18$ for adjusted ICS severity and 0.14 for unadjusted severity).

The disparities between the classification of asthma severity based on symptoms and that based on percentage of predicted FEV₁ and percentage of predicted FEF_{25%-75%} lie mainly in subjects with moderate and severe symptom classification having an FEV₁ of 80% or more predicted or FEF_{25%-75%} of 65% or more predicted. Specifically, 91.2% (31 of 34) of subjects with moderate symptom severity (adjusted for ICS use) had mild asthma ac-

Table 3. Effect of Adding Individual Variables on the Distribution of Asthma Symptom Severity^a

	No. (%)			κ
	Mild	Moderate	Severe	
Wheezing attacks in the last 12 mo	84 (71.2)	23 (19.5)	11 (9.3)	
+ Sleep disturbance due to wheeze in the last 12 mo	81 (68.6)	26 (22.0)	11 (9.3)	
+ Daily β_2 -agonist use in the last 2 wk	66 (55.9)	41 (34.8)	11 (9.3)	
+ Daily symptoms in the last 2 wk	64 (54.2)	43 (36.4)	11 (9.3)	
Regular ICS use	7 (10.9)	16 (37.2)	5 (45.5)	
Symptom severity adjusted for regular ICS use	57 (48.3)	34 (28.8)	27 (22.9)	
FEV ₁ , % predicted	106 (89.8)	11 (9.3)	1 (0.9)	-0.04
FEV ₁ /FVC	75 (63.5)	22 (18.6)	21 (17.8)	0.18
FEF _{25%-75%} , % predicted	102 (86.4)	12 (10.2)	4 (3.4)	0.02
Combined symptoms/ β_2 -agonist use/ICS use+				
FEV ₁ , % predicted	50 (42.4)	40 (33.9)	28 (23.7)	
FEV ₁ /FVC	42 (35.6)	38 (32.2)	38 (32.2)	
FEF _{25%-75%} , % predicted	52 (44.1)	36 (30.5)	30 (25.4)	
FEV ₁ and FEV ₁ /FVC, % predicted	38 (32.2)	42 (35.5)	38 (32.2)	

Abbreviations: FEF_{25%-75%}, forced expiratory flow, midexpiratory phase; FEV₁, forced expiratory volume in the first second of expiration; FVC, forced vital capacity; ICS, inhaled corticosteroid.

^aValues are expressed as percentage of total number of asthmatic adolescents (n = 118) and total number of adolescents. Symptoms were added in a stepwise fashion in order of increasing severity (as listed) to the frequency of baseline wheezing attacks in the last 12 months. Regular ICS use was adjusted for by moving severity up 1 step (ie, mild to moderate and moderate to severe). Each lung function parameter was added to the final combined symptoms/ β_2 -agonist use/regular ICS use severity distribution. For FEV₁, % predicted, mild, moderate, and severe asthma were defined as 80% or more, 60% or more and less than 80%, and less than 60%, respectively; for FEF_{25%-75%}, % predicted, as 65% or more, 50% or more and less than 65%, and less than 50%, respectively; and for FEV₁/FVC, as 80% or more, 75% or more and less than 80%, and less than 75%, respectively. Agreement between severity distribution was based on final combined symptoms/ β_2 -agonist use/regular ICS use and each lung function parameter was assessed using κ values.

cording to the percentage of predicted FEV₁ and percentage of predicted FEF_{25%-75%}, and 100% (27 of 27) and 96.3% (26 of 27) of subjects with severe symptom severity had mild asthma according to the percentage of predicted FEV₁ and percentage of predicted FEF_{25%-75%}, respectively. Of asthmatic adolescents with moderate symptom severity (adjusted for ICS use), 61.8% had mild asthma according to FEV₁/FVC and 63% with severe symptom severity had either mild or moderate asthma according to FEV₁/FVC.

COMMENT

In this study, we found that in the large majority of asthmatic adolescents in this group recruited from the general population lung function was within the normal range, apart from FEV₁/FVC, where median values for all levels of asthma severity were less than 85%. In determining disease severity, a single assessment of percentage of predicted FEV₁ or percentage of predicted FEF_{25%-75%} added little further information to the severity classification obtained using symptom questionnaire responses. In contrast, adding FEV₁/FVC to asthma symptoms altered the distribution of severity for 17% of subjects. For severity distribution based on all these 3 lung function parameters, there was minimal agreement with that based on asthma symptoms and β_2 -agonist use with and without adjustment for ICS use.

This study recruited a random sample of asthmatic adolescents and nonasthmatic adolescents from the general population rather than from subspecialty clinics. The findings of this study are therefore directly relevant to the general population and in particular to the use of spirometry in assessment and management of asthma in ado-

lescents in general practice. The findings may not be applicable to a population of asthmatic adolescents attending specialty clinics, where asthma symptoms and lung function are likely to be more severe than in this population. However, lung function has also been found to be normal in the majority of students recruited from tertiary care asthma clinics,^{4,6} suggesting that lung function is normal in the young asthmatic population throughout all levels of asthma severity. Similarly, a mismatch between asthma symptoms and lung function in adolescents with more severe asthma has been found, suggesting this finding is also true in general for young people with asthma.

In a cross-sectional population where a proportion of subjects are using medication to control their symptoms, classification of asthma severity should also include intensity of treatment, particularly ICS use. Regular ICS use is likely to reduce asthma symptom frequency and categorize an individual with milder asthma than when not using medication. We have adjusted for regular ICS use in this group of asthmatic adolescents by increasing the level of severity one step in those subjects taking their ICS regularly. We have based this adjustment on the findings of Bacharier et al⁴ (ie, that the determination of asthma severity in young people with asthma worsens when medication use is included with symptom frequency in severity assessment) and the recommendations in the 2005 Global Initiative for Asthma (GINA) guidelines.¹² It is possible that the adjustment we have made has defined more subjects with severe asthma than is actually the case. However, we also conducted our analyses without making an adjustment for regular ICS use (ie, using symptom frequency and β_2 -agonist use only in the definition of asthma symp-

tom severity). Naturally, fewer subjects were classified as having severe asthma with this classification (mild, 54%; moderate, 36%; and severe, 9%). The effect of adding the lung function parameters to symptom severity was greater for FEV₁/FVC, with 22.9% of asthmatic adolescents reclassified as having more severe asthma, minimal change for percentage of predicted FEV₁, and less for percentage of predicted FEF_{25%-75%}. Combining percentage of predicted FEV₁ with FEV₁/FVC resulted in a 26.3% change in severity as compared with a 20.3% change using the symptom severity adjusted for regular ICS use. The overall conclusion, therefore, that adding either percentage of predicted FEV₁ or percentage of predicted FEF_{25%-75%} to symptom severity causes minimal change in severity classification, is not altered by adjusting for regular ICS use in this study.

It may be argued that some of our subjects did not have asthma because we did not use a physician diagnosis of asthma to determine asthmatic status. Rather, we defined subjects as having current asthma based on a self-report of wheeze in the preceding 12 months. Using only a physician diagnosis of asthma excludes a significant proportion of the population with wheezing symptoms. This was shown in a large study of children aged 12 to 14 years, where 17% of children with current asthmalike symptoms did not have a diagnosis of asthma.¹² The ISAAC questionnaire, on which the questionnaire used in this study was based, has been shown to have both a high sensitivity and specificity for physician-diagnosed asthma.¹³ Another study of asthma diagnosis in New Zealand has shown a similarly high sensitivity and specificity of wheezing in the last 12 months for a diagnosis of asthma.¹⁴ Although it is possible that we have included some subjects without asthma as asthmatic adolescents, the survey instrument we used to identify those with asthma has been shown to have sufficiently high sensitivity and specificity to be used with confidence for this purpose.

It is possible that asthma severity has been misclassified because of inaccurate recall of symptoms by our subjects. We based our severity classification on both short-term recall (symptoms and β_2 -agonist use in the preceding 2 weeks), as is recommended in the NAEPP EPR-3 guidelines, and long-term recall (symptoms in the preceding 12 months). Long-term recall is likely to be inaccurate, particularly for recall of symptom frequency. However, using only 2 weeks' recall would not take into account the inherent variability of asthma symptoms that results in individuals with asthma having changes in severity over time. In an analysis of five 12-week asthma trials in pediatric subjects, it was found that subjects frequently moved between severity groups.¹⁵ We therefore elected to combine both short- and long-term recall of symptoms to determine symptom severity.

In a large study of 2 cohorts of children aged between 8 and 11 years, Stout et al¹⁶ found that approximately one-third of children were reclassified with higher asthma severity when FEV₁ was added to symptom frequency to assess asthma severity. This contrasts with our finding that using percentage of predicted FEV₁ does little to change severity classification as determined by symptoms. In the Stout et al study, asthma was defined as a physician diagnosis of asthma and either a history of

asthma symptoms or at least 1 hospitalization or at least 2 urgent care visits in the previous 6 months. We used a definition of asthma based on a self-reported history of current wheeze, and although the majority of subjects in our study appeared to have a previous asthma diagnosis, it is likely that our subjects had milder asthma than those in the Stout et al study. However, the unadjusted symptom severity distribution in our study was similar to that in the more severe cohort in the Stout et al study, where 57.4% of subjects had mild intermittent or persistent asthma and 42.6% of subjects had moderate or severe asthma. There was a difference between the 2 studies in the numbers of subjects demonstrating abnormal lung function (ie, a predicted FEV₁ < 80%), with 10.2% of subjects in our study and 16.7% and 28.2% in cohort 1 and cohort 2 of the Stout et al study, respectively. This suggests that overall lung function was better in our subjects and this likely accounts for the different effect of adding percentage of predicted FEV₁ to symptom severity in both studies.

Of the lung function parameters, only FEV₁/FVC and percentage of predicted FEF_{25%-75%} showed any significant difference in asthma groups based on symptom severity (Table 2). Similar results have been found in other studies of lung function in children.^{4,6} The authors of one of these studies have suggested that these parameters have greater sensitivity in measuring airflow obstruction and should be assessed when spirometry is performed.⁶ FEF_{25%-75%} measures the average flow rate over the middle 50% of the FVC and is said to be more sensitive than FEV₁ in detecting small-airway obstruction. However, we found that adding percentage of predicted FEF_{25%-75%} to asthma symptoms resulted in little change in severity classification and that there was also no agreement between asthma symptom severity and percentage of predicted FEF_{25%-75%}. In addition to this, percentage of predicted FEF_{25%-75%} demonstrates high variability, making its use for monitoring asthma progression difficult. These results suggest that percentage of predicted FEF_{25%-75%} is not a useful tool for assessing asthma severity in this age group.

The median value for FEV₁/FVC was less than 85%, the value defined as the cutoff for normality in the NAEPP EPR-3 guidelines for this age group, for the asthmatic population, and also for each symptom severity group. For the nonasthmatic group, the median value was 86%. These results suggest that a level of 85% for FEV₁/FVC may be overestimating the number of young people with asthma with airflow obstruction. The value of 85% is taken from the National Health and Nutrition Examination Survey control data for this age group.¹⁰ Because the percentage of predicted FEV₁ and FVC values for nonasthmatic adolescents in this study were all very close to 100%, suggesting that the reference values from the National Health and Nutrition Examination Survey were appropriate for use in New Zealand adolescents, it also suggests that a ratio of 85% for FEV₁/FVC is appropriate for use in this study. In the large National Jewish Medical and Research Center study of lung function in children, the average value for the FEV₁/FVC ratio was approximately 77% for males and 83% for females.⁶ This group of children was recruited from a tertiary specialist cen-

ter and therefore is likely to have had more severe asthma than the group in our study. The American Thoracic Society/European Respiratory Society task force has recommended using the fifth percentile of predicted values rather than set cutoff values for the FEV₁/FVC ratio to define obstructive lung disease.¹⁷ This is particularly important for younger and older adults, because the set cutoff ratio of 70% has been found to misclassify a significant number of subjects in these age groups.^{18,19} It may also be appropriate to apply this criterion to defining airway obstruction in adolescents rather than using the set cutoff ratio of 85%.

In conclusion, in the vast majority of asthmatic adolescents, a single measure of percentage of predicted FEV₁ or percentage of predicted FEF_{25%-75%} adds little extra information to that obtained from a symptom history in assessing asthma severity. FEV₁/FVC, however, does appear to add value to this assessment, suggesting that this parameter should be used in determining asthma severity in adolescents.

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Author Contributions: Dr van Dalen had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. *Study concept and design:* van Dalen, Pearce, and Douwes. *Acquisition of data:* van Dalen, Harding, Parkin, Cheng, and Douwes. *Analysis and interpretation of data:* van Dalen, Cheng, Pearce, and Douwes. *Drafting of the manuscript:* van Dalen, Pearce, and Douwes. *Critical revision of the manuscript for important intellectual content:* van Dalen, Harding, Parkin, Cheng, Pearce, and Douwes. *Statistical analysis:* van Dalen, Cheng, Pearce, and Douwes. *Obtained funding:* Pearce and Douwes. *Administrative, technical, and material support:* Harding and Parkin. *Study supervision:* van Dalen, Pearce, and Douwes.

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REFERENCES

1. The diagnosis and treatment of adult asthma. Wellington: New Zealand Guidelines Group; 2002. <http://www.nzgg.org.nz/guidelines/0003/Summary.pdf>. Accessed July 2, 2008.
2. GINA report, global strategy for asthma management and prevention [published 2007]. Global Initiative for Asthma Web site. <http://www.ginasthma.org>. Accessed July 2, 2008.
3. Nguyen BP, Wilson SR, German DF. Patients' perceptions compared with objective ratings of asthma severity. *Ann Allergy Asthma Immunol*. 1996;77(3):209-215.
4. Bacharier LB, Strunk RC, Mauger D, White D, Lemanske RF Jr, Sorkness CA. Classifying asthma severity in children: mismatch between symptoms, medication use, and lung function. *Am J Respir Crit Care Med*. 2004;170(4):426-432.
5. Fuhlbrigge AL, Kitch BT, Paltiel AD, et al. FEV₁ is associated with risk of asthma attacks in a pediatric population. *J Allergy Clin Immunol*. 2001;107(1):61-67.
6. Paull K, Covar R, Jain N, Gelfand EW, Spahn JD. Do NHLBI lung function criteria apply to children? A cross-sectional evaluation of childhood asthma at National Jewish Medical and Research Center, 1999-2002. *Pediatr Pulmonol*. 2005;39(4):311-317.
7. Expert Panel Report 3 (EPR-3): guidelines for the diagnosis and management of asthma—summary report 2007. *J Allergy Clin Immunol*. 2007;120(5)(suppl):S94-S138.
8. Finkelstein JA, Lozano P, Shulruff R, et al. Self-reported physician practices for children with asthma: are national guidelines followed? *Pediatrics*. 2000;106(4)(suppl):886-896.
9. Ellwood P, Asher MI, Beasley R, Clayton TO, Stewart AW; ISAAC Steering Committee. International Study of Asthma and Allergies in Childhood (ISAAC II): phase III rationale and methods. *Int J Tuberc Lung Dis*. 2005;9(1):10-16.
10. Hankinson JL, Odencrantz JR, Fedan KB. Spirometric reference values from a sample of the general US population. *Am J Respir Crit Care Med*. 1999;159(1):179-187.
11. Armitage P, Berry G, Matthews JNS. *Statistical Methods in Medical Research*. 4th ed. Oxford, England: Blackwell Science; 2002.
12. 2005 update: workshop report, global strategy for asthma management and prevention. Global Initiative for Asthma Web site. <http://www.ginasthma.org>. Published 2005. Accessed July 2, 2008.
13. Yeatts K, Davis KJ, Sotir M, Herget C, Shy C. Who gets diagnosed with asthma? Frequent wheeze among adolescents with and without a diagnosis of asthma. *Pediatrics*. 2003;111(5, pt 1):1046-1054.
14. Jenkins MA, Clarke JR, Carlin JB, et al. Validation of questionnaire and bronchial hyperresponsiveness against respiratory physician assessment in the diagnosis of asthma. *Int J Epidemiol*. 1996;25(3):609-616.
15. Sisteck D, Wickens K, Armstrong R, D'Souza W, Town I, Crane J. Predictive value of respiratory symptoms and bronchial hyperresponsiveness to diagnose asthma in New Zealand. *Respir Med*. 2006;100(12):2107-2111.
16. Stout JW, Visness CM, Enright P, et al. Classification of asthma severity in children: the contribution of pulmonary function testing. *Arch Pediatr Adolesc Med*. 2006;160(8):844-850.
17. Pellegrino R, Viegi G, Brusasco V, et al. ATS/ERS task force: standardization of lung function testing. Interpretative strategies for lung function tests. *Eur Respir J*. 2005;26(5):948-968.
18. Hansen JE, Sun XG, Wasserman K. Spirometric criteria for airway obstruction: use percentage of FEV₁/FVC ratio below the fifth percentile, not < 70%. *Chest*. 2007;131(2):349-355.
19. Roberts SD, Farber MO, Knox KS, et al. FEV₁/FVC ratio of 70% misclassifies patients with obstruction at the extremes of age. *Chest*. 2006;130(1):200-206.