Utility of ultrasound assessment of diaphragmatic function before and after pulmonary rehabilitation in COPD patients

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Background: Pulmonary rehabilitation (PR) may improve respiratory symptoms and skeletal muscle strength in patients with COPD. We aimed to evaluate changes in ultrasound (US) measurements of diaphragmatic mobility and thickness after PR in COPD patients and to test its correlation with PR outcomes.

Methods: Twenty-five COPD patients were enrolled and underwent a diaphragm US assessment before and after a 12-week PR program.

Results: We found a correlation between the intraindividual percentage of change in the diaphragmatic length of zone of apposition at functional residual capacity (ΔLzapp%) and the change in 6-minute walking distance (6MWD) after PR (r=0.49, P=0.02). ΔLzapp% was significantly higher in patients with improved 6MWD and COPD Assessment Test (CAT) score (mean rank=12.03±2.57 vs 6.88±4.37; P=0.02). A ΔLzapp% of ≥10% was able to discriminate among patients with improved 6MWD, with a sensitivity of 83% and a specificity of 74%. The area under the receiver operating characteristic curve for ΔLzapp% was 0.83. A cutoff value of ≥9% of ΔLzapp% had a positive predictive value in discriminating a reduction in ≥2 points of CAT score after PR, with a sensitivity and a specificity of 80% and 62%, respectively.

Conclusion: Diaphragm US assessment represents a useful prognostic marker of PR outcomes in COPD patients.

Keywords: diaphragm ultrasound, COPD, pulmonary rehabilitation

Introduction

Pulmonary rehabilitation (PR) is a well-recognized intervention in the management of individuals with COPD that is designed to improve patients’ physical and psychosocial conditions by providing tailored interventions including exercise training, education, and behavioral changes given by an interdisciplinary team of healthcare professionals.1

There is strong evidence that exercise training lessens ventilatory requirement and reduces the degree of dynamic lung hyperinflation leading to improved arterial oxygen content and central hemodynamic responses, thus increasing systemic muscle oxygen availability.2 In previous studies, it has been shown that the combination of improved mechanical efficiency and improved respiratory and skeletal muscle strength1 may determine desensitization to dyspnea4 and consequently reduce dynamic hyperinflation.2

These physiological benefits apply to all COPD patients, irrespective of the degree of disease severity,3 and are associated with improved exercise tolerance, functional capacity, and quality of life,6 thereby reducing breathlessness and hospital admissions.
and improving recovery after exacerbation. However, the responses to PR may vary significantly among individuals. Many studies were not able to detect significant changes in forced expiratory volume in the first second (FEV\textsubscript{1}), forced vital capacity (FVC), and FEV\textsubscript{1}/FVC values after PR.\textsuperscript{7,8} Thus, a wide range of outcome measures has been described to assess clinically relevant outcomes after PR. The decision to incorporate more sophisticated tests mainly depends on the available resources and the philosophy of each PR center.\textsuperscript{10} Recently, Smargiassi et al have demonstrated the possible role of ultrasound (US) measurement of diaphragm thickness and thickening at the zone of apposition (zapp) at the end of a maximal inspiration might be a useful tool to estimate lung hyperinflation.\textsuperscript{11} Moreover, US analysis of the diaphragmatic excursion has been correlated with bronchial obstruction.\textsuperscript{12} Thus, we hypothesized that US variation of diaphragm mobility and thickness before and after a PR program may represent good markers of the effects of a successful PR program in COPD patients and that these changes may correlate with positive outcome measures postrehabilitation.

Therefore, we aimed to estimate the role of US assessment of diaphragm function in COPD patients undergoing a PR program and in the detection of postrehabilitation outcomes.

**Materials and methods**

**Study design and participants**

This prospective observational study was conducted in the Pulmonary Rehabilitation Unit at Policlinico University Hospital, Catania, Italy, between November 2017 and March 2018. Inclusion criteria were as follows: a confirmed diagnosis of COPD according to GOLD criteria,\textsuperscript{13} current or former smoke history of at least 10 pack/years with clinical indication for PR according to British Thoracic Society guidelines\textsuperscript{14} and no exacerbation in the past 12 weeks; patients with pacemaker, unstable angina, recent myocardial infarction, lung cancer, neurological diseases, pleural effusion, interstitial lung diseases, and recent major surgery were excluded.

Forty-seven out of fifty-five consecutive COPD patients (41 males, 87.3%) were referred for PR during the study period; thirty-seven of those patients met the inclusion criteria and were enrolled in the present study. Twenty-five patients fully completed the 12-week PR program and were considered for data analysis.

**Methods**

Eligible patients were assessed for pulmonary function, 6-minute walking test (6MWT), and US assessment of diaphragmatic function and of quadriceps femoris transverse section; all the measurements were done before and after PR. Dyspnea and health status were graded using the modified Medical Research Council (mMRC) dyspnea scale\textsuperscript{15} and COPD Assessment Test (CAT) scale,\textsuperscript{16} respectively; BODE index\textsuperscript{17} and Charlson index\textsuperscript{18} were also used for calculation. Written informed consent was obtained from all participants; Institutional Review Board of Policlinico Hospital, Catania, approved the study (IRB # 0017237).

**Pulmonary function tests**

Pulmonary function tests, spirometry and plethysmography (Medical Graphics, St Paul, MN, USA), were performed following standard protocols,\textsuperscript{19,20} based on reference equations.\textsuperscript{21} Arterial blood gas (ABG) analysis was performed in all patients.

**Ultrasonographic measurements**

US measurements were performed using an US machine with probes of 2–6 and 6–15 MHz (Sonoscape A6 e Logic Book GE). Diaphragm US recordings were performed as previously described\textsuperscript{11} by the same investigator (Dr Teresa Augelletti), a physician trained in internal and thoracic echography, who was blinded to patients’ PR outcomes. The excursion of right hemi-diaphragm during quiet breathing, at tidal volume (CV), and during deep breathing, at total lung capacity (TLC), starting from normal end-expiratory volume (functional residual capacity, FRC) was measured using a convex probe of 3.5 MHz with M-mode technique, and measurements were obtained at CV and at TLC. We considered normal values as previously defined.\textsuperscript{22,23} Measurements of diaphragm kinetics at the zapp were performed as previously described,\textsuperscript{24} with a probe of 6–15 MHz. Diaphragm thickness was measured at zapp using B-mode US imaging. The zapp was measured at the closest point to the “curtain sign” in which the two hyperecogenic parallel layers of diaphragm were clearly identified in the right intercostal position. We measured the distance of zapp from the skin, its thickness (Szapp), and its length (Lzapp) at different lung volumes: FRC and TLC. It was not possible to evaluate Lzapp at the end of maximal expiration (residual volume, RV) for which a longer linear probe is required. We considered only US measurements obtained at patients’ right side for statistical analysis, as left side offers a poor acoustic window in the majority of patients, as in previous studies.\textsuperscript{25} Measurements of quadriceps femoris transverse section were performed as previously described,\textsuperscript{26} taking into account the average of three subsequent measurements. The percentages of change
(Δ%) of US measurements before and after PR for diaphragm zapp were defined as ΔLzapp TLC%, ΔLzapp FRC%, ΔSzapp TLC%, ΔSzapp FRC%; the Δ% for diaphragmatic excursion was defined as ΔQuiet Breathing% and ΔDeep Breathing%. The percentage of change of rectus femoris area was outlined as ΔRectus Femoris Area%.

Pulmonary rehabilitation
All subjects participated in a 12-week, 3 sessions per week, out-patient-based PR program. The PR training included upper and lower extremity endurance and strength training. Exercise sessions were conducted for 60 minutes: 10-minute warm-up period, 40-minute aerobic activity, and 10-minute cool-down period. The aerobic activity was composed of the following: 20 minutes of lower limb endurance training by walking on a treadmill or cycling with an exercise intensity target set at a speed of 60% of the speed of their 6MWT, upper extremity endurance training (arm exercise lifting or stretching elastic bands), and strength training (weight lifting, diagonal arm raises, arm abduction into elevation and reverse, forward flexion, and reverse and straight leg rises). Work rate, SpO₂, heart rate, dyspnea scores, and leg fatigue were monitored during the sessions.

Outcome measurements
We wanted to explore patients’ interindividual changes of clinical and US parameters after PR. Moreover, to determine the clinically meaningful responsiveness and effectiveness of PR, we evaluated whether patients achieved the accepted minimal clinically important difference (MCID) for the 6MWT and for the CAT score, after the 12-week PR programs. The MCID considered for 6MWT was 54 m as previously described.²⁷ The MCID for the CAT has not been formally established but, based on its relationship with Saint George Respiratory Questionnaire (SGRQ), where a change of 4 points in the total score has been shown to represent the minimal clinically significant change and that this was equivalent to a decrease of 1.6 points in CAT score, we considered a reduction of at least 2 points in CAT score after PR as the MCID for CAT.²⁸ Furthermore, we wanted to explore if the interindividual percentage of change in Lzapp measured by US was able to discriminate between patients who “markedly improved” after PR reaching the MCID at 6MWT and CAT.

Statistical analysis
Baseline measurements and results were expressed as mean ± SD or medians and interquartile ranges for continuous variables and percentage for categorical variables. Intraindividual changes in pulmonary function, dyspnea scores, US measurements, and 6MWD post-PR were assessed by Wilcoxon-signed rank test. We used the nonparametric analysis because the assumption of normality, assessed with Shapiro-Wilk test, was not fulfilled or the presence of outliers was detected for all the variables.

Spearman rank order correlation (rho) was used to estimate the strength of relationships between the intraindividual change of Lzapp at FRC (ΔLzapp%) and PR outcome expressed as intra-individual change of 6MWD.

Based on the MCID target, at a second stage, we split our patients into two groups:
- Markedly improved: those who improved after PR and reached the minimally clinically important difference of >54 m at 6MWD and a reduction of at least 2 points in CAT score;
- Slightly improved: those who improved after PR but were not able to reach the meaningful cutoff at 6MWD and CAT.

Mann–Whitney U-test was employed for comparisons between the groups of “markedly improved” patients (who reached the MCID for 6MWT and CAT) and “slightly improved” patients (who did not achieve the MCID for 6MWT and CAT).

In addition, the receiver operating characteristic (ROC) curves were constructed to assess the ability of ΔLzapp% to discriminate among “markedly improved” and “slightly improved” patients after PR. Youden index, a function of sensitivity and specificity, was calculated to find its maximal value for the determination of an optimal cutoff point of ΔLzapp% after PR in our population. A P-value <0.05 was considered statistically significant. Statistical analyses were performed using SPSS 15.0.

Results
The general and baseline characteristics of the study population are shown in Table 1.

All patients were clinically stable and in regular treatment with long-acting beta-2 agonist; 96% took a long-acting anticholinergic and 52% (n=13) were treated with inhaled corticosteroids. Patients continued their inhaled drug therapy throughout the study period. All patients attended the entire PR program and completed the 36 sessions (1 hour/session) planned.

All the patients significantly improved after PR in terms of pulmonary function, ABG, 6MWD, mMRC, and CAT score as shown in Table 2; statistically significant improvements after PR were observed also in US parameters of both
Patients’ characteristics before and after pulmonary rehabilitation

Table 2 Patients’ characteristics before and after pulmonary rehabilitation

| Characteristics | Pre-PR | Post-PR | P-value |
|-----------------|--------|---------|---------|
| N=25            |        |         |         |
| Spirometry      |        |         |         |
| FVC%            | 83 (73–101) | 89 (76–102) | 0.020  |
| FEV₁ %          | 43 (34–50) | 48 (39–55) | 0.001  |
| PEF %           | 41 (40–50) | 45 (40–51) | 0.024  |
| FEV₁/FVC%       | 41 (34–46) | 53 (47–63) | 0.001  |
| VR%             | 132 (103–155) | 129 (100–144) | 0.002  |
| TLC%            | 105 (97–115) | 102 (97–112) | 0.040  |
| RV/TLC%         | 48 (38–52) | 45 (36–50) | <0.001 |
| Arterial blood gas |       |         |         |
| pH              | 7.41 (7.40–7.42) | 7.41 (7.40–7.42) | 1       |
| SaO₂%           | 94 (91–95) | 93 (92–95) | 0.008  |
| PaO₂ mmHg       | 65 (59–70) | 67 (60–71) | 0.016  |
| PaCO₂ mmHg      | 45 (41–49) | 43 (40–45) | 0.041  |
| 6MWD (m)        | 250 (150–350) | 300 (210–400) | <0.001 |
| CAT             | 26 (21–34) | 24 (20–30) | <0.001 |
| MRC             | 4 (3–4) | 3 (2–3) | <0.001 |
| BODE             | 6 (4–7) | 4 (3–6) | 0.001  |

Note: Data are presented as median (25th–75th percentiles).

Abbreviations: PFTs, pulmonary function tests; PR, pulmonary rehabilitation; FVC, forced vital capacity; FEV₁, forced expiratory volume in 1 second; PEF, peak forced expiratory flow; FEV₁/FVC, Tiffeneau index; RV, residual volume; TLC, total lung capacity; RV/TLC, Motley index; ABG, arterial blood gas; 6MWD, 6-minute walking distance; CAT, COPD Assessment Test; MRC, Medical Research Council dyspnea score; BODE, BODE index (Body mass index, airflow Obstruction, Dyspnea, and Exercise).

Discussion

The results of our study showed that the US assessment of diaphragmatic function was able to accurately identify COPD patients who meaningfully improved after PR.

To our knowledge, this is the first study investigating the role of diaphragm US as a possible outcome of PR in subjects with COPD.

Since the available nonimaging diagnostic tests to assess diaphragmatic function are complex and relatively
Invasive, several functional imaging techniques have been used over time; the most simple method to perform and easy to interpret is the fluoroscopy, which can assess the dome excursion through the sniff test but it involves a significant exposure to ionizing radiation.

Alternatively, chest X-ray and computed tomography (CT) have been used but the possibility of having a dynamic imaging is limited; dynamic magnetic resonance imaging (MRI) has also been described, but it has the limitation of high costs. Moreover, drawbacks of all these techniques include limited availability and the need for patient transport. Recently, Chun et al showed that fluoroscopy is an effective and cost-saving technique for evaluating pulmonary rehabilitation in a small cohort of COPD patients compared with CT or MRI, but Houston et al had already clearly stated that US has several advantages over fluoroscopy and it should be considered as the method of choice when studying diaphragm function. The US assessment of diaphragmatic function has been widely and successfully used to detect the presence of diaphragm dysfunction as a postsurgical complication, to identify the occurrence of ventilator-induced diaphragm injury, to evaluate diaphragm dome motion during spontaneous breathing weaning trials, to quantify the work of breathing and titrate ventilatory support, and to

### Table 3

| Patients’ ultrasonographic characteristics before and after pulmonary rehabilitation |
|------------------------------|-----------------|-----------------|------|
|                             | Pre-PR | Post-PR | P-value |
| Diaphragm zapp (mm)         |         |         |       |
| Lzapp TLC                   | 26 (18–31) | 23 (19–26) | 0.002 |
| Lzapp FRC                   | 38 (32–41) | 41 (35–44) | <0.001|
| Szapp TLC                   | 5 (3–6)  | 4 (3–5)  | 0.001 |
| Szapp FRC                   | 4 (3–4)  | 3 (3–4)  | 0.027 |
| Diaphragmatic excursion (mm)|         |         |       |
| Quiet breathing             | 23 (16–27) | 27 (22–31) | <0.001|
| Deep breathing              | 36 (25–53) | 50 (35–58) | <0.001|
| Rectus femoris area (m²)    | 4 (3–4)  | 4 (3–5)  |       |

Note: Data are presented as median (25th–75th percentiles).

Abbreviations: PR, pulmonary rehabilitation; zapp, diaphragm zone of apposition; Lzapp, length of diaphragm zone of apposition; Szapp, thickness of diaphragm zone of apposition; TLC, total lung capacity; FRC, functional residual capacity.
predict extubation success.42 This technique has been studied in COPD patients, showing a reduction in diaphragmatic excursion and thickening in the subset of patients with more pronounced air-trapping11,43,44 and to establish diaphragmatic dysfunction,45 but no studies have explored the use of this technique in the PR settings.

The first remarkable finding of this study is that the sonographic evaluation of the diaphragm is able to document a significant change of diaphragmatic excursion after PR in COPD patients. Our study also strengthened the results of previous studies26,46 about the utility of ultrasound measurements of the rectus femoris area as a noninvasive tool of measuring muscle mass change in patients who underwent PR.

The portability and wide availability of US make sonographic evaluation of both diaphragm and quadriceps femoris ideally suited for a routine incorporation into the PR assessment, to complement a more complete appraisal of COPD patients undergoing PR, to measure individual’s progresses, to potentially tailor PR program for each patient, and to evaluate its outcomes. Indeed, the US technique does not require any special effort, coordination, or cooperation, and therefore can be easily used also in older and more severe patients. Moreover, we can speculate that US diaphragmatic assessment may help in identifying the subpopulation of COPD patients with diaphragm muscle dysfunction and consequent ventilatory mechanics alterations that may expose patients at a major risk of reexacerbation due to a low maximum diaphragm excursion;47 therefore, this technique can be helpful in phenotyping subgroups of COPD patients, identifying those who are potentially more frail and probably at higher risk of exacerbations and it might be even used as a criterion for assigning a rehabilitation priority in the waiting list for the inpatient PR program. Furthermore, US is an accurate, reproducible, and relatively easy to learn technique that needs little learning curve; therefore, even physiotherapists could use this tool in their clinical decision-making processes before and after PR, as already reported in the literature.48

The second important finding of our study is that the US assessment of diaphragmatic function may accurately predict the response to PR. In fact, in our study the intraindividual ΔLzapp% was significantly higher in patients who markedly

| Table 4 | Patients’ percentage of change of ultrasonographic measurements after pulmonary rehabilitation between “markedly improved” and “slightly improved” patients based on MCID at 6MWT |
|-----------------|------------------|------------------|------------------------|------------------|
| Ultrasonographic measurements | 6MWD $>$ 54 m N=8 | 6MWD $<$ 54 m N=17 | P-value |
| Diaphragm zapp | | | |
| ΔLzapp TLC% | –15 (–26 to –12) | –18 (–23 to –9) | 0.791 |
| ΔLzapp FRC% | 12 (10–15) | 8 (3–11) | 0.023 |
| ΔSzapp TLC% | –5 (–8 to –2) | –5 (–10 to –3) | 0.733 |
| ΔSzapp FRC% | –6 (–9 to 0) | –7 (–11 to 0) | 0.850 |
| Diaphragmatic excursion | | | |
| ΔQuiet breathing, % | 5 (2–10) | 15 (3–44) | 0.112 |
| ΔDeep breathing, % | 6 (1–12) | 17 (5–47) | 0.132 |
| ΔRectus femoris area, % | 10 (6–15) | 8 (4–12) | 0.850 |

Note: Data are presented as median (25th–75th percentiles).

Abbreviations: MCID, minimally clinical important difference; PR, pulmonary rehabilitation; zapp, diaphragm zone of apposition; Lzapp, length of diaphragm zone of apposition; Szapp, thickness of diaphragm zone of apposition; TLC, total lung capacity; FRC, functional residual capacity; 6MWD, 6-minute walking distance; 6MWT, 6-minute walking test.

Figure 3 Difference in ΔLzapp% between patients with a minimal clinically significant difference for CAT after PR.

Abbreviations: Lzapp, length of diaphragm zone of apposition; FRC, functional residual capacity; CAT, COPD Assessment Test; PR, pulmonary rehabilitation.
Diaphragm ultrasound in pulmonary rehabilitation

Improved after PR in terms of reaching a meaningful target (MCID) at 6MWT and CAT score. Moreover, our study showed that a cut-off value of >9% Δlzapp% showed a positive predictive value in discriminating clinically meaningful improvement both in the 6MWD and CAT scores of COPD patients after PR. Therefore, these findings should remind clinicians about the importance of diaphragm evaluation using thoracic US in daily clinical practice and that the identification of this percentage of change after PR may contribute in phenotyping COPD patients with a fast and non-invasive technique.

Last but not least, a sonographic study can demonstrate to patients the improvement of the muscle function with a simple visual image; therefore, we can speculate that this might be used as a coping strategy for patients, feeling involved in the management of their disease, and hopefully improving adherence to

Table 5 Patients’ percentage of change of ultrasonographic measurements after pulmonary rehabilitation between “markedly improved” and “slightly improved” patients based on MCID at CAT

| Ultrasonographic measurements | CAT ≥ 2 | CAT < 2 | P-value |
|-------------------------------|---------|---------|---------|
| Diaphragm Zapp               |         |         |         |
| Δlzapp TLC%                  | −18 (−26 to −15) | −13 (−23 to −2) | 0.197 |
| Δlzapp FRC%                  | 11 (10–12)    | 4 (2–11)  | 0.043 |
| Δszapp TLC%                  | −4 (−8 to −3) | −6 (−10 to −2) | 0.863 |
| Δszapp FRC%                  | −5 (−10 to 0) | −8 (−12 to 0) | 0.756 |
| Diaphragmatic excursion      |         |         |         |
| ΔQuiet breathing, %          | 10 (1–15) | 38 (3–60) | 0.180 |
| ΔDeep breathing, %           | 14 (6–31) | 17 (3–47) | 0.913 |
| ΔRectus femoris area, %      | 8 (3–17)  | 8 (4–12)  | 1      |

Note: Data are presented as median (25th–75th percentiles).

Abbreviations: MCID, minimally clinical important difference; PR, pulmonary rehabilitation; Zapp, diaphragm zone of apposition; Lzapp, length of diaphragm zone of apposition; Szapp, thickness of diaphragm zone of apposition; TLC, total lung capacity; FRC, functional residual capacity; CAT, COPD Assessment Test.

Figure 4 Receiver operating characteristic (ROC) curve for Δlzapp% in relation to 6MWD.

Note: ROC curves estimate the ability of Lzapp change (as percentage of baseline) to predict a significant improvement in 6MWD after PR (AUC=0.83, cutoff ≥10%, sensitivity=83%, specificity=74%).

Abbreviations: AUC, area under the curve; Lzapp, length of diaphragm zone of apposition; FRC, functional residual capacity; 6MWD, 6-minute walking distance; 6MWT, 6-minute walking test; PR, pulmonary rehabilitation.

Figure 5 Receiver operating characteristic (ROC) curve for Δlzapp% in relation to CAT.

Note: ROC curves estimate the ability of Lzapp change (as percentage of baseline) to predict a significant improvement in CAT after PR (AUC=0.76, cutoff >9%, sensitivity=80%, specificity=62%).

Abbreviations: AUC, area under the curve; Lzapp, length of diaphragm zone of apposition; FRC, functional residual capacity; CAT, COPD Assessment Test; PR, pulmonary rehabilitation.
PR program. There are several limitations to the present study that should be mentioned. First, the relatively small number of patients might have influenced the results; second, we did not measure diaphragmatic thickness during inspiration that has been used as an indirect measurement of muscle fiber contraction; even though several different US techniques have been described so far, at the moment, no standardized approaches are recommended for a comprehensive study of diaphragm function. Third, health-related quality of life measurements such as SGRQ were not recorded for all patients and therefore not suitable for data analysis. Moreover, we did not measure the maximal static inspiratory pressure and maximal static expiratory pressure. Finally, we have to consider that US can be operator dependent; in the present study, US recordings were reported only by one physician, qualified in using ultrasonography in daily practice; therefore, the interobserver variability could not be evaluated. Nevertheless, we really believe that our results have important clinical implications that can still be generalized in the hospital-based settings.

In conclusion, our study supports the routine use of US diaphragmatic assessment before and after PR as an additional tool for the evaluation of clinical effects of PR in COPD patients. It provides a rapid, reliable, noninvasive and relatively easy-to-use approach that allows repeated measures and does not require patient’s effort or coordination, and it should be used in combination with other exams to assess PR outcomes.

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Author contributions
CC is the guarantor of the content of the manuscript, including the data and analysis, and certifies that the methods, data, and analysis set forth in this paper are truthful and accurate. TA has performed diaphragmatic ultrasound recordings to our patient population. All authors made substantial contributions to the conception and design of the analysis and interpretation of data. All authors contributed toward data analysis, drafting and revision of the paper, and agreed to be accountable for all aspects of the work. All authors provided their final approval to the version to be published. The authors are accountable for the accuracy and integrity of this work.

Disclosure
The authors report no conflicts of interest in this work.

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