

Quality of life in patients with chronic obstructive pulmonary disease and severe hypoxaemia

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Abstract

Background – Patients with severe chronic obstructive pulmonary disease (COPD) have impairment in most areas of quality of life, but a relationship between impairment of the partial pressure of oxygen in arterial blood (P_{aO_2}) and quality of life has not been established. Previous studies used general health measures such as the Sickness Impact Profile (SIP). In this study a disease-specific health measure, the St George's Respiratory Questionnaire (SGRQ), was used to examine the relationship.

Methods – Forty one patients (20 men) of median age 71 years (range 47–85) with COPD were assessed. Forced expiratory volume in one second (FEV_1), P_{aO_2} and partial pressure of carbon dioxide in the arterial blood (P_{aCO_2}) were measured, and the patients completed the SGRQ, SIP, and the Hospital Anxiety and Depression Scale (HAD).

Results – The mean (SD) values were: FEV_1 , 0.83 (0.29) litres; P_{aO_2} , 7.49 (1.03) kPa; P_{aCO_2} , 6.30 (1.05) kPa; SGRQ total score, 55.3 (18.2); SIP total score 15.4 (9.2); anxiety score, 5.7 (4.3); depression score, 5.3 (3.4). P_{aO_2} was significantly correlated with the SGRQ total scores but not with the SIP total score. The SGRQ total score also correlated with anxiety and depression. Multivariate analysis showed that depression score and P_{aO_2} were both significant covariates of the SGRQ total score.

Conclusions – These findings suggest that, in patients with severe COPD, quality of life is related to the severity of hypoxaemia, but this relationship is only detectable when using a disease-specific health measure.

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Keywords: chronic obstructive pulmonary disease, quality of life, hypoxaemia.

Patients with severe chronic obstructive pulmonary disease (COPD) complicated by chronic hypoxaemia complain of disabling breathlessness and reduced exercise capacity.^{1,2} In addition, they suffer from depression³ and impairment of cognitive function.⁴ Previous studies have shown that measures of lung function such as forced expiratory volume in one second (FEV_1) do not accurately predict ex-

ercise capacity in these patients.^{5,6} In recent years health-related quality of life questionnaires have been developed to provide standardised measurements of disturbance to daily life and well being due to disease. These have been shown to provide valid measurements of impaired health in COPD. They reflect, and aggregate together, a range of aspects of disease activity and its resulting disability and distress.^{3,7–10} Unlike the FEV_1 , these measures correlate quite well with exercise capacity.^{3,7,11,12}

In patients with COPD the severity of chronic hypoxaemia might be expected to influence quality of life. However, previous studies have shown either a weak correlation between quality of life scores and arterial oxygen tension,⁸ or none at all.^{3,11} These studies all used general health questionnaires such as the Sickness Impact Profile (SIP)¹³ or the Quality of Wellbeing Scale,⁸ rather than measures of health developed specifically for patients with COPD.

In this study a validated disease-specific health measure, the St George's Respiratory Questionnaire (SGRQ),¹⁴ has been used in addition to the SIP, to test whether the previously reported lack of association between hypoxaemia and disturbance of daily life and wellbeing was due to the choice of measurement instrument.

Methods

Patients with COPD recruited from outpatient chest clinics in East London were studied. The 41 patients (20 men) had a median age of 71 years (range 47–85). Inclusion criteria were a diagnosis of COPD, forced expiratory volume in one second (FEV_1) of less than 1.5 litres, and a clinically stable condition for at least two weeks. Exclusion criteria were the presence of other major chronic illness and inability to read. None of the patients was using long term oxygen therapy at the time of this study; patients who fitted the criteria for long term domiciliary oxygen therapy were subsequently commenced on this treatment.

Each patient was asked to complete three questionnaires: the SIP, SGRQ, and Hospital Anxiety and Depression scale (HAD).¹⁴ All questionnaires were self-completed with supervision. The same person supervised administration of the questionnaires in all cases. Non-directive guidance was given on the few occasions when patients had queries about how to respond to the questionnaire.

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The SIP has 136 response items which produce a range of summary component scores, including physical, psychosocial and an overall total score. Each of these scores ranges from 0 to 100%, a score of 0% indicating no disability. In a previous study almost all healthy control subjects matched for age, sex, and education with a group of patients with COPD had SIP summary scores of less than 10%.¹¹

The SGRQ has 76 items which are weighted to produce three component scores (symptoms, activity, and impact) and a total score. Each of these scores ranges from 0 to 100, a score of 100 indicating maximum disability.

The HAD is a 14 item instrument which produces separate scores for anxiety and depression ranging from 0 to 21; a score of over 10 is judged to be a predictor of a clinical diagnosis of anxiety or depression.

The three questionnaires were completed at one sitting in a randomised order. The patients also underwent spirometric tests and blood gas analysis at rest using arterialised earlobe capillary samples.¹⁵ The whole assessment was repeated after an interval of two to three weeks, each patient using the same order of questionnaires as had been used at the first assessment.

The study was approved by the ethics committee of the Royal Brompton National Heart and Lung Hospitals, and all patients gave written informed consent to participation in the study.

STATISTICAL ANALYSIS

The distribution of all variables approximated to normal so parametric tests were used throughout. Significance was accepted at the $p < 0.05$ level.

Results

Physiological data for the 41 patients from arterial blood gas analysis and spirometric tests

Table 1 Mean (SD) physiological variables of study patients ($n = 41$)

Variable	Assessment 1	Assessment 2
FEV ₁ (l)	0.83 (0.29)	0.81 (0.29)
FEV ₁ (% pred)	41.6 (16.7)	40.6 (15.4)
FVC (l)	1.85 (0.69)	1.91 (0.70)
FEV ₁ /FVC (%)	46.8 (13.7)	47.6 (17.2)
PaO ₂ (kPa)	7.49 (1.03)	7.45 (0.76)
Paco ₂ (kPa)	6.30 (1.05)	6.19 (0.85)
Sao ₂ (%)	86.6 (7.0)	87.9 (3.6)

FEV₁ = forced expiratory volume in one second; FVC = forced vital capacity; PaO₂, Paco₂ = partial pressure of oxygen and carbon dioxide in arterial blood; Sao₂ = arterial oxygen saturation.

Table 2 Mean (SD) quality of life scores for study patients ($n = 41$)

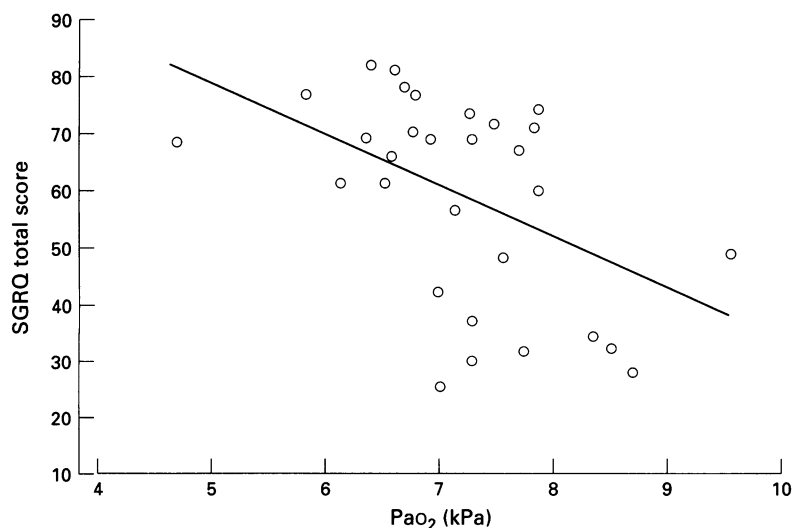
Health measure	Assessment 1	Assessment 2
SGRQ		
Symptoms	64.0 (21.9)	65.8 (23.7)
Activity	65.2 (19.8)	62.8 (18.5)
Impacts	46.5 (23.5)	45.1 (22.2)
Total	55.3 (18.2)	54.0 (16.0)
SIP		
Physical	15.4 (11.3)	17.7 (12.0)
Psychosocial	13.0 (10.6)	13.5 (9.7)
Total	15.4 (9.2)	16.7 (8.6)
HAD		
Anxiety	5.7 (4.3)	5.3 (3.5)
Depression	5.3 (3.4)	5.5 (3.2)

SGRQ = St George's Respiratory Questionnaire; SIP = Sickness Impact Profile; HAD = Hospital Anxiety and Depression scale.

on the two occasions are shown in table 1. All patients had evidence of severe chronic airways obstruction with mean FEV₁ of 41.6% predicted. The mean (SD) scores from the questionnaires on the two occasions are shown in table 2. Paired *t* tests did not demonstrate any significant differences between visits for questionnaire scores, PaO₂, Paco₂, or FEV₁. There were significant correlations between PaO₂ and FEV₁ ($r = 0.38$, $p = 0.01$), PaO₂ and Paco₂ ($r = 0.38$, $p = 0.01$), and between FEV₁ and Paco₂ ($r = 0.36$, $p = 0.02$).

There was a linear relationship between PaO₂ and SGRQ total scores at the first assessment ($r = 0.41$, $p < 0.01$) as shown in the figure. This relationship was confirmed at the second assessment ($r = 0.57$, $p < 0.001$). Analysis of covariance showed no significant difference in the regression slopes on the two occasions. There was no significant correlation between FEV₁ and SGRQ total score on either occasion. An association of marginal significance was found between Paco₂ and SGRQ total score on both assessments (visit 1, $r = 0.32$, $p < 0.05$; visit 2, $r = 0.31$, NS). Correlations between the scores for the component scores of the SGRQ and physiological variables at the first assessment are shown in table 3. The SGRQ symptoms and impacts scores were correlated with level of hypoxaemia, but the activity score was not. Age was not significantly correlated with any SGRQ score ($r \leq 0.1$; NS).

There were no significant correlations between SIP scores and FEV₁, PaO₂, or Paco₂ (table 3). The SGRQ total scores and the SIP total scores were significantly correlated with each other ($r = 0.65$, $p < 0.001$).



Relationship between quality of life (SGRQ total score) and degree of arterial PO₂ hypoxaemia ($n = 41$, $r = 0.41$, $p < 0.01$).

Table 3 Pearson correlation coefficients between quality of life measures and physiological variables ($n = 41$)

	FEV ₁	PaO ₂	PaCO ₂
SGRQ			
Symptoms	0.03	0.34*	0.10
Activity	0.34*	0.27	0.40*
Impacts	0.27	0.37*	0.24
Total	0.30	0.41**	0.32*
SIP			
Physical	0.10	0.20	0.07
Psychosocial	0.02	0.29	0.02
Total	0.14	0.28	0.13
HAD			
Anxiety	0.14	0.36*	0.37*
Depression	0.26	0.17	0.18

FEV₁ = forced expiratory volume in one second; PaO₂, PaCO₂ = partial pressure of oxygen and carbon dioxide in arterial blood; SGRQ = St George's Respiratory Questionnaire; SIP = Sickness Impact Profile; HAD = Hospital Anxiety and Depression scale.

* $p < 0.05$; ** $p < 0.01$.

Anxiety showed significant correlations with SGRQ total score ($r = 0.50$, $p = 0.001$), SIP total score ($r = 0.44$, $p < 0.005$), and was correlated with the PaO₂ ($r = 0.36$, $p < 0.05$) but not with FEV₁ or PaCO₂. Depression was significantly related to SGRQ total score ($r = 0.62$, $p < 0.001$) and the SIP total score ($r = 0.59$, $p < 0.001$). There was no significant correlation found between depression and any physiological variable (table 3).

To test whether the relationship between PaO₂ and SGRQ total score was related to other factors such as FEV₁, PaCO₂ or mood state, a number of multivariate models were generated. In a model incorporating PaO₂ and FEV₁ the PaO₂ was the only significant covariate of the SGRQ total score ($p < 0.05$). A similar model including PaO₂ and PaCO₂ again showed that PaO₂ was the only significant covariate. In a model incorporating PaO₂ and anxiety, PaO₂ was a significant covariate of the SGRQ total ($p < 0.002$) whereas anxiety was not. Finally, in a model incorporating PaO₂ and depression both covariates were significant (PaO₂, $p < 0.05$; depression, $p = 0.0001$).

Discussion

This is the first study to report a correlation between degree of hypoxaemia and quality of life scores in COPD. This correlation was statistically significant even though a relatively small number of patients was studied. The observation appears to have been due to the use of a questionnaire designed specifically for use in patients with airflow limitation. In this study, as in previous ones, attempts to correlate hypoxaemia with impaired health measured with a general health questionnaire failed.³ General health measures were developed to provide an estimate of impaired health due to a wide range of diseases. As a result, the proportion of items relevant to any particular disease may be low. A previous review of the SIP concluded that this instrument is relatively insensitive in patients with asthma and COPD because its scores remain in the mildly abnormal range until the FEV₁ falls below 50% of the predicted normal.¹⁰

Disease-specific questionnaires were developed because of the potential insensitivity

of the general measures. All of the items in such questionnaires are relevant to the condition under study. Those included in the SGRQ were selected because they were relevant to the daily lives and well being of the patients, not because of a demonstrated relationship to a physiological variable. In view of this, we conclude that the correlation between hypoxaemia and SGRQ score was not due to an inbuilt bias in the questionnaire that would unduly have favoured a correlation with hypoxaemia. The detection of a correlation with this questionnaire was presumably due to the high content of items relevant to COPD. The correlation between PaO₂ and quality of life was found on two separate occasions which makes it unlikely that this was a chance observation.

No significant correlations were found between FEV₁ and any quality of life score. Previous studies have also shown such relationships to be weak.^{8,9,11} In addition, there were no patients with an FEV₁ value of more than 1.5 litres in the current study so the range of measurements was small, making the detection of a weak correlation more difficult. Our finding of an association between quality of life and anxiety and depression confirms findings from other studies.^{3,7,11,16} In general, anxiety and depression have been found to be better predictors of quality of life scores than physiological parameters. In this study, multivariate analysis showed that the effect of PaO₂ on quality of life was independent of the effect of mood state.

Quality of life questionnaires provide a standardised method of measuring the impact of disease on the health and well being of patients. They attempt a comprehensive coverage but, for reasons of length and practicality, they cannot address all of the possible disturbances that patients may suffer. Furthermore, they have been developed and validated in populations of patients and thus address areas of commonality between patients with the disease in question and do not provide a description of the precise manner in which each individual's life is disturbed. A distinction should therefore be drawn between the quality of life of an individual patient with COPD and the standardised estimate of their impaired health provided by a quality of life questionnaire. It is perhaps unfortunate that the term "quality of life" has been attached to this type of questionnaire. There are many influences upon the quality of life of an individual with COPD. The conclusion to be drawn from this study is that the level of hypoxaemia is one potentially influential factor and, furthermore, one that appears to be independent of the degree of airways obstruction as measured by the FEV₁.

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