ORIGINAL ARTICLE

One-year change in health status and subsequent outcomes in COPD

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ABSTRACT

Background Poor health status has been associated with morbidity and mortality in patients with COPD. To date, the impact of changes in health status on these outcomes remains unknown.

Aims To explore the relationship of clinically relevant changes in health status with exacerbation, hospitalisation or death in patients with COPD.

Methods Characteristics and health status (St George’s Respiratory Questionnaire, SGRQ) were assessed over a period of 3 years in 2138 patients with COPD enrolled in the Evaluation of COPD Longitudinally to Identify Predictive Surrogate Endpoints (ECLIPSE) study: a longitudinal, prospective, observational study. Associations between change in health status (=4 units in SGRQ score) during year 1 and time to first exacerbation, hospitalisation and death during 2-year follow-up were assessed using Kaplan–Meier plots and log-rank test.

Results 1832 (85.7%) patients (age 63.4±7.0 years, 65.4% male, FEV1 48.7±15.6% predicted) underwent assessment at baseline and 1 year. Compared with those who deteriorated, patients with improved or stable health status in year 1 have a lower likelihood of exacerbation (HR 0.78 (95% CI 0.67 to 0.89), p<0.001 and 0.84 (0.73 to 0.97), p=0.016, respectively), hospitalisation (0.72 (0.58 to 0.90), p=0.004 and 0.77 (0.62 to 0.96), p=0.023, respectively) or dying (0.61 (0.39 to 0.95), p=0.027 and 0.58 (0.37 to 0.92), p=0.019, respectively) during 2-year follow-up. This effect persisted after stratification for age and the number of exacerbations and hospitalisations during the first year of the study.

Conclusions Patients with stable or improved health status during year 1 of ECLIPSE had a lower likelihood of exacerbation, hospitalisation or dying during 2-year follow-up. Interventions that stabilise and improve health status may also improve outcomes in patients with COPD.

Trial registration number NCT00292552, registered at ClinicalTrials.gov.

INTRODUCTION

Patients with COPD have impaired disease-specific health status1 2 that may deteriorate over time.3 The St George’s Respiratory Questionnaire (SGRQ) is a disease-specific health status questionnaire reflecting a broad variety of health impacts in patients with chronic respiratory diseases.4 High baseline SGRQ total scores (ie, poor health status) have been associated with an increased number of exacerbations and hospitalisations in patients with COPD.5 Moreover, higher baseline SGRQ scores were associated with worse survival.6

An increase of four points or more in SGRQ total score can be considered as a meaningful worsening of patient’s health status.7 To date, it remains unknown whether and to what extent clinically relevant changes in disease-specific health status may affect morbidity and mortality in patients with COPD. Nevertheless, it seems reasonable to hypothesise that a worsening of disease-specific health status will be associated with worsening of other important health outcomes.

We sought to explore the impact of clinically relevant 1-year changes in SGRQ total scores on the likelihood of having an exacerbation, hospitalisation or dying in patients with COPD, using data from the Evaluation of COPD Longitudinally to Identify Predictive Surrogate Endpoints (ECLIPSE) study.8

METHODS

The ECLIPSE study is a multicentre, longitudinal, prospective 3-year observational study to identify...
novel endpoints in COPD. Participants were eligible if they were 40–75 years of age, had a smoking history of 10 or more pack-years and a diagnosis of COPD confirmed with post-bronchodilator FEV₁/FVC<0.7. Demographic and clinical characteristics were assessed as described before.9

**Health status**

Disease-specific health status was assessed using the COPD-specific version of the SGRQ (SGRQ-C) to provide SGRQ scores. The SGRQ-C has three domain scores (symptoms, activity and impact) and a total score, ranging from 0 (optimal) to 100 points (worst).4 Health status was assessed at baseline, 1, 2 and 3 years. A change of four units in total score is considered as the minimum clinically important difference.10 Only the SGRQ total scores are analysed here.

**Study outcomes**

Following the initial period of SGRQ total score change determination, during year 1, the following outcomes were determined: survival (=all-cause mortality), time to first moderate to severe exacerbation and time to first COPD hospitalisation. Exacerbations were defined as events diagnosed by a clinician that led to the prescription of antibiotics and/or corticosteroids and/or to hospitalisation, as described elsewhere.11 Hospitalisations were limited to severe exacerbations requiring hospital admissions. In ECLIPSE, adverse outcomes were assessed at each visit plus monthly phone calls.8

**Statistics**

Categorical variables were described as frequencies. Continuous variables were tested for normality and were described as mean and SD. Only patients who had complete SGRQ data at baseline and at 1-year follow-up were included in the current analyses. To compare baseline characteristics between patients who completed the first year and those who dropped out during that time, an independent sample t test or Mann–Whitney U test was used. Further, baseline characteristics were compared between patients with a clinically significant 1-year deterioration in SGRQ score (Δ≥4+ points), non-clinically significant 1-year change in SGRQ score (Δ−3.99 to +3.99 points), or a clinically significant 1-year improvement in SGRQ score (Δ≤−4 points). Continuous variables were compared using a univariate analysis of variance followed by post hoc least significance difference multiple comparisons or Kruskal–Wallis test followed by Mann–Whitney U test, as appropriate. Categorical variables were compared using χ² tests.

For the study outcomes, events were flagged and time to event observed over a period of 2 years starting after the 1-year assessment. Kaplan–Meier plots illustrate a relationship between 1-year change in SGRQ score and study outcomes. Interactions between variables were tested and Kaplan–Meier curves were constructed for the whole group and further stratified by median baseline age (>64 vs ≤64 years) and number of exacerbations (<2 vs ≥2 exacerbations) and hospitalisations (<1 vs ≥1 hospitalisations) during the first year of the study. HRs, accompanied with 95% CIs, represent the ratio of the hazard rates derived from unadjusted survival analysis. The log-rank test was used for pairwise comparisons between groups.

Kaplan–Meier curves were constructed using GraphPad Prism 5. Statistical analyses were performed using IBM SPSS statistics, V19.0.

**RESULTS**

**Baseline characteristics**

A total of 2138 patients with COPD participated in the ECLIPSE study, of whom 1832 (85.7%) underwent the 1-year assessment. They were mostly men, slightly overweight, had a moderate to severe airflow limitation, an impaired exercise capacity, and an impaired health status (table 1). Symptoms of dyspnoea, depression or fatigue occurred in 52.0%, 25.3% and 73.7% of patients, respectively.

Patients who completed the 1-year assessment had a significantly better baseline lung function, 6-min walk distance (6MWD), health status and less symptoms compared with those who dropped out during the first year (n=306, table 1).

**One-year change in health status**

Patients with a 1-year improvement in health status (n=675, 36.8%) had a worse baseline health status compared with those who remained stable or deteriorated (n=630 and n=527, respectively; table 2). Age, baseline lung function, 6MWD, modified Medical Research Council (mMRC) dyspnoea grades, depression scores and fatigue scores were comparable between groups.

**Exacerbations, hospitalisations and mortality**

During the 2-year follow-up period (years 2 and 3 of the ECLIPSE study), 1234 (67.7%) patients reported an exacerbation; of these,

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**Table 1  Baseline patient characteristics: first-year completers versus first-year dropout**

<table>
<thead>
<tr>
<th></th>
<th>1st year completers</th>
<th>1st year dropouts</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>63.4 (7.0)</td>
<td>63.9 (7.8)</td>
</tr>
<tr>
<td>Male, n (%)</td>
<td>1199 (65.4)</td>
<td>195 (63.7)</td>
</tr>
<tr>
<td>Current smokers, n (%)</td>
<td>659 (36.0)</td>
<td>114 (37.3)</td>
</tr>
<tr>
<td>BMI, kg/m²†</td>
<td>26.0 (22.7–29.6)</td>
<td>25.6 (22.4–29.5)</td>
</tr>
<tr>
<td>FEV₁, % predicteda</td>
<td>48.7 (15.6)*</td>
<td>45.8 (16.1)</td>
</tr>
<tr>
<td>FEV₁, L*</td>
<td>1.4 (0.5)*</td>
<td>1.3 (0.5)</td>
</tr>
<tr>
<td>FACIT score, pointsj,k</td>
<td>3.1 (0.9)*</td>
<td>2.9 (1.0)</td>
</tr>
<tr>
<td>CES-D score, pointsh,i</td>
<td>1.6 (1.1)**</td>
<td>1.9 (1.1)</td>
</tr>
<tr>
<td>BODE index, pointsf,g</td>
<td>3.1 (2.1)**</td>
<td>3.8 (2.3)</td>
</tr>
<tr>
<td>CES-D score, pointsl</td>
<td>11.2 (9.3)**</td>
<td>13.2 (9.3)</td>
</tr>
<tr>
<td>FACIT score, pointsk</td>
<td>35.4 (10.6)**</td>
<td>33.1 (10.7)</td>
</tr>
<tr>
<td>SGRQ symptom score, pointsz</td>
<td>60.1 (21.5)**</td>
<td>64.8 (21.1)</td>
</tr>
<tr>
<td>SGRQ activity score, pointsm</td>
<td>61.8 (21.9)**</td>
<td>68.3 (20.9)</td>
</tr>
<tr>
<td>SGRQ impact score, pointsn</td>
<td>35.5 (19.3)**</td>
<td>39.8 (20.0)</td>
</tr>
<tr>
<td>SGRQ total score, points</td>
<td>47.6 (18.3)**</td>
<td>52.7 (18.2)</td>
</tr>
</tbody>
</table>

1 Values expressed as mean (SD), median (25%-75% percentile) or number of patients (n), proportion (%).

*p<0.05; **p<0.001.

†Non-parametric tests were used for skewed data.

‡n=1182 (1st year completers), n=1791 (1st year completers), n=298 (1st year dropout), n=1787 (1st year completers), n=285 (1st year dropout), n=1743 (1st year completers), n=297 (1st year dropout), n=1801 (1st year completers), n=297 (1st year completers), n=281 (1st year dropout), n=251 (1st year dropout), n=249 (1st year dropout), n=229 (1st year dropout).

BMI, body mass index; FEV₁, forced expiratory volume in the first second; FVC, forced vital capacity; 6MWD, 6-minute walk distance; mMRC, modified Medical Research Council dyspnoea scale; BODE, body mass index, airway obstruction, dyspnoea, and exercise tolerance; CES-D, Center for Epidemiologic Studies Depression Scale; FACIT, Functional Assessment of Chronic Illness Therapy fatigue scale; SGRQ, St George’s Respiratory Questionnaire.
Chronic obstructive pulmonary disease

Table 2  Baseline patient characteristics

<table>
<thead>
<tr>
<th>Age, years</th>
<th>Whole group (n=1832)</th>
<th>SGRQ total score</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Improvement (Δ≤−4 points) (n=675)</td>
<td>No change (Δ=−3.99 to 3.99) (n=630)</td>
</tr>
<tr>
<td></td>
<td>63.4 (7.0)</td>
<td>63.0 (7.0)</td>
</tr>
<tr>
<td>Male, n (%)</td>
<td>1199 (65.4)</td>
<td>438 (64.9)</td>
</tr>
<tr>
<td>Current smokers, n (%)</td>
<td>659 (36.0)</td>
<td>258 (38.2)*</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>26.0 (22.7–29.6)</td>
<td>25.9 (22.9–29.3)</td>
</tr>
<tr>
<td>FEV₁, % predicted</td>
<td>48.7 (15.6)</td>
<td>48.8 (15.2)</td>
</tr>
<tr>
<td>FVC, L</td>
<td>3.1 (0.9)</td>
<td>3.1 (0.9)</td>
</tr>
<tr>
<td>FEV₁/FVC (%)</td>
<td>44.9 (11.5)</td>
<td>44.9 (11.4)</td>
</tr>
<tr>
<td>Exacerbations previous 12 months, n†</td>
<td>0.9 (1.2)</td>
<td>0.8 (1.2)</td>
</tr>
<tr>
<td>6MWD, m†</td>
<td>378 (119)</td>
<td>382 (112)</td>
</tr>
<tr>
<td>mMRC dyspnoea score (%)‡</td>
<td>1.6 (1.1)</td>
<td>1.7 (1.0)</td>
</tr>
<tr>
<td>BODE index, points§</td>
<td>3.1 (2.1)</td>
<td>3.0 (1.9)</td>
</tr>
<tr>
<td>CES-D score, points¶</td>
<td>11.2 (9.3)</td>
<td>11.7 (9.3)</td>
</tr>
<tr>
<td>FACIT score, points‖</td>
<td>35.4 (10.6)</td>
<td>34.8 (10.5)</td>
</tr>
<tr>
<td>SGRQ symptom score, points</td>
<td>60.1 (21.5)</td>
<td>64.1 (20.4)<strong>,</strong></td>
</tr>
<tr>
<td>SGRQ activity score, points</td>
<td>61.8 (21.9)</td>
<td>66.5 (19.6)<strong>,</strong></td>
</tr>
<tr>
<td>SGRQ impact score, points</td>
<td>35.5 (19.3)</td>
<td>40.0 (18.8)<strong>,</strong></td>
</tr>
<tr>
<td>SGRQ total score, points</td>
<td>47.6 (18.3)</td>
<td>51.8 (16.9)<strong>,</strong></td>
</tr>
<tr>
<td>One-year change SGRQ total score, points</td>
<td>−1.3 (10.7)</td>
<td>−11.8 (7.5)<strong>,</strong></td>
</tr>
</tbody>
</table>

Values expressed as mean (SD), median (25–75% percentile) or number of patients (n), proportion (%).

*p<0.05 versus ‘No change’; **p<0.05 versus ‘Deterioration’.
†Non-parametric tests were used for skewed data.
‡n=1828 (whole group), n=671 (improvement), n=1791 (whole group), n=617 (no change), n=512 (deterioration), n=1787 (whole group), n=658 (improvement), n=611 (no change), n=518 (deterioration), n=1743 (whole group), n=641 (improvement), n=667 (no change), n=504 (deterioration), n=1801 (whole group), n=598 (improvement), n=614 (no change), n=520 (deterioration), n=665 (improvement).
§BMI, body mass index; FACIT, Functional Assessment of Chronic Illness Therapy fatigue scale; FEV₁, forced expiratory volume in the first second; FVC, forced vital capacity; 6MWD, 6-min walk distance; mMRC, Medical Research Council dyspnoea scale; BODE, body mass index, airway obstruction, dyspnoea, and exercise tolerance; CES-D, Center for Epidemiologic Studies Depression Scale; SGRQ, St George’s Respiratory Questionnaire.

hospitalisation occurred in 486 (26.7%) patients. There were 113 (6.2%) deaths during 2-year follow-up, with an average time to death of 754 (183) days from baseline assessment.

Patients with a 1-year improvement had better survival than those who deteriorated (HR=0.61, 95% CI 0.39 to 0.95, p=0.027), similarly those who remained stable had better survival (HR=0.58, 95% CI 0.37 to 0.92, p=0.019 vs deterioration; figure 1A and table 3). Patients who improved or remained stable also had a lower likelihood of having a hospitalisation (HR=0.72, 95% CI 0.58 to 0.90, p=0.004 improvement vs deterioration; HR=0.77, 95% CI 0.62 to 0.96, p=0.023 no change vs deterioration; figure 1B and table 3). A similar observation was noted with risk of exacerbation (HR=0.78, 95% CI 0.67 to 0.89, p<0.001 improvement vs deterioration; HR=0.84, 95% CI 0.73 to 0.97, p=0.016 no change vs deterioration; figure 1C and table 3). The association between 1-year change in health status and exacerbation, hospitalisation and dying was affected by age and number of exacerbations and hospitalisations during year 1. Nevertheless, patients with a 1-year deterioration in health status had worst outcomes in most scenarios (see figure 1, online supplementary figures E1–E3 and table E1). The results retrieved from the SGRQ total score are comparable with the results retrieved from the SGRQ impact domain scores (see online supplementary Figure E4 and table E2).

DISCUSSION

This is the first study exploring the association between changes in health status and the likelihood of having an exacerbation, hospitalisation and dying in a large cohort of patients with COPD. Patients with a 1-year improvement or stable health status had a lower likelihood of having an exacerbation, hospitalisation or dying compared with patients who worsened.

Recent studies have identified numerous determinants of change in health status, but knowledge about the importance of such changes is scarce. In the current study, patients with a deterioration in health status during 1 year had a higher chance of an exacerbation, hospitalisation or dying during 2 years of follow-up. These findings support the recommendation by the GOLD committee to include the improvement of health status as an objective in COPD management. As stated before, ‘health status measurement is a means of quantifying, in a standardised and objective manner, the impact of disease on patients’ daily life, health, and wellbeing. It is a process that is essentially similar to a highly structured clinical history, although the end product is not a clinical impression but an objective measurement that can be used for scientific purposes. It is no more “soft” or “touchy-feely” than any well taken clinical history.’

The current study suggests a prognostic role for health status measurement that may help professionals to identify patients at increased risk of adverse health events.

The current study showed that the results retrieved from the SGRQ total score are comparable with those retrieved from the SGRQ impact domain score. This is not surprising since the SGRQ impact domain score behaves in a similar way to the total score whereas the SGRQ symptom domain score has been identified as the weakest correlate with overall health. The SGRQ impact domain covers aspects concerning social functioning and psychological disturbances related to the disease. It is...
the broadest domain with the most items that correlates best with outcomes in COPD.17 Thus, health status is rather a measure of the impact of the disease and the illness perception of the patient than a measure of general well-being. For instance, a previous study showed that patients with a poor health status were more likely to be referred to the respiratory specialist.18 Furthermore, patients whose health status improved had a decreased consultation and hospitalisation rate.19 Consequently, health status may be closely related to self-management and coping strategies. Indeed, patients with COPD specified coping as a relevant component of health status.20 Previously, psychological distress and difficulty in coping with their disease has been shown to impact mortality in patients with severe COPD.21 Therefore, self-regulation could significantly assist participants to control their individual symptoms and avoid acute exacerbations.22 Consequently, a worsening in health status may be associated with important health outcomes, such as hospitalisation and death. The current study demonstrates that patients’ perceptions and changes of patients’ perceptions about their health as assessed by SGRQ can play a meaningful role in COPD prognosis.

These data suggest that monitoring for trends in health status over time are important, but they are confined to health status measured using the SGRQ, which is too complex for use in routine practice. However there is already evidence that the shorter instruments such as the COPD Assessment Test, which were designed for routine use, can also predict exacerbations in patients already known to be at high risk.23

Even in younger patients (age ≤64 years) whose health status deteriorated during the first year, there was a greater risk of dying or an exacerbation compared with patients whose health status remained stable or improved. Bentsen and colleagues24 recently showed that younger patients and those with higher anxiety scores reported worse health status. The authors explained this phenomenon by the fact that younger patients are still learning how to cope with their disease and that older patients experience functional impairment more often as a part of getting older.24 In the current study, younger patients reported significantly higher Center for Epidemiologic Studies Depression Scale scores (ie, more symptoms of depression) and worse health status at baseline compared with older patients (see online supplementary table E3). The current study underlines the importance to pay special attention to the health status of younger patients with COPD and their perceptions of the impact of the disease.

COPD as a leading cause of death is increasing25 and the economic burden of the disease is substantial.26-28 Indeed, COPD exacerbations requiring hospitalisation are a ‘major cost driver’,29 that continue to increase.30 Consequently, prognostic measurements for the progression of the disease are important. Moreover, prevention and early treatment of exacerbations are needed. The current study indicates a need for assessing health status as a possible part of exacerbation prevention strategies. Furthermore, the current study supports the interest in stabilising health status.29 Since the current study identified a decline in health status as a contributing risk factor for getting an exacerbation, even after stratification for the exacerbation and hospitalisation frequency during year 1, it is important to interrupt this vicious circle and intervene on time. Stabilising and/or improving health status can be achieved by providing optimal pharmacological,10 medical/surgical29 and non-pharmacological treatment, including pulmonary rehabilitation.30 Finally, the enhancement of patient’s compliance with treatment is important since patients who withdrew from treatment showed a more rapid decline in health status.29 Consequently, prognostic measurements for the progression of the disease are important. Moreover, prevention and early treatment of exacerbations are needed. The current study indicates a need for assessing health status as a possible part of exacerbation prevention strategies. Furthermore, the current study supports the interest in stabilising health status.29 Since the current study identified a decline in health status as a contributing risk factor for getting an exacerbation, even after stratification for the exacerbation and hospitalisation frequency during year 1, it is important to interrupt this vicious circle and intervene on time. Stabilising and/or improving health status can be achieved by providing optimal pharmacological,10 medical/surgical29 and non-pharmacological treatment, including pulmonary rehabilitation.30 Finally, the enhancement of patient’s compliance with treatment is important since patients who withdrew from treatment showed a more rapid decline in health status.29 Consequently, prognostic measurements for the progression of the disease are important. Moreover, prevention and early treatment of exacerbations are needed. The current study indicates a need for assessing health status as a possible part of exacerbation prevention strategies.

**Figure 1** Impact of 1-year change in disease-specific health status on (A) mortality (n=1832), (B) hospitalisation (n=1823) and (C) exacerbation (n=1823).

**Table 3** HRs, accompanied with 95% CIs for the whole group

<table>
<thead>
<tr>
<th>HR (95% CI)</th>
<th>Improvement vs no change</th>
<th>No change vs deterioration</th>
<th>Improvement vs deterioration</th>
</tr>
</thead>
<tbody>
<tr>
<td>SGRQ total score</td>
<td>1.05 (0.65 to 1.69)</td>
<td>0.58 (0.37 to 0.92)</td>
<td>0.61 (0.39 to 0.95)</td>
</tr>
<tr>
<td>Survival</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospitalisation</td>
<td>0.94 (0.76 to 1.17)</td>
<td>0.77 (0.62 to 0.97)</td>
<td>0.72 (0.58 to 0.90)</td>
</tr>
<tr>
<td>Exacerbation</td>
<td>0.93 (0.82 to 1.07)</td>
<td>0.84 (0.73 to 0.97)</td>
<td>0.78 (0.67 to 0.90)</td>
</tr>
</tbody>
</table>

Bold printed numbers represent significant (p≤0.05) results. SGRQ, St George’s Respiratory Questionnaire.
Although the ECLIPSE study is a multicentre, longitudinal, prospective 3-year study that included a large number of patients worldwide, thus allowing an in-depth characterisation of the COPD population, there are relevant limitations that should be taken into account. First, the study consisted of a convenience sample of secondary care patients. Patients were eligible if they had moderate to very severe COPD, excluding patients with mild airflow limitation (GOLD stage I). Furthermore, patients who reported a better health status at baseline had a greater chance to worsen during 1-year follow-up and vice versa. Consequently, part of the effect of change in health status could be explained by a statistical artefact such as regression to the mean; however, that would not explain the association between change in health status and the other outcomes. Moreover, there was no interaction between change in SGRQ score and baseline SGRQ score. Finally, the current findings need to be interpreted in the light of the number of comparisons that were made in the present study. Nonetheless, multiple findings in the same direction, rather than a single statistically significant result, suggest that these are not due to chance alone.

To conclude, ECLIPSE COPD subjects with a 1-year improvement or stable health status had a lower likelihood of having an exacerbation, hospitalisation or dying during 2-year follow-up compared with those whose health status deteriorated. The current study supports the concept of routine monitoring health status of patients with COPD with the aim of improving or at least stabilising their health status.

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One-year change in health status and subsequent outcomes in COPD

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