Prediction of exercise-induced desaturation in COPD patients without resting hypoxemia: a retrospective study

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Abstract

Background: There is no universally accepted criterion for assessing exercise-induced desaturation (EID). The purpose of this study is to compare the two methods regularly used for determining EID in COPD patients, as well as to explore the risk factors and predictors related to EID.

Methods: The 6MWT was performed with continuous SpO2 monitoring on patients with stable COPD. Using two methods (method A: “SpO2rest–SpO2min ≥ 4% and/or SpO2min < 90%”, method B: “SpO2rest–SpO2end ≥ 4% and/or SpO2end < 90%”) as EID determination criteria to assess the incidence of EID. The differences and consistency of the two methods are compared. Moreover, we collected data through the pulmonary function test, mMRC dyspnea score, COPD assessment test, BODE index and CT-defined emphysema. Univariate and multivariate logistic regression analyses were used to identify factors affecting the EID. For the parameters that predict EID in 6MWT, a receiver operating characteristic (ROC) curve analysis was employed.

Results: The analysis included 124 patients. The overall incidence of EID was 62.1% by using method A as the criterion and 51.6% by method B. All of the EID patients found by method B were included in the EID patients identified by method A, as well as 13 new-EID patients. The difference in diagnostic outcomes between the two approaches was not statistically significant (P > 0.05), but they were in excellent agreement (Kappa = 0.807, P = 0.001). Logistic regression analyses found that DLCO SB% pred, DLCO/VA% pred, CAT score, mean density, PD15, emphysema volume and %LAA were significant determinants of the EID. For predicting EID, the ROC analysis produced AUC and cutoffs of 0.689 and 50.45% (DLCO SB% pred), 0.707 and 75.0% (DLCO/VA% pred), 0.727 and 15 points (CAT score), 0.691 and −955.00HU (PD15), 0.671 and −856.46HU (mean density), 0.668 and 338.14 ml (emphysema volume) and 0.656 and 7.63% (%LAA), respectively.

Conclusions: Two methods evaluating EID in this research are in a good agreement, method A can find more EID patients by focusing on SpO2min. When conditions are constrained, it is also sufficient to assess EID in COPD patients by method B. In terms of the predictors of EID, DLCO SB% pred, DLCO/VA% pred, CAT score and CT-defined emphysema are all statistically significant test variables to determine EID.
Introduction
Chronic obstructive pulmonary disease (COPD) is a chronic respiratory disease characterized by irreversible persistent airflow limitation, with high prevalence, mortality and disease burden [1]. Exercise-induced desaturation (EID) is a term used to describe patients with COPD who do not have hypoxemia at rest but have a desaturation during exercise. In contrast to the classical cardiopulmonary exercise test, the 6-min walk test (6MWT) does not require large professional medical equipment, is simple to perform, is acceptable to most patients in terms of exercise intensity, and is closer to patients' daily lives in terms of exercise form, and is now more widely used to monitor EID than the classical cardiopulmonary exercise test [2].

Different clinical investigations utilize different criteria to determine EID because there are no common standards or definitions. The following are commonly used: (1) the difference between the oxygen saturation (SpO₂) at the beginning of the test (SpO₂rest) and the SpO₂ at the end of the test (SpO₂end) (SpO₂rest–SpO₂end) ≥ 4% and/or the SpO₂end < 90% [3–7], (2) the difference between the SpO₂rest and the minimum SpO₂ during the test (SpO₂min) (SpO₂rest–SpO₂min) ≥ 4% and/or the SpO₂min < 90% [8], (3) SpO₂min ≤ 88% [9], (4) SpO₂end ≤ 88% [7, 10]. A study compared “SpO₂end ≥ 88%” and “SpO₂end ≤ 88%” and “SpO₂rest–SpO₂end ≥ 4% and/or the SpO₂end < 90%” in 507 patients with COPD, the incidence of EID was 5.1% (26/507) by the former method and 13% (66/507) by the latter. Although the former found a lower incidence of EID in COPD patients, it had a stronger prognostic value for long-term mortality than the latter after 162 months of follow-up [7]. It is certain that EID is prevalent in COPD patients, regardless of the determination method. However, few studies have focused on the SpO₂min during 6MWT [8, 9], potentially misclassifying some patients who desaturate during the 6MWT but do not desaturate at the conclusion as "non-EID patients". Therefore, continuous SpO₂ monitoring is required during the 6MWT to record the SpO₂min during the trial.

Emphysema plays a role in the onset and course of EID [10]. CT-defined emphysema is more sensitive and repeatable than subjective eye scoring systems, and it also allows for emphysema spatial distribution and localization. Most studies used — 950 Hounsfield units (HU) as the ideal threshold for diagnosis of CT defined emphysema [11–13]. The most widely used indicators for evaluating emphysema were lung volumes with X-ray attenuation values below — 950 HU, percentage low attenuation regions (% LAA), the 15th percentile point of density (PD 15), and the mean density of the lung [13–15]. Because it is less affected by picture noise, PD15 is more commonly utilized in longitudinal studies of emphysema [14, 15]. Studies shows that %LAA is independently associated with EID [16, 17]. According to the findings, each 1% increase in percent LAA raises the relative risk of EID in the 6MWT by 10% and the relative risk of EID recurrence by 20%, each 1% increase in %LAA increases the relative risk of EID in 6MWT by 10% and the relative risk of reoccurrence of EID by 20% [18]. Few studies, however, have looked at the relative predictive value and threshold of the mean density, PD15, emphysema volume, and %LAA for EID at the same time.

In patients with COPD, EID is linked to lower quality of life, decreased exercise tolerance, higher readmission rates for acute exacerbations, and increased morbidity and mortality [3, 19–21]. On the one hand, identifying whether markers indicate EID occurrences may assist physicians in assisting this group of patients in conducting the 6MWT with oxygen, which may help them avoid terminating the trial when the patients' blood oxygen levels are too low (less than 85% or 80%) [22, 23]. On the other hand, it can reflect the patient's most realistic exercise capacity while also reducing the danger of hypoxia-related exercise.

Therefore, the primary goal of this research was to answer the following three questions: (1) Obtain SpO₂min by continuously measuring SpO₂ during 6MWT. Using two methods (method A: “SpO₂rest–SpO₂min ≥ 4% and/or SpO₂min < 90%”, method B: “SpO₂rest–SpO₂end ≥ 4% and/or SpO₂end < 90%”) as EID determination criteria to assess the incidence of EID in patients with COPD during 6MWT. Comparing the differences and consistency between the two methods. (2) What are the clinical features of patients with EID who were diagnosed using the former technique but not the latter? (3) To investigate the predictive value of different indicators for EID by comparing questionnaire scores, pulmonary function and degree of emphysema in patients with or without EID. To find more sensitive predictors of EID and timely detection of EID events in COPD.

Methods
Patients
A total of 124 consecutive patients with COPD who met the inclusion criterion and did not meet the exclusion...
criteria were enrolled in the study between January 2019 and December 2021 in the Department of Respiratory and Critical Care Medicine of the China-Japan Friendship Hospital. The inclusion criterion was a diagnosis of stable COPD according to the Global Initiative for Chronic Obstructive Lung Disease (GOLD), as updated in 2020. The exclusion criteria were as follows: (1) SpO$_{2\text{rest}}$ < 94%; (2) unstable angina pectoris or myocardial infarction; (3) recently diagnosed pulmonary embolism and severe pulmonary hypertension in 1 month; (4) resting heart rate > 120 beats/min; (5) systolic blood pressure > 180 mmHg (1 mmHg = 0.133 kPa) and/or diastolic blood pressure > 100 mmHg; (6) malignant arrhythmia; (6) severe valvular disease; (7) limitations on walking including predominant neurological or musculoskeletal limitation; (8) Combined with a malignant tumor [22].

6MWT
The 6MWT was performed on a 30 m, flat, straight indoor walking course, supervised by at least two experienced investigators following the ATS guidelines [22]. Continuous measurements of SpO$_2$ were performed using a finger pulse oximeter that transmits data in real time via Bluetooth mentioned from 1 min before the beginning of the 6MWT to the fourth minute after the 6MWT. The 6MWD, SpO$_{2\text{rest}}$, SpO$_{2\text{end}}$, SpO$_{2\text{min}}$ was measured, then calculate SpO$_{2\text{rest}}$–SpO$_{2\text{end}}$ and SpO$_{2\text{rest}}$–SpO$_{2\text{min}}$. Use two methods: A: “SpO$_{2\text{rest}}$–SpO$_{2\text{min}}$ ≥ 4% and/or SpO$_{2\text{min}}$ < 90%” and B: “SpO$_{2\text{rest}}$–SpO$_{2\text{end}}$ ≥ 4% and/or SpO$_{2\text{end}}$ < 90%” as EID determination criteria to assess the incidence of EID in 6MWT. Patients determined to be EID by method A were defined as “EID patients”, patients determined to be EID by method A but not by method B were defined as “new-EID patients”, patients determined to be EID by both methods were defined as “original-EID patients”, and patients determined to be non-EID by both methods were defined as “non-EID patients”.

Pulmonary function
Pulmonary function was performed by well-trained hospital staff according to the American Thoracic Society (ATS) and the European Respiratory Society (ERS) guidelines using a Jaeger® MasterScreen system (Jaeger®, Viasys Healthcare GmbH, Hochberg, Germany) [24].

Questionnaires
Dyspnea was measured using the modified Medical Research Council (mMRC) dyspnea score [25], health-related quality of life and symptom burden of COPD was measured using the COPD Assessment Test (CAT) [26]. The BODE index was calculated according to BMI (B), airflow obstruction (O), dyspnea (D) and exercise ability (E) [27].

CT defined emphysema
All patients underwent CT scans using GE Healthcare multidetector-row CT scanner. The scans were done at suspended full inspiration. Exposure settings were 120 kVp and 40 mAs, images were reconstructed at 1.0 or 5.0 mm contiguous slices. The CT scans were analyzed using FACT Medical Imaging System 1.2.0 software. Briefly, the lungs were segmented from the thorax wall, heart, and main pulmonary vessels, followed by segmentation of the individual lobes. A density of less than − 950HU is used as the threshold for emphysema reconstruction [11–13], the lung volumes with X-ray attenuation values below − 950 HU, %LAA, the PD 15 and the mean density of the lung were calculated to assess the degree of emphysema [13–15] (Fig. 1).

Statistical analysis
The statistical analyses were performed with SPSS statistics software (version 23.0; SPSS Inc., Chicago, IL). Categorical variables were expressed as compositional ratios,
and the chi-square test was used. We used the Shapiro–Wilk test for the analysis of the normality of the data; results were described as mean and standard deviation or median and interquartile range (IQR) according to the data distribution. To determine whether differences were present between groups, selected characteristics were assessed using a chi-square test or independent t-test. Kappa verified the agreement of the two determination methods. Univariate and multivariate logistic regression analyses were used to identify factors affecting the EID. Then a receiver operating characteristic (ROC) curve analysis and the area under the curve (AUC) were used to determine the optimal cutoff value for the factors predicting EID. P values < 0.05 were considered statistically significant.

Results

Clinical characteristics

This study included 101 men and 23 women with stable COPD; among them, 37 patients underwent CT in 5 mm and 87 patients in 1 mm. The severity of airway limitation was classified as GOLD stage: mild (n = 24), moderate (n = 43), severe (n = 33) and very severe (n = 24). The mean \(\text{SpO}_{2_{\text{rest}}}-\text{SpO}_{2_{\text{min}}}\) in this study was 7.05 ± 6.79%, and the mean \(\text{SpO}_{2_{\text{rest}}}-\text{SpO}_{2_{\text{end}}}\) was 5.28% ± 6.66%. By using method A as the criterion, the overall incidence of EID was 62.1% (EID patients, n = 77) and 51.6% (original-EID patients, n = 64) by method B. All of the EID patients found by method B were included in the EID patients identified by method A, as well as 13 new EID patients. There was no statistically significant difference between the diagnostic outcomes of the two procedures (P > 0.05, Table 1). The diagnostic results of the two approaches were in good agreement in terms of consistency (Kappa = 0.807, P < 0.001). Table 2 lists the features of these individuals in various groups. In terms of age, height, body mass, BMI, FVC, FEV1, FVC% pred, FEV1%pred, and \(\text{SpO}_{2_{\text{rest}}'}\) no statistically significant changes were identified between EID and non-EID patients (P > 0.05). However, there were statistically significant differences in gender, 6MWD, \(\text{SpO}_{2_{\text{min}}'}\), \(\text{SpO}_{2_{\text{end}}'}\) diffusion function, questionnaire scores and degree of emphysema (P < 0.05, Table 2). Only the degree of \(\text{SpO}_{2}\) change in 6MWT differed statistically significant between the new-EID patients (n = 13) who were not detected by method B but detected by method A and original-EID patients (n = 64) who were detected by both methods (Table 2).

Correlation of EID with other outcome measures

With single regression analysis, the EID was significantly associated with 6MWD, \(\text{D}_{15}\text{CO SB}\) pred, \(\text{D}_{15}\text{CO/VA}\) pred, mMRC score, CAT score, CT-defined emphysema measurements, such as the mean density, PD15, emphysema volume, and LAA (Table 3). Only \(\text{D}_{15}\text{CO SB}\) pred, \(\text{D}_{15}\text{CO/VA}\) pred, CAT score, mean density, PD15, emphysema volume, and LAA were significant drivers of the EID in a multiple regression analysis (Table 3).

Different factors predict EID

In terms of diffusion function, the AUC of the \(\text{D}_{15}\text{CO SB}\) percent pred predicting EID was 0.689, whereas the AUC of the \(\text{D}_{15}\text{CO/VA}\) percent pred was 0.707, according to the ROC curve (Fig. 2A). The \(\text{D}_{15}\text{CO SB}\) pred and \(\text{D}_{15}\text{CO/VA}\) pred cutoff values were 50.45% and 80.9%, respectively (Table 4). The AUC of the CAT score predicting EID was 0.727 (Fig. 2B), and the cutoff value was 15 points, according to the ROC curve (Table 4). The AUCs for PD15, mean density, emphysema volume, and LAA in CT-defined emphysema were 0.691, 0.671, 0.668, and 0.656 (Fig. 2C, D), respectively, for PD15, mean density, emphysema volume, and LAA: − 955.00HU, − 856.46HU, 338.14 ml, and 7.63% were the cutoff values (Table 4).

Discussion

EID is prevalent in COPD patients and is a phenomenon that persists over time [9, 28]. Both the 2014 ERS/ATS and the 2021 Chinese version 6MWT guidelines require continuous oxygen saturation monitoring during exercise.

Table 1: Comparison of the incidence of EID by GOLD classification under different determination methods

| Determination methods | GOLD classification [n (%)] |   |   |   |   | Total (n = 124) |
|-----------------------|-----------------------------|---|---|---|---|----------------|
|                       | Stage I (n = 24)            | Stage II (n = 43) | Stage III (n = 33) | Stage IV (n = 24) |   |
| Method A              | 12 (50.0)                   | 28 (65.1)        | 20 (60.6)          | 17 (70.8)         | 77 (62.1) |
| Method B              | 10 (41.7)                   | 24 (55.8)        | 15 (45.5)          | 15 (62.5)         | 64 (51.6) |
| \(\chi^2\)           | 0.336                       | 0.778            | 1.521              | 0.375             | 2.778    |
| \(P\)                | 0.562                       | 0.378            | 0.218              | 0.540             | 0.096    |

EID: Exercise-induced desaturation; method A \(\text{SpO}_{2_{\text{min}}}-\text{SpO}_{2_{\text{end}}} > 4\%\) and/or \(\text{SpO}_{2_{\text{rest}}} < 90\%\); method B \(\text{SpO}_{2_{\text{rest}}}-\text{SpO}_{2_{\text{end}}} > 4\%\) and/or \(\text{SpO}_{2_{\text{rest}}} < 90\%\); \(\text{SpO}_{2_{\text{rest}}}\) resting blood oxygen saturation at the beginning of the test; \(\text{SpO}_{2_{\text{min}}}\) minimum blood oxygen saturation during the test; \(\text{SpO}_{2_{\text{end}}}\) blood oxygen saturation at the end of the test.
| Variables | new-EID patients (n = 13) | original-EID patients (n = 64) | P | EID patients (n = 77) | non-EID patients (n = 47) | P | P* |
|-----------|---------------------------|--------------------------------|---|----------------------|---------------------------|---|---|
| Male (n, %) | 11 (84.62%) | 56 (87.5%) | 0.778 | 67 (87.01%) | 34 (72.34%) | 0.041 | 0.063 |
| Age (y) | 63.85 ± 9.08 | 64.44 ± 10.10 | 0.846 | 64.34 ± 9.88 | 63.21 ± 11.30 | 0.562 | 0.117 |
| Height (m) | 1.67 (1.58, 1.76) | 1.64 (1.62, 1.72) | 0.623 | 1.64 ± 0.20 | 1.66 ± 0.73 | 0.544 | 0.112 |
| Weight (kg) | 62.00 (57.00, 72.00) | 64.25 (55.73, 72.30) | 0.739 | 65.46 ± 13.18 | 64.06 ± 11.68 | 0.550 | 0.003 |
| BMI (kg/m²) | 21.51 (20.84, 24.45) | 23.09 (20.58, 25.13) | 0.545 | 23.14 ± 4.24 | 23.94 ± 3.53 | 0.280 | 0.291 |
| GOLD stage (I: II: III: IV) | 2:4:5:2 | 2:10:24:15:13:17 | 0.778 | 67 (87.01%) | 34 (72.34%) | 0.041 | 0.063 |
| SpO₂end (%) | 94.00 (93.50, 96.00) | 86.00 (84.00, 91.00) | < 0.001 | 88.00 (84.00, 92.00) | 95.00 (94.00, 97.00) | < 0.001 | < 0.001 |
| Baseline 6MWT | 6MWD (m) | 397.85 ± 112.56 | 415.78 ± 92.18 | 0.540 | 411.45 ± 94.54 | 464.85 ± 100.28 | 0.003 | 0.042 |
| Lung function index | FEV₁ (L) | 1.47 (0.76, 2.60) | 1.35 (0.84, 1.87) | 0.786 | 1.43 ± 0.68 | 1.42 ± 0.59 | 0.953 | 0.650 |
| | FVC (%) | 94.69 ± 14.28 | 87.04 ± 22.83 | 0.295 | 88.24 ± 23.10 | 88.70 ± 21.46 | 0.915 | 0.046 |
| | DLCO SB (mmol/min/kPa) | 5.01 | 4.63 ± 1.74 | 0.497 | 4.69 ± 1.76 | 5.72 ± 1.52 | 0.002 | 0.117 |
| | DLCO/VA% pred (%) | 0.30 ± 0.24 | 0.30 ± 0.24 | 0.189 | 0.30 ± 0.24 | 0.30 ± 0.24 | 0.189 | 0.117 |
| | | 0.32 | 0.17 | 0.74 | 0.80 | 0.003 | 0.021 |

EID Exercise-induced desaturation; BMI Body mass index; 6MWT 6 min-walk test; 6MWD 6 min-walk distance; SpO₂ Reth blood oxygen saturation at the beginning of the test; SpO₂min Minimum blood oxygen saturation during the test; SpO₂max Blood oxygen saturation at the end of the test; FEV₁ Forced vital capacity; FEV₁%pred Forced vital capacity as a percentage of predicted value; FVC%pred Forced expiratory volume in the first second; FVC%pred Forced expiratory volume in the first second as a percentage of predicted value; DLCO SB Lung carbon monoxide diffusion; DLCO SB%pred Lung carbon monoxide diffusion as a percentage of predicted value; DLCO/VA Lung carbon monoxide diffusion per liter of alveolar air volume; DLCO/VA%pred Lung carbon monoxide diffusion per liter of alveolar air volume as a percentage of predicted value; mMRC Modified Medical Research Council; CAT COPD Assessment Test; PD15 15th percentile point; %LAA % Low attenuation area

P * means the comparison between the new-EID and non-EID groups.
The remarks about EID, on the other hand, said “drop 4% or SpO2 < 90%,” without indicating whether the decline was SpO2_{rest}–SpO2_{end} or SpO2_{rest}–SpO2_{min}. The SpO2_{end} was shown to be similar to the SpO2_{min}, but it did not reliably predict the SpO2_{min} in individuals who slowed down during exercise due to “exertion” or even rested to compensate for ventilation [30]. Chuang et al. discovered that patients who were desaturated and subsequently re-saturated had a higher FEV1/FVC ratio (P=0.01) and reduced air retention [31]. A corresponding trend can be seen in our study too. Patients who were partially desaturated and re-saturated (n=13) had lower blood oxygen variability (P<0.001). They had improved diffusion, ventilation, and less emphysema when it came to lung function and CT defined emphysema. Thus, they may be more likely to return to non-hypoxic levels in a relatively “effortless” condition at the end of exercise. Although no statistical differences were observed for the appeal indices, a numerical trend may be seen (Table 1), and perhaps a larger sample size might yield an appreciable outcome. When compared the EID cohorts functional outcomes with those without EID, there were statistically significant variations in the SpO2 changes, 6MWD, and questionnaire scores (mMRC and CAT). However, the differences in pulmonary function and CT-defined emphysema were not statistically significant. This indicates that even while patients have not yet reached cutoff levels for lung function or CT-defined emphysema, clinicians still need to consider those with a more prominent “symptom burden” and reduced walking distance (which may be causally connected to EID).

After the pandemic, the following economic recession and shrank fiscal allotment, our local community hospitals can not afford Bluetooth real-time monitoring devices, as we learned from primary physician colleagues in National COPD network. Besides, most COPD patients we encountered in clinic were in low social-economic-status and 6MWT is not covered by health insurance in some low-income provinces of China. Many physicians executed the 6MWT by having the patient wear a portable finger clip oximeter to measure SpO2 throughout the test. Since it is against the guideline for the assessor to walk alongside the patient being evaluated, it is nearly impossible to accurately obtain SpO2_{min}. In our study, Bluetooth real-time transmission technology was used so that SpO2 data could be sent to the iPad in real-time while the patient was being assessed. When the test is over, the assessor may immediately see changes in the patient’s SpO2 over the course of the 6MWT. More EID events can be discovered by finding SpO2_{min} rather than collecting the difference between SpO2_{rest} and SpO2_{end}. This also implies that by focusing on SpO2_{min}, it is mostly possible to prevent misclassifying some desaturated-resaturated patients as “non-EID patients”, hence underestimating the risk of EID in this group. The rate of EID increased with GOLD classification in both methods, with the highest rate of EID at 6MWT in patients with pulmonary function stage IV, which is consistent

| Variables | EXP(B) | 95%C.I | P  | EXP(B) | 95%C.I | P  |
|-----------|-------|--------|----|-------|--------|----|
| gender    | 0.752 | 0.300–1.883 | 0.542 |       |       |    |
| Age (y)   | 1.011 | 0.976–1.047 | 0.559 |       |       |    |
| BMI (kg/m²) | 0.951 | 0.862–1.100 | 0.282 |       |       |    |
| SpO2_{rest} (%) | 1.069 | 0.907–1.260 | 0.428 |       |       |    |
| 6MWD (m) | 0.994 | 0.990–0.998 | 0.005 | 0.996 | 0.992–1.000 | 0.071 |
| FVC% pred | 0.999 | 0.983–1.016 | 0.914 |       |       |    |
| FEV1% pred | 0.993 | 0.979–1.008 | 0.356 |       |       |    |
| D_{CO} SB pred (%) | 0.973 | 0.956–0.990 | 0.002 | 0.970 | 0.952–0.988 | 0.001 |
| D_{CO} VA pred (%) | 0.967 | 0.950–0.985 | <0.001 | 0.962 | 0.940–0.985 | 0.002 |
| mMRC dyspnea scale | 1.453 | 1.007–2.097 | 0.046 | 1.195 | 0.693–2.061 | 0.522 |
| CAT score | 1.121 | 1.055–1.3192 | <0.001 | 1.142 | 1.052–1.241 | 0.002 |
| BODE index | 1.193 | 0.999–1.424 | 0.051 |       |       |    |
| mean density (HU) | 0.983 | 0.968–0.999 | 0.036 | 0.985 | 0.969–1.000 | 0.048 |
| PD15 (HU) | 0.982 | 0.967–0.997 | 0.016 | 0.983 | 0.968–0.999 | 0.033 |
| emphysema volume (ml) | 1.001 | 1.000–1.001 | 0.019 | 1.001 | 1.000–1.001 | 0.049 |
| %LAA (%) | 1.052 | 1.011–1.093 | 0.012 | 1.045 | 1.004–1.089 | 0.033 |

EID Exercise-induced desaturation; BMI Body mass index; 6MWD 6 min-walk distance; SpO2_{rest} Resting blood oxygen saturation at the beginning of the test; FVC% pred Forced vital capacity as a percentage of predicted value; FEV1% pred Forced expiratory volume in the first second as a percentage of predicted value; D_{CO} SB pred Lung carbon monoxide diffusion as a percentage of predicted value; D_{CO} VA pred Lung carbon monoxide diffusion per liter of alveolar air volume as a percentage of predicted value; mMRC Modified Medical Research Council; CAT COPD Assessment Test; PD15 15th percentile point; %LAA % low attenuation areas
with other national and international studies [6–10]. It’s worth mentioning that even in patients with stage I, the prevalence of EID was as high as 50% (12/24) in this study. Although the short sample size could be one of the causes contributing to the high prevalence, it also implies that patients with minor airflow restriction are at risk for EID and should be of concern to doctors.

It is worth noting that a high agreement between the two approaches was reported in our study (kappa = 0.807), although method A identified more patients with EID. This highlights the significance of

![Fig. 2](image_url)

**Fig. 2.** ROC curve analysis of the $D_{1,CO}\,SB\%_{pred}$, the $D_{1,CO/VA}\%_{pred}$ (A), the CAT score (B), the mean density, the PD 15 (C), the emphysema volume, and the %LAA (D) for predicting EID. ROC receiver operating characteristic; EID exercise-induced desaturation; $D_{1,CO}\,SB\%_{pred}$ lung carbon monoxide diffusion as a percentage of predicted value; $D_{1,CO/VA}\%_{pred}$ lung carbon monoxide diffusion per liter of alveolar air volume as a percentage of predicted value; CAT COPD Assessment Test; PD15 15th percentile point; %LAA % low attenuation areas.
continuous SpO2 measurement, which is the first choice for financially accessible healthcare facilities. However, SpO2rest−SpO2end measurements alone, without the aid of sophisticated continuous monitoring devices, can effectively detect EID in hospitals with limited resources, especially when assessing those without resting hypoxemia. This is practical and economical.

Several factors contribute to the formation of EID during exercise, including increased oxygen demand during exercise, dynamic lung hyperinflation, and ventilation-perfusion mismatch [32]. Intermittent hypoxia in patients during exercise may have the same long-term consequences as persistent hypoxemia. Patients with EID exhibited a 38 m loss in 6MWD at 1-year follow-up [33], as well as a faster rate of lung function decline, more frequent acute exacerbations [3, 34], more frequent acute exacerbations [3, 6] and a higher incidence of nocturnal hypoxemia than those without EID [35]. As a result, prompt evaluation of the onset of EID and interventional treatment are critical components of conventional long-term COPD care.

Furthermore, the 6MWT guidelines consider SpO2 of less than 85% or 80% as a criterion for trial termination to limit the risk of malignant cardiac events or other adverse events [23, 29]. However, it has been suggested that SpO2 less than 80% in stable COPD patients undergoing 6MWT under the supervision of an experienced physical therapist is not associated with adverse events [36]. As a result, it’s thought that terminating the test due to a SpO2 of less than 80% in stable COPD patients isn’t necessary. When examining 6MWT in clinical practice, some COPD patients are likely to have tolerated hypoxia and believe they can complete the test despite a reduction in SpO2. Stopping the test due to a drop in SpO2 would not accurately reflect this group of patients’ true exercise capacity. Therefore, it’s critical to identify the clinical characteristics of COPD patients at risk of EID during the 6MWT so that this group of patients can be instructed to walk with oxygen and prevent having the test terminated due to hypoxia. It can reflect their true exercise capacity and lessen the risk of hypoxia-related complications. Furthermore, it aids in the early initiation of oxygen therapy prescriptions, thereby delaying the adverse prognosis induced by hypoxia.

Most studies consider diffusion function as a good predictor of EID. Patients with low D1,CO SB% pred had a higher risk of EID with a threshold of 62%, according to Hadeli et al. [37] in a large cohort (n = 8000). The ECLIPSE cohort study showed that age, female, SpO2rest ≤ 95%, D1,CO SB% pred < 50%, and FEV1% pred < 45% had a high predictive value for EID [6, 9]. Both D1,CO SB% pred and D1,CO/VA% pred were found to be good predictors of EID in our study. Furthermore, we discovered that D1,CO/VA% pred had somewhat superior predictive power (AUC = 0.707) than D1,CO SB% pred (AUC = 0.689), despite the fact that few research have examined their predictive value for EID. D1,CO SB% pred was marginally better than D1,CO/VA% pred in predicting EID in a retrospective investigation of 97 patients with respiratory disease (58 with interstitial lung disease), with both having a threshold of 55% [38], contradicting to the findings of the current study. One of the explanations for the contradictory conclusions could be the various research populations primarily targeted in the current study compared to prior study. The leading indicator of diffusion function is usually D1,CO SB% pred. However, when diffusion volume alveolar (VA) falls, D1,CO SB% pred drops, therefore the VA is commonly used in the evaluation of diffusion function to rule out the effect of lung volume on diffusion volume, i.e. D1,CO/VA. Under exercise, oxygen exchange is complicated, involving changes in pulmonary blood flow, lung capacity, and metabolism. D1,CO/VA% pred is a better predictor of EID than D1,CO SB% pred owing to the fact that D1,CO/VA% pred considers both intrinsic and volumetric diffusing capacity, making it a more comprehensive measure

Table 4  ROC curve analysis for predicting EID

| Variables                  | AUC  | Cut-off value | Sensitivity | Specificity | P     |
|----------------------------|------|---------------|-------------|-------------|-------|
| CAT score                  | 0.727| 15            | 0.753       | 0.66        | <0.001|
| D1,CO SB% pred (%)         | 0.689| 50.45         | 0.889       | 0.425       | 0.001 |
| D1,CO/VA% pred (%)         | 0.707| 80.90         | 0.733       | 0.589       | <0.001|
| mean density (HU)          | 0.671| −856.46       | 0.808       | 0.593       | 0.033 |
| PD15 (HU)                  | 0.691| −955.00       | 0.692       | 0.704       | 0.017 |
| emphysema volume (ml)      | 0.668| 338.14        | 0.689       | 0.667       | 0.003 |
| %LAA (%)                   | 0.656| 7.63%         | 0.635       | 0.667       | 0.005 |

ROC Receiver operating characteristic; EID Exercise-induced desaturation; CAT COPD Assessment Test; AUC Area under curve; D1,CO SB% pred Lung carbon monoxide diffusion as a percentage of predicted value; D1,CO/VA% pred Lung carbon monoxide diffusion per liter of alveolar air volume as a percentage of predicted value; PD15 15th percentile point; %LAA % Low attenuation areas
of diffusing capacity. In our investigation, the D<sub>1</sub>CO SB% pred threshold value for diagnosing EID was similar to the results of earlier studies. The threshold value for D<sub>1</sub>CO/VA% pred to diagnosis EID was 80.9%, indicating that included VA in the diffusing capacity study is more indicative of the likelihood of EID in COPD patients with mild diffusion impairment. It also emphasizes that in order to be more keenly aware of the likelihood of EID, doctors should pay attention to both D<sub>1</sub>CO SB% pred and D<sub>1</sub>CO/VA% pred declines in pulmonary function indices.

In this study, patients with EID in 6MWT had significantly higher mMRC scores, CAT scores, and BODE indexes than non-EID patients. It suggests that individuals with EID had more severe dyspnea, a higher symptom burden, and a worse illness outcome than patients without EID. Surprisingly, the CAT score had a stronger predictive value for EID than the other two questionnaire scores (AUC=0.727). Within 2–3 min, the CAT score measured health-related quality of life and illness symptom load. For patients who are unable to complete the 6MWT or conduct continuous SpO<sub>2</sub> measurements; who do not have access to a diffusion function test in primary care settings; or who live in low-income areas, the CAT score may be used to screen a subset of patients at high risk of developing EID. It is easy to perform, time-consuming, and has no staffing or site requirements, so it can be an option for primary and community care units as a more convenient and rapid method to assess patients with risk of EID.

CT-defined emphysema is a useful complement to subjective visual assessment. It was found significantly correlated with D<sub>1</sub>CO SB% pred, D<sub>1</sub>CO/VA% pred and FEV<sub>1</sub>% pred [17, 39]. PD15 is also the most significantly connected with D<sub>1</sub>CO/VA% pred in diffusion function, followed by %LAA [39], which explains the strong correlation of these two indicators with EID in multifactorial regression analysis. EID initiation and development are determined by the degree of emphysema [18], and investigations have demonstrated that %LAA is independently associated with EID [16, 17]. According to Marie Waattevik et al. the median %LAA in patients with COPD who formed EID in 6MWT for the first time was 12.6%, and the median %LAA in patients with COPD who developed EID in 6MWT for the second time was 21.7% [18]. In our study, the median %LAA of COPD patients who developed EID in the 6MWT was 11%, and the cut-off value for predicting EID was 7.63%. Many COPD patients who acquired EID only had a low %LAA, according to research. The lower degree of airflow limitation (higher FEV<sub>1</sub>% pred) in our population may have resulted in a relatively low %LAA, implying that even mild to moderate emphysema (%LAA 5–25%) [10] is enough to impact the development of EID. In our study, we found a more excellent value of PD15 (AUC=0.691) than %LAA (AUC=0.656) in predicting EID, which may be due to the fact that PD15 is influenced not only by the volume of emphysema but also by the amount of residual lung tissue available for gas exchange [39] and is more stable than %LAA in assessing the extent of emphysema [40]. Moreover, PD15 also diminishes with age, making it a stronger indicator of the existence and progression of emphysema [15].

Despite the fact that CT is becoming more widespread in modern medical practice, this does not indicate that all patients with COPD should have a CT scan to determine the amount of emphysema and the risk of EID; after all, CT is still an expensive and invasive procedure. However, for COPD patients who have undergone CT either as part of disease surveillance or for other comorbidities, the use of imaging information from CT to effectively assess all and localized pulmonary changes in a short period and to accurately quantify the risk of emphysema can predict the risk of EID, which can guide future individualized treatment and prognostic assessment of patients with COPD. This has implications for COPD patients’ future individualized treatment and prognostic evaluation.

There are several limitations to this study as well. First, the study’s sample size was tiny, and the data came from a single center. Second, we did not analyse cardiopulmonary comorbidities in our patients. Atrial fibrillation and hypertension have been linked to lower SpO<sub>2</sub> during the 6MWT in some studies [6, 8]. It’s unclear whether comorbidities were present in the study’s participants, which could affect the accuracy of the findings. Third, several factors influence CT-defined emphysema, including the thickness of the scan layer, the degree of obesity, and the depth of deep inspiration [41]. Other research has found that when utilizing the threshold approach to measure emphysema, the degree of emphysema increases as the layer thickness gets thinner. This pattern reduces as the threshold is raised [42], whereas, 37 individuals were scanned with a 5-mm layer thickness in the current study, which may have influenced the CT quantification results.

Conclusions
This study emphasizes the need of continuous SpO<sub>2</sub> monitoring to determine SpO<sub>2min</sub> during the 6MWT in order to capture more EID in COPD patients. Standardized criteria for EID assessment and determination should be identified for in-depth investigation in future clinical trial studies. When conditions are limited, it is also sufficient to assess EID in COPD patients by measuring SpO<sub>2rest</sub>–SpO<sub>2end</sub> ≥ 4% and/or SpO<sub>2end</sub> < 90%. In terms of predictors of EID, D<sub>1</sub>CO SB% pred and D<sub>1</sub>CO/VA%
pred, as well as CAT score and CT-defined emphysema, are all statistically significant test factors for determining EID. The above indicators were used to find COPD patients with a high risk of EID, who were advised to take oxygen in 6MWT to avoid the exercise termination due to hypoxia and to reflect the best exercise capacity; meanwhile, early intervention treatment was performed to improve patient prognosis by identifying the risk of developing EID in these patients.

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Author contributions

TY and WQ made substantial contributions to the conception or design of the work. All authors contributed toward acquisition of data for the work. LY, WQ, HS and ZH performed analysis of data for the work. LY, WQ and MS carried out interpretation of data for the work. LY drafted the work. All authors read and approved the final manuscript.

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Availability of data and materials

The data that support the findings of this study are available from Dr. Ting Yang. These data were used under license for the current study, so these data are not publicly available. However, the data are available from the authors upon reasonable request and with permission from corresponding authors.

Declarations

Ethics approval and consent to participate

This study was approved by the Institutional Review Board of China-Japan Friendship Hospital (2019-161-K110), all methods were performed in accordance with the relevant guidelines and regulations in ethics approval. All participants had signed informed consent.

Consent for publication

Not applicable.

Competing interests

All the authors state that there are no conflicts of interest related to this study.

Abbreviations

6MWT: 6-Minute-walk test; 6MWD: 6-Minute-walk distance; %LAA: % Low attenuation areas; AUC: Area under the curve; BMI: Body mass index; COPD: Chronic obstructive pulmonary disease; CAT: COPD assessment test; DlCO SB: Lung carbon monoxide diffusion; DlCO-SB% pred: Lung carbon monoxide diffusion per liter of alveolar air volume; DlCO/VA% pred: Lung carbon monoxide diffusion per liter of alveolar air volume as a percentage of predicted value; EID: Exercise-induced desaturation; FVC: Forced vital capacity; FVC% pred: Forced vital capacity as a percentage of predicted value; FEV1: Forced expiratory volume in the first second; FEV1% pred: Forced expiratory volume in the first second as a percentage of predicted value; HU: Hounsfield units; mMRC: Modified Medical Research Council; PD15: 15th percentile point; ROC: Receiver operating characteristic; SpO2: Blood oxygen saturation; SpO2r: Resting blood oxygen saturation at the beginning of the test; SpO2min: Minimum blood oxygen saturation during the test; SpO2end: Blood oxygen saturation at the end of the test; VA: Alveolar volume.

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