

The Evolving Role of Pulmonary Function Interpretation: Clinical Implications of the New ERS/ATS Standards in Asthma Care

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■ Abstract

Asthma remains a significant public health challenge, requiring precise diagnostic and management strategies. Pulmonary function tests (PFTs) are essential in assessing disease severity, guiding treatment decisions, and monitoring disease progression. The 2022 ERS/ATS technical standards introduced critical updates to enhance the accuracy and standardization of interpretation of pulmonary function findings. These modifications include the adoption of Global Lung Initiative reference values, the transition from race-based to race-neutral equations, the replacement of percent-predicted values with z-scores, and a redefinition of bronchodilator responsiveness criteria. Additionally, new spirometric patterns such as dysanapsis and preserved ratio impaired spirometry have been recognized, improving the detection and characterization of airflow limitation. These updates significantly impact asthma management by refining disease phenotyping, improving diagnostic precision, and tailoring treatment strategies. Furthermore, advancements in artificial intelligence are expected to enhance predictive analytics and early intervention strategies in assessment of pulmonary function. However, challenges remain with respect to the adoption of these modifications in clinical practice, particularly regarding the classification of disease severity and the impact of race-neutral equations on diagnostic thresholds. Future research is necessary to validate the long-term implications of these changes on asthma outcomes. Clinicians must familiarize themselves with evolving standards to optimize patient care and reduce health disparities. The 2022 ERS/ATS guidelines represent a substantial advancement in PFT, with the potential to improve both clinical decision-making and patient prognosis in asthma management.

Key words: Asthma. Lung function. Spirometry. Reference equations. Guidelines.

■ Resumen

Las pruebas de función pulmonar son fundamentales para evaluar la gravedad del asma, guiar las decisiones terapéuticas y monitorizar su progresión. Las recomendaciones técnicas de la ERS/ATS de 2022 introdujeron actualizaciones clave para mejorar la precisión y estandarización en la interpretación de la función pulmonar. Estas modificaciones incluyen la adopción de los valores de referencia de la *Global Lung Initiative* (GLI), la transición hacia ecuaciones neutras, la sustitución de los valores porcentuales del predicho por los *z-scores* y la redefinición de los criterios de respuesta broncodilatadora. Además, se han reconocido nuevos patrones espirométricos, como la alteración disanápica y la espirometría con cociente preservado y alteración de la función pulmonar (PRISm), lo que mejora la detección y caracterización de la limitación del flujo aéreo. Estas actualizaciones refinan la caracterización fenotípica del asma, mejoran la precisión diagnóstica y facilitan estrategias terapéuticas más personalizadas. No obstante, persisten desafíos en la adopción clínica de estas modificaciones, en particular en la clasificación de la gravedad de la enfermedad y en el impacto de las ecuaciones neutras sobre los umbrales diagnósticos. Aunque se necesita conocer las implicaciones a largo plazo de estos cambios en los pacientes con asma, resulta conveniente la familiarización con estos estándares para optimizar la atención del paciente y reducir las disparidades en salud. Las guías ERS/ATS de 2022 representan un avance significativo en la evaluación de la función pulmonar, con potencialidad para mejorar la toma de decisiones clínicas y el pronóstico de los pacientes con asma.

Palabras clave: Asma. Función pulmonar. Espirometría. Ecuaciones de referencia. Guías clínicas.

Introduction

Asthma remains a major public health concern, affecting millions of individuals worldwide and contributing significantly to morbidity, health care utilization, and economic burden [1,2]. Given its chronic nature and the inherent variability in disease expression, effective monitoring tools are essential for optimizing disease control and guiding therapeutic interventions. Among these, pulmonary function testing (PFT) is a cornerstone in the evaluation and management of asthma, providing critical insights into pathophysiology [3]. Spirometry remains the primary tool for detecting bronchial obstruction, assessing both baseline lung function and responsiveness to bronchodilators, while additional assessments, including lung volume measurements and diffusing capacity of the lungs for carbon monoxide (DLCO), offer further insights into disease profile and severity [4].

Recognizing the need for standardization and accuracy in the interpretation of PFT results, leading scientific societies such as the European Respiratory Society (ERS) and the American Thoracic Society (ATS) have continuously refined guidelines to improve the reliability and clinical applicability of PFTs [5,6]. These efforts culminated in the 2022 ERS/ATS technical standards [7], which introduced critical updates aimed at addressing persistent challenges in the interpretation of lung function parameters, including the refinement of reference equations and diagnostic thresholds (Table 1). Given the pivotal role of lung function assessment in asthma care, these modifications may have significant implications for diagnosis, phenotyping, and treatment strategies.

This review aims to critically examine the recent updates in the ERS/ATS standards for interpreting PFT results and

to assess their impact on the clinical approach to asthma management. By analyzing the rationale behind the changes and their potential influence on diagnostic and therapeutic decision-making, we seek to provide clinicians with a comprehensive perspective on how these evolving guidelines shape the landscape of asthma care.

Clinical Implications of Key Updates in ERS/ATS Standards

Adoption of Global Lung Initiative Reference Values

A major update in the new ERS/ATS standards is the recommendation to use Global Lung Initiative (GLI) reference values for spirometry and other pulmonary function parameters. The GLI project provides age-, sex-, and height-specific reference equations derived from a large, multiethnic population. Currently, reference equations are available for spirometry [8], lung volumes [9], DLCO [10], and multiple breath washout [11], with additional equations in various stages of development.

The universal adoption of GLI reference equations offers several advantages, including improved standardization of PFT reporting and interpretation, consistency across different PFTs—preventing discordant results [12]—and applicability across all age groups, thereby eliminating the need for transitional equations. While some controversy remains regarding the impact of transitioning from traditional pediatric reference equations (eg, Zapletal), available evidence confirms that this change has a minimal effect [13] or, at most, results in a slight overestimation of pulmonary function impairment [14]. Additionally, the availability of algorithms and open-source

Table 1. Summary of the Main Modifications Introduced in the 2022 ERS/ATS Recommendations for Interpretation of Pulmonary Function Results.	
Parameter/Test	Recommendations
Reference equations	To use GLI reference equations Clarify that biological sex, not gender be used To assess race-neutral equations
Defining normal range	General use of LLN (5 th percentile) and ULN (95 th percentile) Use of fixed ratio FEV ₁ /FVC <0.7 or 80% predicted not recommended
Classification of physiological impairments	Spirometry: airflow obstruction; use lung volumes to detect hyperinflation/air trapping; dyanapsis; nonspecific pattern; PRISm Lung volumes: restrictive disorder (simple vs complex); hyperinflation/air trapping; mixed disorder Gas transfer: use of VA and KCO to classify low DLCO
Severity of lung function impairment	For all measures use z-score: Mild: −1.65 to −2.5 Moderate: −2.51 to −4.0 Severe: <−4.1
Bronchodilator response	>10% of predicted value in FEV ₁ or FVC
Interpretation of change over time	FEV ₁ Q in adults Conditional change score in children

Abbreviations: ATS, American Thoracic Society; DLCO, diffusing capacity of the lungs for carbon monoxide FEV₁, forced expiratory volume at 1 second; ERS, European Respiratory Society; FEV₁Q, FEV₁ quotient; FVC, forced vital capacity; GLI, Global Lung Initiative; KCO, DLCO/VA ratio; LLN, lower limit of normal; PRISm, preserved ratio impaired spirometry; ULN, upper limit of normal; VA, alveolar volume.

software for applying GLI equations in calculating functional parameters provides further advantages (<https://www.lungfunction.org/>; <https://gli-calculator.ersnet.org>).

The main implications of adopting GLI references in asthma patients can be summarized as follows: 1) The GLI reference values enhance accuracy in identifying airflow limitation across different demographic groups; 2) By incorporating data from diverse populations, GLI reference equations reduce bias in PFT, particularly for underrepresented ethnic groups; 3) Clinicians must transition from previously used reference values to the GLI system, ensuring appropriate classification of lung function abnormalities; 4) The use of GLI reference values may require updated training for health professionals to facilitate accurate interpretation and application in diverse clinical settings; and 5) Comparative studies between older reference equations and the GLI system suggest improved detection of subclinical airflow limitation in pediatric and elderly populations, necessitating adjustments in early asthma diagnosis and intervention strategies.

The year following the publication of the 2022 ERS/ATS recommendations [7] saw an additional, significant development, with the introduction of race-neutral reference equations [15]. The GLI-2023 global equations were derived from the same dataset as conventional GLI equations but incorporated inverse probability weighting to ensure equal contribution from all racial and ethnic groups. According to the ERS, the new GLI-2023 global equations, designed to encompass the full spectrum of lung function across all populations, should be applied with careful consideration of symptoms and medical history, particularly in clinical, occupational, and insurance contexts [16]. Accordingly, various academic societies have endorsed these new equations for interpretation of PFT results. In April 2023, the ATS issued an official statement [17] recommending the use of race-neutral equations in interpretation of PFT results to improve diagnostic accuracy and mitigate potential harms, such as delayed diagnoses and inappropriate clinical decisions. The statement also underscores the need for further research and education to understand the impact of this shift, emphasizing that race should not be used to infer biological characteristics [17].

However, transitioning to race-neutral reference equations requires careful consideration, particularly in children, to avoid unintended consequences. An analysis of 8719 North American children aged 5-12 years showed that race-neutral equations overestimate FEV₁ and FEV₁/FVC reference values in Black and Hispanic children, leading to a nearly 14% increase in asthma diagnoses. This shift was also associated with higher emergency visit rates and hospitalizations [18]. Another study of 24 630 children and adolescents confirmed that race-neutral equations generate lower predicted percentages and z-scores in Black children, while causing minimal differences in White children [19]. Consequently, adjusted models indicate that Black children are nearly 3 times more likely to present abnormal spirometry findings when transitioning from race-based to race-neutral equations [19].

Current evidence suggests similar findings in children and adolescents with asthma, where race-neutral equations yield lower predicted percentages and z-scores for FEV₁ in Black

children, resulting in a higher percentage of abnormal spirometry findings in both controlled and uncontrolled asthma cases [20].

The clinical implications of employing race-neutral reference equations in asthma care include the following: 1) Eliminating race-based adjustments in interpretation of PFT results promotes equity in respiratory medicine; 2) Transitioning to race-neutral equations ensures that lung function impairment is not underestimated in certain racial groups, leading to earlier and more accurate diagnosis; 3) Clinicians must be aware of this change and educate patients and colleagues on its significance in reducing health care disparities; 4) Adopting race-neutral equations may improve health outcomes in minority populations previously at risk of underdiagnosis or undertreatment; 5) The removal of race-specific reference values aligns with broader global initiatives aimed at eliminating implicit biases in medical practice, thus reinforcing ethical principles of equitable health care delivery; and 6) Studies examining the clinical impact of this shift suggest that race-neutral equations may lead to increased diagnosis of restrictive lung disease in previously overlooked populations, prompting more aggressive early interventions.

Replacement of Percent Predicted With Lower Limit of Normal or z-score

The 2022 ERS/ATS recommendations for interpreting PFT results propose replacing the traditional percent predicted value (%Pred) with the lower limit of normal (LLN) and the z-score [7]. While this shift has been discussed for some time, the current guidelines reinforce its adoption to enhance diagnostic precision and reduce misclassification of ventilatory disorders, allowing for a more individualized assessment that aligns with the statistical distribution of spirometry findings in the general population.

Historically, %Pred has been the most widely used criterion for assessment of pulmonary function. However, it has significant limitations [21]. Notably, it does not account for the natural variability in values within a healthy population, potentially leading to misclassification of individuals with expected variability as abnormal or failing to detect impairments in individuals whose values fall within the reference range but at the lower end of the distribution. Moreover, %Pred lacks uniform applicability across age groups, as lung function values change with age; thus, %Pred tends to overestimate impairment in older adults and underestimate it in children and adolescents. Additionally, %Pred relies on arbitrary cut-off points (typically 80% of the predicted value), which do not reflect the true population distribution and may result in inaccurate classifications.

In contrast, the 2022 ERS/ATS recommendations emphasize the use of LLN and z-score, both of which statistical approaches allow for a more precise interpretation of pulmonary function [7]. The LLN is defined as the value below which approximately 5% of the healthy population falls, after adjustment for age, sex, height, and ethnicity. This approach more accurately differentiates pathological values from those that simply reflect individual variability within normal limits, preventing misclassification of individuals with low but physiologically normal lung function.

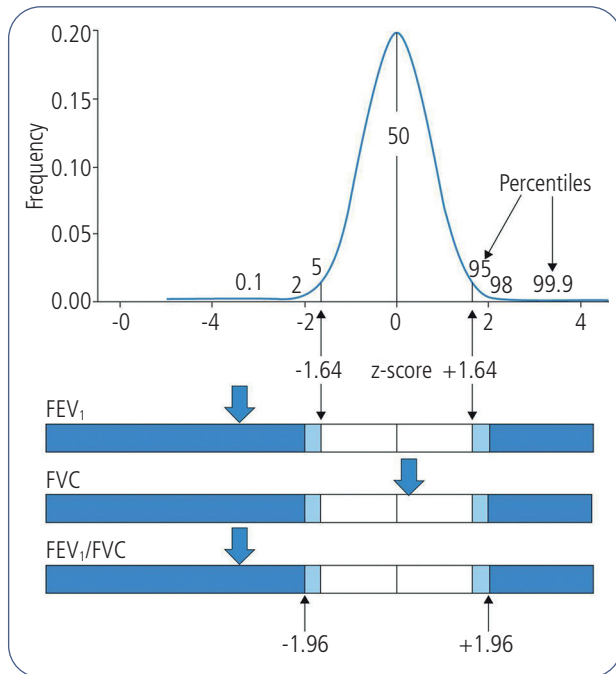


Figure 1. Schematic representation of the normal distribution curve of a respiratory function parameter and the correspondence between percentiles and z-score values. In the lower panel, the blue arrows represent the z-score values of a patient with obstructive impairment. FEV₁ indicates forced expiratory volume at 1 second; FVC, forced vital capacity.

The z-score expresses the deviation of a measured value from the population mean in terms of standard deviations. A z-score ≤ -1.64 indicates that the functional parameter falls at or below the 5th percentile (Figure 1), suggesting clinically relevant impairment. Unlike %Pred, the z-score enables standardized comparisons across individuals of different ages and physiological backgrounds [21], improving diagnostic sensitivity and specificity while reducing both overdiagnosis and underdiagnosis [22].

Implementing LLN and z-score instead of %Pred has significant clinical implications, particularly for vulnerable populations such as children, older adults, and individuals with anthropometric characteristics outside the standard reference range [7,23]. More accurate classification of obstructive or restrictive ventilatory patterns facilitates better therapeutic decision-making, preventing unnecessary medication use in individuals with low but normal pulmonary function and enabling earlier intervention in patients with incipient decline who still fall within the traditional %Pred reference range.

However, despite the advantages of LLN and z-score, it is important to acknowledge that much of the available evidence on asthma prognosis and treatment is based on patient stratification using %Pred. This includes defining study populations in clinical trials as well as establishing diagnostic and prognostic scales [24,25]. Therefore, further scientific research is needed to validate these new functional assessment parameters in clinical practice.

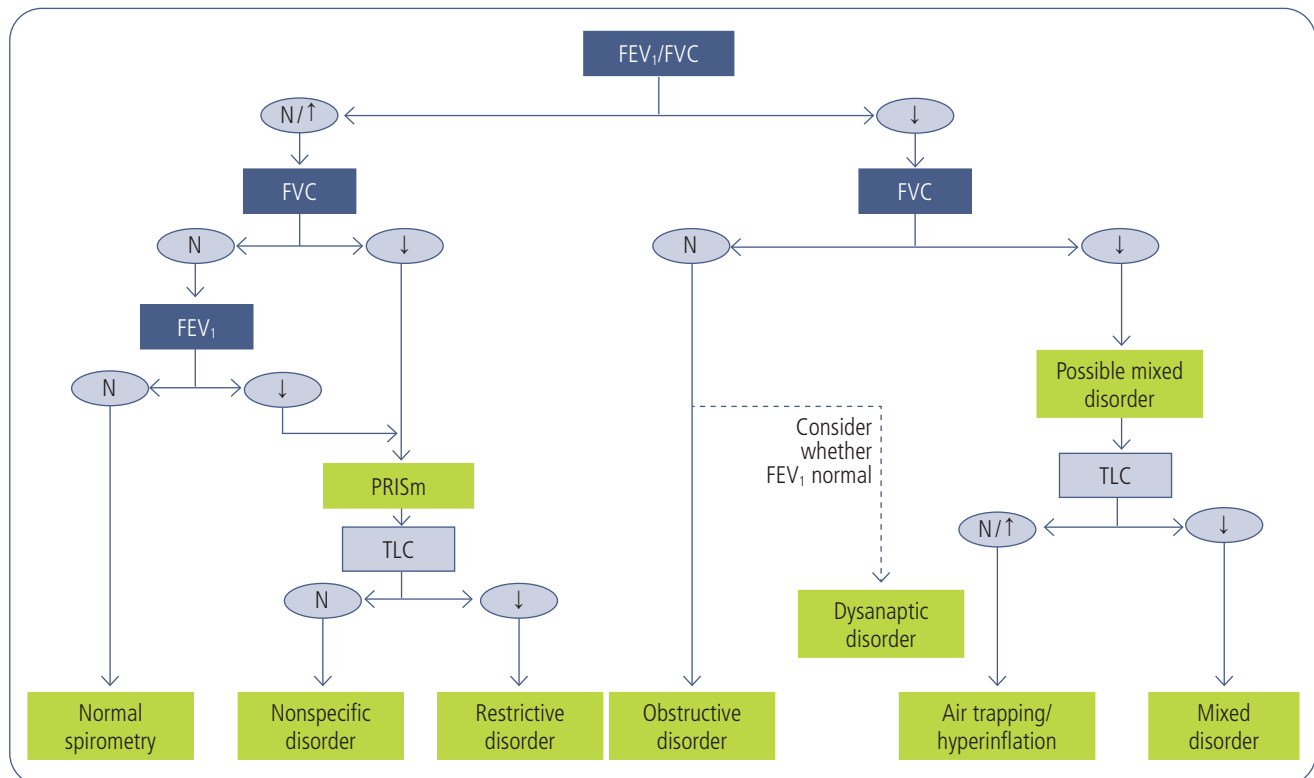


Figure 2. Algorithm for the interpretation of spirometry results proposed by the 2022 ERS/ATS recommendations for interpretation of pulmonary function results. The algorithm includes the assessment of static lung volumes for the definitive characterization of specific patterns. FEV₁ indicates forced expiratory volume at 1 second; FVC, forced vital capacity; N, normal value (between both limits of normality); PRISm, preserved ratio impaired spirometry; TLC, total lung capacity; ↓, reduced value (<lower limit of normality); ↑, increased value (>upper limit of normality).

Recognition of New Spirometric Patterns

An important novelty in the 2022 ERS/ATS guidelines on interpretation of PFT findings [7] is the identification of new spirometric impairment patterns, including the dysanaptic and nonspecific patterns (Figure 2).

Obstructive impairment is once again defined based on the LLN of the ratio of forced expiratory volume at 1 second to forced vital capacity (FEV_1/FVC) rather than using a fixed threshold of 0.70. While this criterion for obstructive ventilatory impairment aligns with the 1991 ATS [26] and 2005 ATS/ERS [5] guidelines, it differs from definitions provided by major obstructive disease management guidelines. Notably, the Global Initiative for Chronic Obstructive Lung Disease (GOLD) [27] and American and European guidelines on chronic obstructive pulmonary disease (COPD) [28] continue to use a fixed FEV_1/FVC threshold of 0.70 to identify obstruction, whereas asthma guidelines such as GEMA [29] and GINA [1] acknowledge the possibility of defining airflow limitation based on the LLN. Increasing evidence supports the definitive dismissal of the fixed 0.70 threshold for diagnosis of airflow limitation [30]. This is primarily because the FEV_1/FVC ratio declines with age, leading to underestimation of the prevalence of obstructive impairment in younger individuals and an overestimation in the elderly [30]. In fact, the FEV_1/FVC ratio decreases with age and height, even in nonsmokers, where the LLN drops below the fixed 0.70 threshold beyond age 45 [16]. Consequently, using the 0.70 threshold results in misclassification rates of up to 50% in older adults [16]. Furthermore, the fixed ratio fails to accurately distinguish mild obstruction and introduces significant age- and sex-related biases [30].

Dysanapsis is a newly recognized spirometric pattern. Under maximal effort, a low FEV_1/FVC ratio with a normal FEV_1 in an otherwise healthy individual may result from disproportionate growth between the airways and lung parenchyma [7]. Dysanapsis involves an unequal growth pattern, where lung parenchyma and airway length expand more than airway caliber [7]. While this profile may represent a normal variant in healthy individuals, it can also indicate a predisposition to obstructive disease [31-34]. This functional profile should be considered a potentially normal variant, particularly in healthy, asymptomatic, tall young males, especially if FVC is increased and distal flows remain normal [31]. In children, dysanaptic growth is associated with obesity or rapid weight gain in early childhood and predicts expiratory flow limitation, serving as an indicator of susceptibility to obstruction [32-34]. Determining whether dysanaptic growth signifies obstruction or is a normal variant requires assessment of the clinical context and additional evaluations, such as bronchodilator response testing, DLCO measurements, respiratory muscle strength evaluation, and cardiopulmonary exercise testing [7]. Furthermore, proper execution of forced expiratory maneuvers must be verified, as submaximal effort can overestimate FEV_1 and, consequently, lead to misinterpretation [31].

Another recognized pattern is preserved ratio impaired spirometry (PRISm). In the absence of plethysmography to confirm reduction in total lung capacity (TLC), a decreased

FVC or FEV_1 with a normal FEV_1/FVC ratio corresponds to PRISm [35,36]. This impairment can be observed in restrictive lung disease and small airway disease or result from suboptimal effort, where incomplete inspiration or expiration leads to overestimated FEV_1 and FVC values [7]. In such cases, the flow-volume curve may display a downward concavity at the end of expiration [7]. Bronchodilator testing may prove useful under optimal effort and when TLC cannot be determined [7,37]. A significant response to bronchodilation could indicate a degree of bronchial reactivity [7,37]. Evaluating slow vital capacity (SVC) can also help. An FVC of at least 200 mL lower than SVC may suggest small airway collapse with air trapping during forced expiration [12,16,38].

PRISm is a relatively common entity, with a prevalence ranging from 1.4% to 10% in the general population [35,36,39]. In Norwegian males, identified risk factors for PRISm include obesity, smoking history, and respiratory symptoms such as cough, sputum production, wheezing, asthma, and bronchitis [39]. Furthermore, in this population, PRISm is associated with an increased risk of respiratory mortality (HR, 4.00 [95%CI, 1.22-13.16]) [39]. A study of individuals aged 35-65 years from primary care centers confirms that asthma history and smoking are risk factors for PRISm, which, in turn, represents an independent risk factor for airflow limitation over the following 5 years [40]. Thus, PRISm could reflect preobstructive impairment. It has also been linked to an increased risk of small airway dysfunction, as defined by oscillometry and imaging techniques [41].

When TLC measurement is available, spirometric impairment can be better characterized. The presence of normal TLC, a normal FEV_1/FVC ratio, and a reduced FVC or FEV_1 characterizes nonspecific ventilatory impairment [37,42]. The significance of this pattern remains unclear. It may be a precursor to either a restrictive or an obstructive process [42]. Long-term follow-up of patients with nonspecific impairment revealed that two-thirds remained stable, while one-third progressed to either restriction or obstruction [42]. In obstructive processes, small airway collapse may lead to reduced FVC reduction and increased residual volume (RV) before a decline in FEV_1/FVC is observed.

Finally, the presence of a low FEV_1/FVC ratio with reduced TLC allows for the diagnosis of mixed ventilatory impairment, which, while less frequent, may be present in some asthma patients. This pattern represents the combination of airflow limitation and parenchymal or extraparenchymal lung disease. Although less common than obstruction, it can be observed in asthma patients with comorbid congestive heart failure or obesity.

In summary, the incorporation of these new spirometric patterns enables a more precise assessment of functional impairments in asthma, leading to improved diagnostic accuracy and tailored therapeutic strategies. While PRISm is more commonly associated with restrictive diseases, in asthma patients it may indicate significant airflow obstruction with air trapping, warranting static lung volume measurements for comprehensive evaluation. In the context of asthma, identifying a nonspecific pattern may suggest obstruction with reduced concurrent vital capacity, potentially due to hyperinflation or air trapping. Lastly, evidence of mixed impairment in asthma

patients may reflect substantial obstruction accompanied by restrictive changes, possibly secondary to bronchial remodeling or comorbid conditions.

Interpretation of Lung Volumes and Diffusion Capacity

Recent international recommendations have been published regarding the measurement of static lung volumes [43], encompassing plethysmography, dilution techniques, and multiple breath inert gas washout procedures, while also introducing, for the first time, a classification of quality levels for these measurements.

The 2 most relevant contributions of lung volume assessment in asthma are the identification of restriction and evaluation of hyperinflation/air trapping. Restriction is defined as a reduced TLC (below the LLN) and is typically due to decreased muscle strength, increased elastic recoil pressure, or reduced chest wall compliance.

The new consensus on interpretation differentiates between simple and complex restriction. Simple restriction involves a proportional decrease in FVC and TLC and is characteristic of diffuse interstitial lung diseases. In contrast, complex restriction is characterized by a disproportionate decline in FVC relative to TLC, leading to an increased RV and RV/TLC ratio, suggesting air trapping without an associated decrease in the FEV₁/FVC ratio [44]. Air trapping may reflect hidden obstruction or mechanical inability to reduce thoracic cavity volume, as seen in neuromuscular diseases and obesity.

Another novelty in the updated interpretation strategy is the unification of the terms air trapping and hyperinflation into a single abnormality, characterized by an increased RV/TLC ratio or functional residual capacity (FRC)/TLC above the upper limit of normal [7]. Both air trapping and pulmonary hyperinflation can be present in asthma patients. Therefore, static lung volume measurements have significant clinical implications in asthma, aiding in differentiating between the reversible obstruction characteristic of asthma and permanent structural changes, as well as guiding the use of specific therapies to reduce hyperinflation and improve respiratory mechanics.

Regarding DLCO, 4 possible abnormalities are recognized, namely, low DLCO with low alveolar volume (VA) and reduced DLCO/VA (KCO), low DLCO and VA with normal KCO, low DLCO with normal VA, and elevated DLCO [7]. Elevated DLCO may result from erythrocytosis, alveolar hemorrhage, and increased pulmonary blood flow (as occasionally seen in asthma patients, as well as in obese individuals and persons with left-to-right shunts). The interpretation of the 3 scenarios with reduced DLCO is based on evaluating VA and, in cases with reduced VA, the KCO [7]. However, Presti and Johnson [45] identified important issues in the proposed algorithm for interpreting DLCO. Although the algorithm acknowledges that KCO increases with lower VA, it does not account for the predictable relationship between KCO, DLCO, and VA [45,46]. Additionally, the algorithm overlooks the fact that patients with interstitial lung disease may have low, normal, or high KCO, and that patients with reduced VA due to incomplete

lung expansion may exhibit normal DLCO after adjustment for VA [45].

Beyond these considerations, the updated guidelines emphasize the need to interpret DLCO in conjunction with other parameters, such as VA and the DLCO/VA ratio, to differentiate between various phenotypes and comorbid conditions. This approach has multiple clinical implications in asthma management, including the following: 1) Differentiation from other respiratory conditions, as a reduced DLCO suggests the presence of concomitant diseases such as asthma-COPD overlap and early interstitial lung damage, which influence therapeutic strategies; 2) Monitoring pulmonary vascular involvement, since in severe asthma or patients with secondary pulmonary hypertension, assessment of DLCO allows for early detection of vascular impairment, facilitating timely interventions; 3) Evaluation of treatment response, assuming that normalization of DLCO once inflammation is controlled may indicate functional improvement in patients with difficult-to-control asthma, guiding adjustments in anti-inflammatory or biologic therapy; and 4) Optimized phenotypic stratification of asthma, given that an elevated DLCO value may be associated with specific phenotypes, such as eosinophilic asthma.

Ultimately, the updates to the 2022 ERS/ATS guideline reinforce the role of DLCO as a key complementary tool in the comprehensive assessment of asthma patients, enabling more precise diagnosis and optimization of treatment based on underlying pathophysiology.

Classification of the Severity of Functional Impairment

Another significant innovation introduced in the 2022 ERS/ATS document on interpretation of PFT findings is the unified classification of severity for all assessments based on the z-score. According to this system, any impairment is considered mild when the z-score of the corresponding parameter is between -2.5 and -1.64 , moderate when the z-score ranges from -4.0 to -2.5 , and severe when the z-score is below -4.0 [7].

It is important to note that this classification is based solely on mortality risk as a reference point and may not necessarily reflect the severity of symptoms, the risk of exacerbations, or social consequences. Neder [12] asserts that such a classification should primarily reflect current functional impairment rather than future risk, as the latter is a complex construct that extends beyond pulmonary function in individual patients. Nevertheless, some uncertainties persist regarding the suitability of the ERS/ATS-recommended classification system for stratifying the severity of obstructive ventilatory impairment [47].

Assessing the severity of ventilatory impairment is often challenging and uncertain [48]. Traditionally, this evaluation has relied on arbitrary thresholds to categorize results into 3 to 5 levels, which correlate only loosely with disease symptoms and mortality rates [49]. It remains unclear whether the 3-tier severity scale derived from the ERS/ATS z-score classification will prove more effective than previous scales based on percent predicted values [48]. In fact, studies in COPD patients have

demonstrated that mortality is better predicted using %Pred-based classifications rather than z-score-based systems, particularly in individuals over 65 years of age [50].

Other classification models based on absolute FEV₁ values in relation to the square or cubic power of height [51] or on the FEV₁/FVC ratio [52,53] have also demonstrated high prognostic value in both healthy individuals and patients with obstructive ventilatory impairments.

Given these uncertainties and considering that the new classification has yet to be adopted by major national and international guidelines for the management of asthma and other obstructive diseases, its real impact on clinical practice and decision-making remains to be determined. This is particularly relevant considering that nearly all available evidence derives from severity stratification schemes based on %Pred FEV₁ values.

Bronchodilator Test

The 2022 ERS/ATS consensus document [7] introduces significant modifications in the interpretation of bronchodilator test results. The assessment remains based on changes in both FEV₁ and FVC. FEV₁ primarily reflects the degree of airflow limitation and is considered the most sensitive marker of reversible airway obstruction, as it quantifies the maximal expiratory flow in the first second of a forced maneuver. An increase in FEV₁ after administration of bronchodilator is indicative of bronchodilation and improved airway patency. At the same time, FVC provides additional information, particularly in patients with obstructive lung diseases characterized by air trapping. An increase in postbronchodilator FVC suggests a reduction in dynamic hyperinflation and improved lung emptying, which may not be captured solely by FEV₁. This response is particularly relevant in conditions where air trapping contributes significantly to respiratory symptoms. The differential behavior of FEV₁ and FVC highlights the importance of a comprehensive assessment of bronchodilator response, not only in terms of airflow

improvement, but also with respect to changes in lung volume and ventilatory mechanics.

Instead of considering the absolute change associated with the percentage change relative to the baseline value [5], the consensus now defines a positive test result as an increase in FEV₁ or FVC >10% compared to the predicted value [7]. This new interpretation criterion aims to mitigate the impact of baseline pulmonary function on the response expressed as a percentage of the baseline value or in absolute terms, recognizing that patients with high baseline FEV₁ values are penalized when assessing percentage changes, while those with very low baseline FEV₁ values face notable limitations in achieving an increase greater than 200 mL [7]. Evaluating the bronchodilator test based on the predicted value also minimizes the effects of sex and height. Additionally, evidence suggests improved survival rates in patients with obstructive ventilatory impairment and reversibility of more than 8% of the predicted FEV₁ [54].

Table 2 summarizes the different scenarios considered in the interpretation of the bronchodilator test, as well as the subsequent steps in clinical and functional assessment.

However, applying the new criteria may not always confirm a bronchodilator response compared to classical criteria [45,55], although the test result is not definitive for selecting asthma treatment [24]. Various studies in asthma patients indicate that the frequency of a positive bronchodilator test is slightly lower when applying the 2022 ERS/ATS criteria than the 1991 ERS/ATS recommendation [55-58]. Nevertheless, at the individual level, there is strong concordance between test results under both recommendations, with the Cohen κ indices ranging from 0.78 to 0.89, regardless of whether GLI reference equations or those of other reference groups are applied [57,58].

In any case, these potential discrepancies have little impact on the overall value of the bronchodilator test. In fact, although a bronchodilator response may indicate changes in a patient's clinical status, its utility in differentiating between various airway diseases remains imprecise [7,59]. Therefore, some experts have suggested using nonbinary reversibility criteria [60], particularly in the pediatric population [16].

Table 2. Interpretation of Bronchodilator Response and Subsequent Clinical Steps According to ERS/ATS 2022 Criteria.

Post-BD Spirometry Pattern	Interpretation	ERS/ATS 2022 Positive Response Criteria	Next Steps
Increase in FEV ₁ and FVC (FEV ₁ /FVC ≥LLN)	Reversible airflow limitation (asthma, ACO without fixed obstruction)	FEV ₁ and/or FVC increase ≥10% of predicted value	Assess FeNO, blood eosinophils, or allergy markers to confirm asthma or ACO
Increase in FEV ₁ and FVC (FEV ₁ /FVC <LLN)	Reversible airflow limitation with persistent obstruction (asthma, ACO with fixed component)	FEV ₁ and/or FVC increase ≥10% of predicted value	Evaluate for fixed obstruction using HRCT or DLCO testing
Stable FEV ₁ , increased FVC (FEV ₁ /FVC ≥LLN)	Bronchodilator-reversible air trapping (early COPD, small airway dysfunction, hyperinflation improvement)	FEV ₁ remains stable, FVC increases ≥10% of predicted value	Assess RV, IC/TLC ratio for hyperinflation; consider impulse oscillometry (IOS), DLCO, or HRCT
Stable FEV ₁ , increased FVC (FEV ₁ /FVC <LLN)	Bronchodilator-reversible air trapping with persistent airway obstruction	FEV ₁ remains stable, FVC increases ≥10% of predicted value	Evaluate for fixed obstruction in addition to hyperinflation studies

Abbreviations: ACO, asthma COPD overlap; ATS, American Thoracic Society; BD, bronchodilation; COPD, chronic obstructive pulmonary disease; DLCO, diffusing capacity of the lungs for carbon monoxide; ERS, European Respiratory Society; FeNO, fractional exhaled nitric oxide; FEV₁, forced expiratory volume at 1 second; FVC, forced vital capacity; HRCT, high-resolution computed tomography; IC, inspiratory capacity; LLN, lower limit of normal; RV, residual volume; TLC, total lung capacity.

In summary, the revised bronchodilator test interpretation system has several clinical implications, as follows: 1) It improves the standardization of assessment based on predicted values, minimizing variability due to patient-specific factors such as height, sex, and baseline pulmonary function; 2) A slightly more restrictive bronchodilation criterion enhances the identification of a true bronchodilator response; and 3) Shifting to a predicted value-based evaluation reduces the misclassification of asthma severity in patients with mild airflow limitation.

Longitudinal Assessment of Pulmonary Function

Longitudinal assessment of PFT results can reveal excessive decline in pulmonary function due to exposure to harmful agents, underlying disease, or disease progression [61]. Ideally, an individual's pulmonary function before disease onset would serve as the reference point [61]. However, since this information is often unavailable, comparisons are made with the physiological decline observed in the healthy population [7], after taking into account biological variability and measurement errors [7,16]. Given that variability between tests (eg, up to 150 mL for FEV₁) far exceeds even an accelerated annual decline rate, multiple measurements over an extended period are required to establish a valid decline rate for an individual [5,62].

Longitudinal assessment of pulmonary function requires parameters that enable precise analysis of changes over time. Traditionally, %Pred has been widely used. However, it has a key limitation, namely, it is calculated in relation to a predicted value that changes with age and other factors, thus complicating its interpretation in long-term follow-ups. To address this limitation, the z-score provides a more robust alternative, reflecting the standard deviation of the observed value relative to the expected value at any given time, allowing for more precise comparisons.

The 2022 ERS/ATS guidelines on interpreting PFT findings recommend using the FEV₁ quotient (FEV₁Q) in adults and change score in children [7]. In adults, FEV₁Q is an interesting method for evaluating decline in pulmonary function [63]. It expresses FEV₁ in relation to a lower limit that represents the “survival threshold” below which the risk of mortality increases significantly [63]. Consequently, FEV₁Q is calculated as the ratio of FEV₁ (in liters) divided by 0.5 in men and 0.4 in women (values correspond to the first percentile) [63]. Under normal conditions, FEV₁Q decreases by 1 unit every 18 years in healthy individuals and by 1 unit every 10 years in smokers and older adults [63]. Therefore, FEV₁Q should remain stable for up to a year, while a rapid decline indicates a significant change in pulmonary function [7]. However, specific thresholds for defining stability or rapid decline in FEV₁Q do not yet exist [64], and in practice, detecting excessive changes reliably can be challenging. Additionally, Neder [12] highlights the complexities of FEV₁Q, noting that the first percentile can vary significantly depending on age, body size, and underlying diseases.

Several considerations must be taken into account for pediatric populations [7]. A child or adolescent is not simply

a miniature version of an adult [65,66], and thus, longitudinal assessment of pulmonary function during a period of rapid growth and development cannot be extrapolated from adult studies [16]. Consequently, interpreting decline in children and adolescents must account for the complexity of pulmonary function during this stage of life [23]. In 2020, a change score was developed to evaluate the decline in pulmonary function in children and adolescents [23]. This index considers longitudinal changes in the z-score of FEV₁ using a specific formula. While this is a promising tool for assessing decline in pulmonary function in pediatric populations, further studies are needed to validate its relevance.

Impact on Asthma Care

The recent modifications for the interpretation of PFT results introduced by the ERS and the ATS have far-reaching implications for asthma care. These changes refine diagnostic precision, optimize disease monitoring, personalize treatment approaches, and contribute to health equity. Below, we explore the most relevant aspects of these updates and their impact on asthma management.

Improved Diagnostic Accuracy

Accurate lung function assessment is essential for diagnosing asthma, particularly in patients with borderline or mild disease. The adoption of GLI reference values enhances diagnostic precision by providing more representative and standardized baseline values across different populations. This update minimizes misclassification errors that could lead to unnecessary treatments or missed diagnoses. The improved accuracy is particularly relevant for distinguishing between asthma and other obstructive or restrictive pulmonary diseases, thereby optimizing patient management.

Enhanced Disease Monitoring

Asthma is a chronic condition that requires continuous monitoring to assess disease progression and treatment efficacy. The updated bronchodilator response criteria and standardized interpretation strategies ensure that changes in lung function are detected with greater reliability over time. These modifications allow clinicians to identify subtle declines in pulmonary function, prompting timely adjustments in therapy. Standardized interpretation of spirometry results also facilitates longitudinal comparisons, improving the ability to track disease course and response to treatment across various clinical settings.

Personalized Treatment Adjustments

The redefinition of the bronchodilator response has direct implications for therapeutic decision-making, particularly regarding the initiation or intensification of bronchodilator therapy. Patients who were previously in a diagnostic gray area can now be classified more precisely [57], potentially facilitating the implementation of personalized therapeutic interventions. This refinement may be particularly relevant when considering the optimal timing for the introduction

of inhaled corticosteroids or combination therapies, as it could help ensure that patients receive treatment that is both evidence-based and appropriately tailored to their needs. Additionally, the updated guidelines encourage a more nuanced approach to bronchodilator testing, potentially reducing the overuse of medications in cases where reversibility criteria are less clear.

Addressing Health Disparities

Historically, pulmonary function reference values have incorporated race-based adjustments, a practice increasingly recognized as problematic because of its potential to worsen health disparities. The transition to race-neutral GLI reference equations represents a significant step toward equitable asthma care. This change ensures that interpretation of spirometry findings is not influenced by race-based biases, which could previously have led to underdiagnosis or undertreatment in certain populations. Clinicians must be aware of these updates and advocate for their implementation to promote equitable care across diverse patient groups.

Implications for Pediatric Asthma Management

The use of GLI reference values holds particular significance in pediatric asthma care, where lung function trajectories change dynamically with growth and development. More accurate and age-appropriate reference equations improve early detection of abnormal pulmonary function patterns, enabling earlier and more effective interventions. This refinement supports clinicians in differentiating between transient wheezing, persistent asthma, and other obstructive conditions in children. Additionally, standardized lung function interpretation can guide treatment adjustments during key developmental stages, optimizing long-term respiratory health outcomes.

Integration with Emerging Technologies

The growing incorporation of artificial intelligence (AI) and machine learning in pulmonary medicine presents new opportunities for enhancing asthma management. The updated ERS/ATS standards provide a robust framework for integrating predictive analytics into clinical practice. By leveraging refined reference values and bronchodilator response criteria, AI-driven algorithms can enhance the detection of subtle changes in lung function, facilitate personalized treatment recommendations, and improve risk stratification. These advancements hold promise for early intervention strategies, potentially reducing exacerbations and improving long-term disease control.

Conclusion

The revised ERS/ATS standards for interpreting PFT results mark a significant advancement in asthma diagnosis and management. By improving diagnostic accuracy, enhancing disease monitoring, supporting personalized treatment approaches, addressing health disparities, refining pediatric asthma care, and integrating with emerging technologies,

the updates have the potential to transform asthma care. Widespread adoption and implementation of the ERS/ATS standards will be critical in ensuring optimal outcomes for patients across different clinical settings and demographic groups.

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Conflicts of Interest

The author declares that he has no conflicts of interest.

References

1. Global Initiative for Asthma. Global Strategy for Asthma Management and Prevention, 2024. Updated May 2024. Available from: www.ginasthma.org.
2. Guía Española para el Manejo del Asma (GEMA 5.2). Madrid: Luzán S; 2022.
3. Chung KF, Wenzel SE, Brozek JL, Bush A, Castro M, Sterk PJ, et al. International ERS/ATS guidelines on definition, evaluation and treatment of severe asthma. *Eur Respir J*. 2014;43(2):343-73.
4. Mosnaim G. Asthma in Adults. *N Engl J Med*. 2023;389(11):1023-31.
5. Pellegrino R, Viegi G, Brusasco V, Crapo RO, Burgos F, Casaburi R, et al. Interpretative strategies for lung function tests. *Eur Respir J*. 2005;26(5):948-68.
6. Miller MR, Crapo R, Hankinson J, Brusasco V, Burgos F, Casaburi R, et al. General considerations for lung function testing. *Eur Respir J*. 2005;26(1):153-61.
7. Stanojevic S, Kaminsky DA, Miller MR, Thompson B, Aliverti A, Barjaktarevic I, et al. ERS/ATS technical standard on interpretive strategies for routine lung function tests. *Eur Respir J*. 2022;60(1):2101499.
8. Quanjer PH, Stanojevic S, Cole TJ, Baur X, Hall GL, Culver BH, et al. Multi-ethnic reference values for spirometry for the 3-95-yr age range: the global lung function 2012 equations. *Eur Respir J*. 2012;40(6):1324-43.
9. Hall GL, Filipow N, Ruppel G, Okitika T, Thompson B, Kirkby J, et al. Official ERS technical standard: Global Lung Function Initiative reference values for static lung volumes in individuals of European ancestry. *Eur Respir J*. 2021;57(3):2000289.
10. Stanojevic S, Graham B, Cooper B, Thompson B, Carter K, Francis R, Hall G. Official ERS Technical Standards: Global Lung Function Initiative Reference Values for the Carbon Monoxide Transfer Factor for Caucasians. *Eur Respir J*. 2017;50(3):1700010.
11. Ramsey KA, Stanojevic S, Chavez L, Johnson N, Bowerman C, Hall GL, et al. Global Lung Function Initiative reference values for multiple breath washout indices. *Eur Respir J*. 2024;64(6):2400524.
12. Neder JA. The new ERS/ATS standards on lung function test interpretation: some extant limitations. *Eur Respir J*. 2022;60(2):2200252.
13. Quanjer PH, Weiner DJ. Interpretative consequences of adopting the Global Lungs 2012 reference equations for

- spirometry for children and adolescents. *Pediatr Pulmonol*. 2014;49(2):118-25.
14. Raaijmakers L, Zwitserloot A, Merkus P, Gappa M. Implications of the Transition From Zapletal to GLI Reference Values for Spirometry. *Pediatrics*. 2016;137(1). doi: 10.1542/peds.2015-0033.
 15. Bowerman C, Bhakta NR, Brazzale D, Cooper BR, Cooper J, Gochicoa-Rangel L, et al. A Race-neutral Approach to the Interpretation of Lung Function Measurements. *Am J Respir Crit Care Med*. 2023;207(6):768-74.
 16. Barkous B, Briki C, Boubakri S, Abdesslem M, Ben Abbes N, Ben Hmid W, et al. Routine pulmonary lung function tests: Interpretative strategies and challenges. *Chron Respir Dis*. 2024;21:14799731241307252.
 17. Bhakta NR, Bime C, Kaminsky DA, McCormack MC, Thakur N, Stanojevic S, et al. Race and Ethnicity in Pulmonary Function Test Interpretation: An Official American Thoracic Society Statement. *Am J Respir Crit Care Med*. 2023;207(8):978-95.
 18. Non AL, Li X, Jones MR, Oken E, Hartert T, Schoettler N, et al. Comparison of Race-neutral Versus Race-specific Spirometry Equations for Evaluation of Child Asthma. *Am J Respir Crit Care Med*. 2025;211(3):464-76.
 19. Forno E, Weiner DJ, Rosas-Salazar C. Spirometry Interpretation After Implementation of Race-Neutral Reference Equations in Children. *JAMA Pediatr*. 2024;178(7):699-706.
 20. Burbank AJ, Atkinson CE, Espallat AE, Schworer SA, Mills K, Rooney J, et al. Race-specific spirometry equations may overestimate asthma control in Black children and adolescents. *Respir Res*. 2023;24(1):203.
 21. Stanojevic S, Wade A, Stocks J. Reference values for lung function: past, present and future. *Eur Respir J*. 2010;36(1):12-19.
 22. Dinh-Xuan AT, Graham BL, Thompson B, Miller MR, Stanojevic S. Reconciling the past and considering the future of pulmonary function test interpretation. *Eur Respir J*. 2024;63(2):2302225.
 23. Stanojevic S, Filipow N, Ratjen F. Paediatric reproducibility limits for the forced expiratory volume in 1 s. *Thorax*. 2020;75(10):891-6.
 24. Pérez de Llano L, Muñoz Fernández MC, Dávila I. Impact of the ERS/ATS 2022 Guidelines for Interpretation of Lung Function Test Results When Assessing the Response to Biologics in Asthma. *J Investig Allergol Clin Immunol*. 2024;34(1):73-4.
 25. Pérez de Llano L, Cisneros C, Domínguez-Ortega J, Martínez-Moragón E, Olaguibel JM, Plaza V, et al. Response to Monoclonal Antibodies in Asthma: Definitions, Potential Reasons for Failure, and Therapeutic Options for Suboptimal Response. *J Investig Allergol Clin Immunol*. 2023;33(1):1-13.
 26. Lung function testing: selection of reference values and interpretative strategies. American Thoracic Society. *Am Rev Respir Dis*. 1991;144(5):1202-18.
 27. Agustí A, Celli BR, Criner GJ, Halpin D, Anzueto A, Barnes P, et al. Global Initiative for Chronic Obstructive Lung Disease 2023 Report: GOLD Executive Summary. *Arch Bronconeumol*. 2023;59(4):232-48.
 28. Qaseem A, Wilt TJ, Weinberger SE, Hanania NA, Criner G, van der Molen T, et al. Diagnosis and management of stable chronic obstructive pulmonary disease: a clinical practice guideline update from the American College of Physicians, American College of Chest Physicians, American Thoracic Society, and European Respiratory Society. *Ann Intern Med*. 2011;155(3):179-91.
 29. Plaza V, Blanco M, Ferreira J, García G, Morete A, Quirce S, et al. Highlights of the Spanish Asthma Guidelines (GEMA), Version 5.4. *Open Respir Arch*. 2024;6(4):100356.
 30. Choi JY, Rhee CK. It is high time to discard a cut-off of 0.70 in the diagnosis of COPD. *Expert Rev Respir Med*. 2024;18(9):709-19.
 31. Dos Santos Andreato L, Soares MR, Pereira CA. Reduced FEV(1)/FVC and FEV(1) in the Normal Range as a Physiological Variant. *Respir Care*. 2019;64(5):570-5.
 32. Peralta GP, Abellan A, Montazeri P, Basterrechea M, Esplugues A, González-Palacios S, et al. Early childhood growth is associated with lung function at 7 years: a prospective population-based study. *Eur Respir J*. 2020;56(6):2000157.
 33. Arismendi E, Bantulà M, Perpiñà M, Picado C. Effects of Obesity and Asthma on Lung Function and Airway Dysanapsis in Adults and Children. *J Clin Med*. 2020;9(11):3762.
 34. Forno E, Weiner DJ, Mullen J, Sawicki G, Kurland G, Han YY, et al. Obesity and Airway Dysanapsis in Children with and without Asthma. *Am J Respir Crit Care Med*. 2017;195(3):314-23.
 35. Higbee DH, Granell R, Davey Smith G, Dodd JW. Prevalence, risk factors, and clinical implications of preserved ratio impaired spirometry: a UK Biobank cohort analysis. *Lancet Respir Med*. 2022;10(2):149-57.
 36. Wan ES, Balte P, Schwartz JE, Bhatt SP, Cassano PA, Couper D, et al. Association Between Preserved Ratio Impaired Spirometry and Clinical Outcomes in US Adults. *JAMA*. 2021;326(22):2287-98.
 37. Hyatt RE, Cowl CT, Bjoraker JA, Scanlon PD. Conditions associated with an abnormal nonspecific pattern of pulmonary function tests. *Chest*. 2009;135(2):419-24.
 38. Saint-Pierre M, Ladha J, Berton DC, Reimao G, Castelli G, Marillier M, et al. Is the Slow Vital Capacity Clinically Useful to Uncover Airflow Limitation in Subjects With Preserved FEV(1)/FVC Ratio? *Chest*. 2019;156(3):497-506.
 39. Cestelli L, Johannessen A, Gulsvik A, Stavem K, Nielsen R. Risk Factors, Morbidity, and Mortality in Association With Preserved Ratio Impaired Spirometry and Restrictive Spirometric Pattern: Clinical Relevance of Preserved Ratio Impaired Spirometry and Restrictive Spirometric Pattern. *Chest*. 2025;167(2):548-60.
 40. Miura S, Iwamoto H, Omori K, Yamaguchi K, Sakamoto S, Horimasu Y, et al. Preserved ratio impaired spirometry with or without restrictive spirometric abnormality. *Sci Rep*. 2023;13(1):2988.
 41. Zhao N, Wu F, Peng J, Zheng Y, Tian H, Yang H, et al. Preserved ratio impaired spirometry is associated with small airway dysfunction and reduced total lung capacity. *Respir Res*. 2022;23(1):298.
 42. Iyer VN, Schroeder DR, Parker KO, Hyatt RE, Scanlon PD. The nonspecific pulmonary function test: longitudinal follow-up and outcomes. *Chest*. 2011;139(4):878-86.
 43. Bhakta NR, McGowan A, Ramsey KA, Borg B, Kivastik J, Knight SL, et al. European Respiratory Society/American Thoracic Society technical statement: standardisation of the measurement of lung volumes, 2023 update. *Eur Respir J*. 2023;62(4):2201519.
 44. Clay RD, Iyer VN, Reddy DR, Siontis B, Scanlon PD. The "Complex Restrictive" Pulmonary Function Pattern: Clinical

- and Radiologic Analysis of a Common but Previously Undescribed Restrictive Pattern. *Chest*. 2017;152(6):1258-65.
45. Presti TP, Johnson DC. Improving pulmonary function test interpretation. *Eur Respir J*. 2023;61(1):2201858.
 46. Johnson DC. Importance of adjusting carbon monoxide diffusing capacity (DLCO) and carbon monoxide transfer coefficient (KCO) for alveolar volume. *Respir Med*. 2000;94(1):28-37.
 47. Bhatt SP, Bodduluri S, Nakhmani A. ERS/ATS spirometry interpretation standards: a gap in grading severity of airflow obstruction. *Eur Respir J*. 2024;63(2):2301910.
 48. Calverley PMA. A Rising STAR in Chronic Obstructive Pulmonary Disease or More Deckchair Rearrangement? *Am J Respir Crit Care Med*. 2024;210(11):1285-7.
 49. Huang TH, Hsiue TR, Lin SH, Liao XM, Su PL, Chen CZ. Comparison of different staging methods for COPD in predicting outcomes. *Eur Respir J*. 2018;51(3):1700577.
 50. Tejero E, Prats E, Casitas R, Galera R, Pardo P, Gavilan A, et al. Classification of Airflow Limitation Based on z-Score Underestimates Mortality in Patients with Chronic Obstructive Pulmonary Disease. *Am J Respir Crit Care Med*. 2017;196(3):298-305.
 51. Pedone C, Scarlata S, Scichilone N, Forastiere F, Bellia V, Antonelli-Incalzi R. Alternative ways of expressing FEV1 and mortality in elderly people with and without COPD. *Eur Respir J*. 2013;41(4):800-5.
 52. Calverley PMA. A STAR Is Born: A New Approach to Assessing Chronic Obstructive Pulmonary Disease Severity. *Am J Respir Crit Care Med*. 2023;208(6):647-8.
 53. Bhatt SP, Nakhmani A, Fortis S, Strand MJ, Silverman EK, Wilson CG, et al. STAR Has Better Discrimination for Mortality than ERS/ATS Chronic Obstructive Pulmonary Disease Severity Classification. *Am J Respir Crit Care Med*. 2024;210(11):1376-9.
 54. Ward H, Cooper BG, Miller MR. Improved criterion for assessing lung function reversibility. *Chest*. 2015;148(4):877-86.
 55. Li Y, Lin J, Wang Z, Tan L, Liu S, Huang J, et al. Bronchodilator Responsiveness Defined by the 2005 and 2021 ERS/ATS Criteria in Patients with Asthma as Well as Chronic Obstructive Pulmonary Disease. *Int J Chron Obstruct Pulmon Dis*. 2022;17:2623-33.
 56. Betancor D, Villalobos-Vilda C, Olaguibel JM, Rodrigo-Muñoz JM, Puebla MJA, Arismendi E, et al. The New ERS/ATS 2022 Bronchodilator Response Recommendation: Comparison With the Previous Version in an Asthma Cohort. *Arch Bronconeumol*. 2023;59(9):608-11.
 57. Trepčić N, Nemet M, Vukoja M. Assessing the Impact of the Updated 2021 European Respiratory Society/American Thoracic Society Criteria on Bronchodilator Responsiveness in Asthma. *Cureus*. 2024;16(8):e66844.
 58. Chaiwong W, Deesomchok A, Pothirat C, Duangjit P, Liwsrisakun C. Impact of the new European respiratory (ERS)/American Thoracic Society (ATS) pulmonary function test interpretation guidelines 2021 on the interpretation of bronchodilator responsiveness in subjects with airway obstruction. *Respir Med*. 2023;220:107460.
 59. Halpin DMG. Bronchodilator Responsiveness in Asthma and Chronic Obstructive Pulmonary Disease: Time to Stop Chasing Shadows. *Am J Respir Crit Care Med*. 2024;209(4):349-51.
 60. Ioachimescu OC, Ramos JA, Hoffman M, McCarthy K, Stoller JK. Assessing bronchodilator response by changes in per cent predicted forced expiratory volume in one second. *J Investig Med*. 2021;69(5):1027-34.
 61. Redlich CA, Tarlo SM, Hankinson JL, Townsend MC, Eschenbacher WL, Von Essen SG, et al. Official American Thoracic Society technical standards: spirometry in the occupational setting. *Am J Respir Crit Care Med*. 2014;189(8):983-93.
 62. Miller MR, Hankinson J, Brusasco V, Burgos F, Casaburi R, Coates A, et al. Standardisation of spirometry. *Eur Respir J*. 2005;26(2):319-38.
 63. Miller MR, Pedersen OF. New concepts for expressing forced expiratory volume in 1 s arising from survival analysis. *Eur Respir J*. 2010;35(4):873-82.
 64. Rurak K, Schotland H. A query on FEV(1)Q: it may be useful, but is it helpful? *Eur Respir J*. 2023;61(1):2201646.
 65. Ben Saad H. In 2023, it is vital to standardize the interpretation of spirometry in children. *Pediatr Pulmonol*. 2023;58(8):2187-8.
 66. Guezguez F, Ben Saad H. What constitutes a "clinically significant" bronchodilator response in children? *Eur Respir J*. 2020;55(5):2000207.

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