

Associations between the psychological health of patients and carers in advanced COPD

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Objective: Anxiety and depression are highly prevalent in patients with COPD and their informal carers, and associated with numerous risk factors. However, few studies have investigated these in primary care or the link between patient and carer anxiety and depression. We aimed to determine this association and factors associated with anxiety and depression in patients, carers, and both (dyads), in a population-based sample.

Materials and methods: This was a prospective, cross-sectional study of 119 advanced COPD patients and their carers. Patient and carer scores ≥ 8 on the Hospital Anxiety and Depression Scale defined symptoms of anxiety and depression, χ^2 tests determined associations between patient and carer symptoms of anxiety/depression, and χ^2 and independent *t*-tests for normally distributed variables (otherwise Mann–Whitney *U* tests) were used to identify other variables significantly associated with these symptoms in the patient or carer. Patient–carer dyads were categorized into four groups relating to the presence of anxious/depressive symptoms in: both patient and carer, patient only, carer only, and neither. Factors associated with dyad symptoms of anxiety/depression were determined with χ^2 tests and one-way analysis of variance for normally distributed variables (otherwise Kruskal–Wallis tests).

Results: Prevalence of symptoms of anxiety and depression was 46.4% ($n=52$) and 42.9% ($n=48$) in patients, and 46% ($n=52$) and 23% ($n=26$) in carers, respectively. Patient and carer symptoms of anxiety/depression were significantly associated. Anxious and depressive symptoms in the patient were also significantly associated with more physical comorbidities, more exacerbations, greater dyspnea, greater fatigue, poor mastery, and depressive symptoms with younger age. Symptoms of carer anxiety were significantly associated with being female and separated/divorced/widowed, and depressive symptoms with younger age, higher educational level, and more physical comorbidities, and symptoms of carer anxiety and depression with more unmet support needs, greater subjective caring burden, and poor patient mastery. Dyad symptoms of anxiety/depression were significantly associated with greater patient fatigue.

Conclusion: Symptoms of anxiety and depression in COPD patients and carers are significantly associated. Given their high prevalence, considerable impact on mortality, impact on quality of life and health care use, and associations with each other, screening for and addressing patient and carer anxiety and depression in advanced COPD is recommended.

Keywords: COPD, anxiety, depression, informal carers, patient–carer dyad

Plain language summary

Anxiety and depression are very common in patients with COPD and family or friends who provide care and support (carers). We wanted to find out whether these psychological health problems in patients and carers were linked and what the risk factors are for patients, carers, or both being anxious or depressed. Knowing these might help us target patients and carers most at risk and improve the management of psychological health in COPD. We collected information from 119 patients with advanced COPD and their carers. Up to 40% of these patients and

carers had symptoms of anxiety and/or depression. Anxious and depressed patients were more likely to have anxious and depressed carers. Symptoms of anxiety and depression were more common in patients who were more symptomatic and those with poor control of their lung problem and other physical health problems. Symptoms of depression alone were found more in patients who were younger. More anxious carers were female and separated and divorced or widowed, and more depressed carers were younger, more highly educated, and had physical health problems themselves. Carers who were either more anxious or depressed had more unmet support needs, felt they had a higher caring burden, and had patients with less control of their lung problem. Symptoms of anxiety or depression in both a patient and their carer were linked to patient fatigue. Our results suggest that it is important to screen for and address anxiety and depression not only in patients with COPD but also carers, especially of patients with anxiety or depression.

Introduction

Anxiety and depression are common in COPD, with reported prevalence ranges of 10%–55% and 7%–42%, respectively.^{1–3} They decrease treatment adherence and are associated with higher mortality,^{2–4} poorer health-related quality of life,^{1–8} and increased hospital admissions.^{2,3,9} Previous studies have identified numerous risk factors for anxiety and depression in patients, including younger age,^{4,7,8,10} being female,^{4,5,8,10} living alone,⁶ current smoking,^{2–4,8,10} and comorbidities.^{3,4} Poor psychological health preferentially coexists with frequent exacerbations,^{4,8,9} dyspnoea,^{4,5,8,10–12} fatigue,⁴ impaired physical functioning and reduced exercise capacity,^{2–4,6,8,10} and low self-efficacy.^{2,9}

As COPD progresses and becomes advanced, patients become increasingly dependent on informal carers (family members or friends). Anxiety and depression are known to be as prevalent in carers as in patients, with up to 63% reporting clinically significant anxiety symptoms and 51% reporting depressive symptoms.^{13,14} Older age,^{13,15,16} being female,^{13,15,17,18} and spousal patient–carer relationship^{13,14} have been associated with carer psychological morbidities. Low educational level,^{13,16} lack of support,^{19,20} and poor carer health¹⁸ are also risk factors, as are disease factors such as severity,^{13–15} exacerbations,²⁰ and dyspnea.^{13,14} Subjective carer burden (carer's appraisal of the impact of the role on their health, social and occupational life, finances, and reactions to caring)^{13–15,18,21} has been significantly associated with anxiety and depression, but the evidence for objective carer burden (such as time spent caregiving) is mixed.^{13,14,21}

However, the majority of studies have investigated psychological health in secondary-care cohorts (eg, patients

admitted to hospital for exacerbations), which does not represent the whole population of people living with advanced COPD.^{14,18,19} Few studies have been conducted in the primary-care setting.⁷ Also, while they have been studied separately, less is known about the association between patient and carer anxiety and depression, and factors associated with both patient and carer anxiety and depression in the dyad are unknown. We aimed to determine the association between patient and carer anxiety and depression and factors associated with anxiety and depression in patients, carers, and both in the dyad in a population-based cohort, sampled from primary care.

Materials and methods

Study design and participants

This cross-sectional study was an analysis of data from patients and carers who participated in baseline interviews for the longitudinal interview-study component of the larger prospective Living with Breathlessness study.²² In the longitudinal interview study, 652 patients with advanced COPD were identified by 63 primary-care practices in the east of England and mailed a recruitment pack: 248 patients agreed to participate, of whom 235 consented and participated (36% participation rate). Targeting of practice types maximized their variability to achieve representativeness. Patients with well-characterized advanced COPD were recruited, who met at least two of six inclusion criteria: forced expiratory volume in 1 second (FEV_1) <30%, two or more exacerbations requiring prednisolone or antibiotics in the previous year, one or more hospital admission for COPD in the previous 2 years, long-term oxygen therapy, cor pulmonale and Medical Research Council dyspnea scale score ≥ 4 . Exclusion criteria were serious mental health problem, serious learning difficulty, active cancer, and active alcoholism. Patients were asked to identify their primary informal carer, who was also invited to participate. Carer-exclusion criteria were under 18 years of age and unable to give informed consent; 119 carers consented and effectively participated (51% participation rate). The study received ethics approval from the National Research Ethics Service Committee East of England – Cambridge South (12/EE/0163). All participants gave written informed consent.

Data collection

Data were collected during face-to-face interviews with patients and carers in their home setting. Interviews with patients and carers were conducted separately wherever

possible; occasionally, the physical layout of the home made separate interviews impossible. Sociodemographics and clinical characteristics of participants were obtained. Data on the caring situation (patient–carer kin relationship, cohabitation, duration of caring, and hours of caring per week) were also collected. Patients completed the 20-item Chronic Respiratory Questionnaire (CRQ; self-administered standardized version)²³ to measure disease-specific health-related quality of life: items scored on a 7-point scale contribute to four subdomains of dyspnea, fatigue, emotional function, and mastery (how in control the patient feels of their condition), each scored 0–7, with higher scores indicating better health-related quality of life. Carers rated subjective caring burden on the Family Appraisal of Caregiving Questionnaire – palliative care, addressing caregiver strain (feelings of fatigue, isolation, lack of control, negative impact on own health, social life, occupational life, and finances), caregiver distress (anxiety and depression), and positive caregiving appraisals (commitment, satisfaction, stronger relationship), with each statement scored on a 5-point Likert scale ranging from strong agreement to strong disagreement.²⁴ Carers identified unmet support needs with the Carer Support Needs Assessment Tool, which covers 14 support domains in two groupings of support to enable the carer to provide care for the patient (eg, managing symptoms, including giving medicines) and direct support for carers' own health and well-being (eg, dealing with feelings and worries), with the need for more support indicated on a 4-point categorical scale from 0 (none) to 3 (very much more).²⁵ Patient and carer psychological health was assessed with the 14-item Hospital Anxiety and Depression Scale (HADS), a well-validated and reliable self-report screening tool for symptoms of anxiety and depression, which is analyzed as two separate 7-item subscales for anxiety (HADS-A) and depression (HADS-D), each with a total score of 0–21.²⁶ Of the 119 patients who had a participating carer, 112 patients completed the HADS questionnaire; of the 119 carers, 113 carers provided responses to the questionnaire. Symptoms of anxiety and depression were defined as HADS-A and HADS-D score ≥ 8 , the commonly used threshold for mild symptoms of anxiety and depression.²⁷

Statistical analysis

To determine associations between patient and carer symptoms of anxiety and depression, χ^2 tests were used; χ^2 and independent *t*-tests for normally distributed variables (otherwise Mann–Whitney *U* tests) were used to identify other

variables significantly associated with anxious or depressive symptoms in the patient or carer. Dyads were then categorized into four groups relating to the presence of anxious or depressive symptoms: both patient and carer, patient only, carer only, and neither. Factors significantly associated with symptoms of anxiety or depression in both (dyad symptoms of anxiety or depression) were determined with χ^2 tests and one-way analysis of variance (with Bonferroni post hoc tests) for normally distributed variables (otherwise Kruskal–Wallis tests). Statistical significance was set at $P \leq 0.05$. All statistical analyses were performed in SPSS version 23 (IBM, Armonk, NY, USA).

Results

The mean age (\pm SD) of patients and carers was 71.4 \pm 8.7 years and 64.2 \pm 14.5 years; 39.3% of patients and 72.6% of carers were female. The prevalence of symptoms of anxiety and depression was 46.4% ($n=52$) and 42.9% ($n=48$) in patients, and 46% ($n=52$) and 23% ($n=26$) in carers, respectively. Symptoms of patient anxiety were significantly associated with carer anxiety ($\chi^2=6.475$, $P=0.011$) and depression ($\chi^2=3.86$, $P=0.049$). Symptoms of patient depression were also significantly associated with both carer anxiety ($\chi^2=6.249$, $P=0.012$) and depression ($\chi^2=11.032$, $P=0.011$). Additional factors associated with patient and carer anxious and depressive symptoms are shown in Tables 1 and 2.

Symptoms of patient depression were significantly associated with younger age, and symptoms of anxiety and depression with each other, more patient physical comorbidities, more exacerbations at home, greater dyspnea, greater fatigue, and poor mastery. Symptoms of carer anxiety were associated with being female and separated/divorced/widowed, symptoms of depression with younger age, higher educational level, and more carer physical comorbidities, and symptoms of anxiety and depression with each other, more unmet support needs, greater subjective caring burden, and poor patient mastery.

The number of dyads in which both patient and carer had symptoms of anxiety or depression was 37, in which only the patient had these symptoms 28, in which only the carer had these symptoms 15, and in which neither had these symptoms 29. Dyad symptoms of anxiety or depression were significantly associated with lower mean patient CRQ fatigue score (representing higher fatigue) compared to dyads in which only the patient (2.5 vs 3, $P=0.049$), the carer (2.5 vs 4.2, $P<0.0005$), or neither (2.5 vs 4, $P<0.0005$) had symptoms of anxiety or depression (Figure 1).

Table 1 Factors associated with symptoms of patient anxiety and depression

Factors	Patients with symptoms of anxiety (n=52)	Patients without symptoms of anxiety (n=60)	P-value	Patients with symptoms of depression (n=48)	Patients without symptoms of depression (n=64)	P-value
Age, years	70 (65–74)	73 (66–78.8)	0.135	66 (62–73)	74.5 (69.3–80)	<0.0005*
Sex (female), n (%)	20 (38.5)	24 (40)	0.868	18 (37.5)	26 (40.6)	0.738
Marital status, n (%)						
Single	1 (1.9)	0	0.581	0	1 (1.6)	1
With partner	45 (86.5)	50 (84.7)		41 (85.4)	54 (85.7)	
Separated/divorced/widowed	6 (11.5)	9 (15.3)		7 (14.6)	8 (12.7)	
Living alone, n (%)	5 (9.6)	6 (10)	0.946	5 (10.4)	6 (9.4)	1
Educational level, n (%)						
None	27 (67.5)	25 (52.1)	0.084	24 (66.7)	28 (53.8)	0.485
A level/O level	9 (22.5)	9 (18.8)		6 (16.7)	12 (23.1)	
Higher	4 (10)	14 (29.2)		6 (16.7)	12 (23.1)	
Income, n (%)						
<£220/week	10 (20)	9 (16.7)	0.903	11 (25)	8 (13.3)	0.3
£221–£350/week	20 (40)	22 (40.7)		17 (38.6)	25 (41.7)	
>£350/week	20 (40)	23 (42.6)		16 (36.4)	27 (45)	
Patient–carer spousal relationship, n (%)	39 (76.5)	50 (84.7)	0.271	36 (76.6)	53 (84.1)	0.32
Patient–carer cohabitation, n (%)	44 (84.6)	54 (90)	0.39	41 (85.4)	57 (89.1)	0.564
Smoking status, n (%)						
Current smoker	9 (17.3)	9 (15)	0.92	11 (22.9)	7 (10.9)	0.22
Former smoker	42 (80.8)	49 (81.7)		36 (75)	55 (85.9)	
Never smoked	1 (1.9)	2 (3.3)		1 (2.1)	2 (3.1)	
Duration of COPD	10 (6.3–15)	8 (5–13)	0.129	7 (4–13)	6.5 (3–12)	0.366
Disease-severity measures						
% predicted FEV ₁ , mean (SD)	43.7 (22.3)	34.9 (19.2)	0.154	37.8 (26.5)	38.3 (17.2)	0.941
mMRC dyspnea scale	4 (3–5)	4 (3–4)	0.507	4 (3–5)	4 (3–4)	0.226
Exacerbations requiring admission, n	0 (0–1)	0 (0–1)	0.652	0 (0–1)	0 (0–1)	0.972
Exacerbations at home, n	3 (2–5.5)	2 (0–3)	0.001*	3 (2–6)	2 (0–3)	0.011*
CRQ domains						
Dyspnea	1.8 (1.4–2.8)	2.8 (2.2–4)	<0.0005*	1.8 (1.4–2.6)	3 (2.2–4)	<0.0005*
Fatigue	2.5 (2–3)	4.3 (3.1–4.9)	<0.0005*	2.5 (2–3)	3.8 (3–4.9)	<0.0005*
Mastery, mean (SD)	3.4 (1)	5.1 (1)	<0.0005*	3.6 (1.2)	4.8 (1.2)	<0.0005*
Patient comorbidities						
Physical comorbidities, ^a n	4 (3–6)	3 (2–4)	0.002*	4 (3–6)	3.5 (2–4)	0.002*
Anxiety symptoms, n (%)	NA	NA	NA	34 (70.8)	18 (28.1)	<0.0005*
Depression symptoms, n (%)	24 (65.4)	14 (23.3)	<0.0005*	NA	NA	NA

Notes: ^aAny of allergies, arthritis, asthma, back pain, cancer, diabetes, epilepsy, cardiac disease, hypertension, renal disease, hepatic disease, or stroke. All values medians (IQR) unless otherwise stated. * $P \leq 0.05$.

Abbreviations: FEV₁, forced expiratory volume in 1 second; mMRC, modified Medical Research Council; CRQ, Chronic Respiratory Questionnaire; NA, not applicable.

Discussion

To our knowledge, this is the first study to show that symptoms of anxiety and depression in patients and carers living with advanced COPD are significantly associated with each other in a population-based sample. One mechanism for this may be emotional contagion, where individuals in close relationships mimic each other's emotions (seen in facial expressions), with stronger associations in more severe distress.²⁸

It is possible to interpret these findings within Rolland's family-systems illness model to help our understanding.^{29–32} The relapsing unpredictable disease course of COPD may be particularly taxing for families to adjust to, resulting in

substantial psychological incongruity between periods of chronic disease and relapse, and anticipatory anxiety. There is also the risk of families coming together to deal with exacerbations and hospital admissions, but remaining “enmeshed”, which may be maladaptive in the chronic phase of COPD, leading to family burnout and permanent distortion of the patient–family relationship. As the patients in our study had advanced COPD, families may also have been living in the terminal phase, with issues of loss and need for family reorganization. Taken together, these factors may contribute to the high prevalence of symptoms of anxiety and depression in both the patient and carer cohorts.

Table 2 Factors associated with symptoms of carer anxiety and depression

	Carers with symptoms of anxiety (n=52)	Carers without symptoms of anxiety (n=61)	P-value	Carers with symptoms of depression (n=26)	Carers without symptoms of depression (n=87)	P-value
Age, years	63 (53–71)	68.5 (60.3–77)	0.079	60 (51–69.3)	69 (60–76)	0.017*
Sex (female), n (%)	44 (84.6)	38 (62.3)	0.008*	20 (76.9)	62 (71.3)	0.57
Marital status, n (%)						
Single	3 (5.9)	4 (6.7)	0.023*	1 (3.8)	6 (7.1)	0.635
With partner	40 (78.4)	55 (91.7)		22 (84.6)	73 (85.9)	
Separated/divorced/widowed	8 (15.7)	1 (1.7)		3 (11.5)	6 (7.1)	
Living alone, n (%)	1 (1.9)	1 (1.6)	1	0 (0)	2 (2.3)	1
Educational level, n (%)						
None	28 (66.7)	23 (52.3)	0.131	12 (60)	39 (59.1)	0.044*
A/O level	5 (11.9)	13 (29.5)		1 (5)	17 (25.8)	
Higher	9 (21.4)	9 (18.2)		7 (35)	10 (15.2)	
Income, n (%)						
<£220/week	12 (26.1)	5 (9.8)	0.109	5 (21.7)	12 (16.2)	0.797
£221–£350/week	16 (34.8)	22 (43.1)		8 (34.8)	30 (40.5)	
>£350/week	18 (39.1)	24 (47.1)		10 (43.5)	32 (43.2)	
Patient–carer spousal relationship, n (%)	40 (76.9)	51 (83.6)	0.371	21 (80.8)	70 (80.5)	1
Patient–carer cohabitation, n (%)	43 (84.3)	55 (93.2)	0.135	22 (84.6)	76 (90.5)	0.473
Duration of caring	6 (2.5–12)	7 (3–13)	0.395	9.5 (3–13.5)	7 (3–12)	0.728
Hours of caring/week, n (%)						
<19	18 (37.5)	23 (41.8)	0.687	9 (36)	32 (41)	0.143
20–49	15 (31.3)	13 (23.6)		4 (16)	24 (30.8)	
>50	15 (31.3)	19 (34.5)		12 (48)	22 (28.2)	
Patient disease-severity measures						
% predicted FEV ₁ , mean (SD)	39.1 (22.2)	37.5 (19.1)	0.774	41.8 (25.2)	40 (18.7)	0.463
mMRC dyspnea scale	4 (3–4.3)	4 (3–5)	0.602	4 (3–5)	4 (3–4)	0.143
Exacerbations requiring admission, n	0 (0–1)	0 (0–1)	0.939	0 (0–1)	0 (0–1)	0.545
Exacerbations at home, n	2 (1–4.5)	2 (0–4)	0.085	2 (1.5–5)	2 (0–4)	0.214
Subjective caring burden (FACQ)						
Not enough time for myself	3 (2–4)	2 (2–3)	0.002*	3 (2.5–4)	2 (2–3)	0.003*
Losing control over my own life	3 (2–4)	2 (1–2)	<0.0005*	3 (2.5–4)	2 (1–2)	<0.0005*
Feel tired and run-down	3 (2–4)	2 (1–3)	<0.0005*	4 (2.5–5)	2 (1.5–3)	<0.0005*
Isolated and alone	3 (2–4)	2 (1–2)	0.002*	3 (2–4)	2 (1–3)	0.001*
Guilty about not being able to do more	4 (2–4)	2 (2–3)	0.001*	4 (3–5)	2 (2–4)	<0.0005*
My own health has suffered	3 (2–4)	2 (1–2)	<0.0005*	3 (3–4.5)	2 (1–2)	<0.0005*
Had to give up social life	2 (2–3)	2 (1–2)	0.028*	3 (2–4)	2 (1–3)	0.001*
Not been able to work or study as well	2 (2–4)	2 (1–2)	0.004*	3 (2–4.5)	2 (1–2)	<0.0005*
Financial difficulties	2 (2–3)	2 (1–3)	0.013*	3 (2–4)	2 (1–3)	<0.0005*
Unmet support needs (CSNAT)						
CSNAT domains, n	4 (2–8)	2 (0–3)	<0.0005*	4.5 (2.3–8.8)	2 (0–4)	0.001*
Patient self-management						
CRQ mastery	4 (1.3)	4.5 (1.3)	0.047*	4.1 (1.5)	4.4 (1.3)	0.31
Carer comorbidities						
Physical comorbidities, ^a n	1 (1–2)	1 (0–2)	0.08	2 (1–3)	1 (0–2)	0.005*
Anxiety symptoms, n (%)	NA	NA	NA	23 (88.5)	29 (33.3)	<0.0005*
Depression symptoms, n (%)	23 (44.2)	3 (4.9)	<0.0005*	NA	NA	NA

Notes: ^aAny of allergies, arthritis, asthma, back pain, cancer, diabetes, epilepsy, cardiac disease, hypertension, renal disease, hepatic disease, or stroke. All values medians (IQR) unless otherwise stated. * $P \leq 0.05$.

Abbreviations: FEV₁, forced expiratory volume in 1 second; mMRC, modified Medical Research Council; FACQ, Family Appraisal of Caregiving Questionnaire; CSNAT, Carer Support Needs Assessment Tool; CRQ, Chronic Respiratory Questionnaire; NA, not applicable.

Patient depression was significantly associated with younger age, possibly due to greater difficulty adjusting to disease limitations.^{7,10} The link between exacerbations, dyspnea and anxiety may be explained by the dyspnea–anxiety–dyspnea

cycle¹¹ and cognitive model of panic,¹² a catastrophic misinterpretation of the sensation of dyspnea, possibly due to chemoreceptor hypersensitivity to CO₂ or processing of the affective dimension of dyspnea and anxiety in common

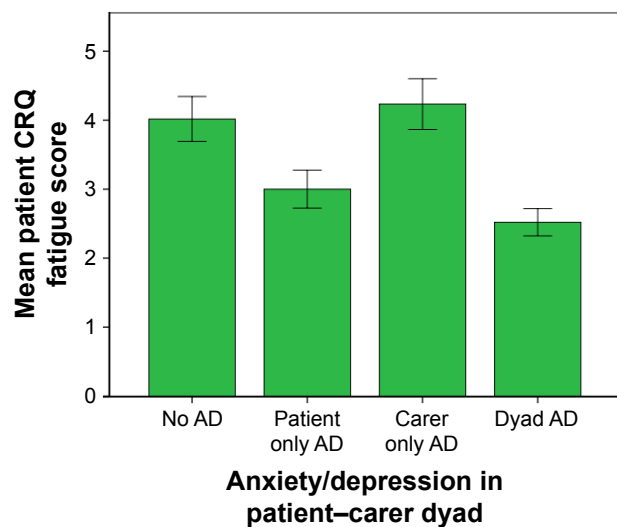


Figure 1 Dyad symptoms of anxiety or depression associated with patient fatigue. **Notes:** No AD (n=29), patient only AD (n=28), carer only AD (n=15), dyad AD (n=37). Data analyzed with one-way analysis of variance with Bonferroni post hoc test. Error bars represent 95% CIs. **Abbreviations:** AD, anxiety/depression; CRQ, Chronic Respiratory Questionnaire.

neuroanatomical regions.^{1,12} Interestingly, anxiety was only associated with exacerbations at home (not requiring hospital admission), which may support the hypothesis of overinterpretation of dyspnea⁹ or reflect higher anxiety from managing exacerbations in a nonclinical environment. There is evidence that systemic inflammation in exacerbations, with elevated interleukin-6 and C-reactive protein, leads to depressive symptoms.³³ Inversely, anxiety and depression can increase the risk of exacerbations, through hypothalamic–pituitary–adrenal axis activation suppressing immune function.⁹ Anxiety and depression were also associated with fatigue and more physical comorbidities, which may be because these lead to physical inactivity, social isolation, and thus depression. Psychological morbidities may impair the ability to acquire skills and lower confidence in self-management, leading to poor mastery.^{3,9} Finally, greater subjective symptom burden (CRQ scores) were significantly associated with anxiety and depression, but objective disease severity (% predicted FEV₁) was not.

Being female was strongly associated with carer anxiety, in concordance with previous studies.^{13,15,17,18} Women may use more emotion-based coping strategies, or view the caregiving role negatively as a continuation of their sex role.^{13,17} The number of unmet carer-support needs had a robust association with carer anxiety and depression; the link between anxiety and carers being separated/divorced/widowed may also be related to lack of social support for the carer. Carers with depression were younger and of higher educational level, which (as with patients) may be due to difficulty adjusting

to forced changes in lifestyle with the caring role, which would be greater for carers having their own physical comorbidities. Finally, anxiety and depression were significantly associated with greater subjective caring burden and anxiety with poor mastery by the patient (which could increase carer burden).

Importantly, we found that dyad anxiety or depression was significantly associated with greater patient fatigue. This may be explained by fatigue diminishing patient capacity to perform activities of daily living, thus increasing carer burden and reducing patients' sense of independence, leading to psychological morbidity in both. Conversely, psychological morbidity in either patient or carer may impact on capacity for patient activity, resulting in deconditioning and fatigue.

Within Rolland's family-system illness model, the associations found between younger age and both patient and carer depressive symptoms may reflect the conflict between this chronic disease having a centripetal effect on the family (greater focus on internal family life) and a younger patient or carer being part of a family in a centrifugal period (which is when individual family members are more engaged in the extrafamilial environment). Living with advanced COPD as a patient or carer may necessitate sacrifices in personal autonomy, which can have profound permanent effects on the family life cycle. Additionally, advanced COPD usually occurs in late adulthood, when having or caring for illness is more expected: younger patients and carers may be in a life-structure-building/maintenance period and may be forced back into an "out of phase" transition period that is disruptive. This is especially the case for diseases with a relapsing course, where the constant state of readiness required maintains the family in an out-of-phase state. Carers who were separated, divorced, or widowed were more likely to have symptoms of anxiety. This may be because separation represents a life-structure-changing period. When illness is concurrent with this, it can have a long-lasting impact on preparation for the next period of the life cycle. Finally, the link between patient mastery and patient and carer symptoms of anxiety and depression may be explained by mastery of illness being a key belief in the family's belief system, which may influence the family's account of illness and their strategies to adapt.

Clinical implications

As disease severity itself is not a barrier to psychological health, it is important to identify and address anxiety and depression in patients with advanced COPD. This study shows that it is feasible to use a simple tool, such as the

HADS, to screen for anxious and depressive symptoms in a clinical setting, which can indicate those who need more detailed assessment. Primary-care health professionals would be best placed to undertake this, to ensure coverage of the whole population of people living with advanced COPD. Clinicians should encourage all patients to participate in pulmonary rehabilitation, especially the exercise-training component, as this may represent the most effective way to interfere with the anxiety–dyspnea cycle.³⁴ Education for patients and carers on managing breathlessness³⁵ and exacerbations may reduce the anxiety associated with exacerbations at home. Pulmonary rehabilitation could also be an opportunity to increase physical activity and social interactions, which may help to prevent depression. It is important that although patients have advanced COPD, clinicians appropriately manage (and do not neglect) comorbid physical conditions and do not attribute all symptoms to COPD.

Given their high prevalence and associations, it may be appropriate to screen carers (particularly of COPD patients with anxiety and depression) for anxiety and depression. Particular attention must be given to carers who are female, younger, separated, divorced, or widowed, who have physical medical conditions themselves, who report a higher caring burden or more unmet support needs, and who have patients who struggle to manage their own COPD. Possible interventions include enabling carers to participate in care planning to improve understanding of COPD, education for carers on symptom control, and inclusion of carers in pulmonary rehabilitation programs, to reduce unmet support needs of the carer.^{35,36} Patients and carers may need access to respite care, and carers need time to attend to their own health (including their own medical conditions), work, and social life. Given the link between patient mastery and carer anxiety and depression, supporting patients in self-management of COPD could reduce psychological morbidity in carers.

Regarding the relationship between dyad anxiety and depression and patient fatigue, patients and carers could be supported to optimize their delegated dyadic coping: carers could offer balanced social support (so that they do not become overwhelmed themselves) and encourage patients to do as many everyday activities as they are able to before becoming too fatigued (to maintain a degree of self-efficacy).³⁷

Strengths and limitations

Important strengths of this study were the prospective design and recruitment of a population-based cohort sampled from primary care (which is most representative of the whole

population of patients living with advanced COPD). Patients typically receive shared care, with the majority of care being undertaken in primary care and the involvement of secondary care signifying either severe exacerbations or progressive symptoms (a distinct experience), which some with COPD do not experience. The patients in our cohort were well characterized in terms of their advanced COPD and sex-balanced. Research in COPD has historically focused on males, but females are increasingly affected by the condition, so are an important study group.

However the cross-sectional approach limited analyses to associations: longitudinal studies and analyses are needed to clarify the direction of causality and investigate changes in determinants of psychological health over time. The HADS is a self-report screening tool for symptoms of anxiety and depression, and not a formal physician diagnosis of anxiety or depression. Further, it asks about somatic symptoms that may equally be attributed to COPD. In addition, we did not control for any treatments for anxiety and depression that may have been received. Interviews with patients and carers were occasionally conducted together, where the physical layout of the home made separate interviews impossible. Previous studies have shown that combined and separate clinics with patients and carers in end-of-life care yield distinct findings. It has been shown that clinically, patients and carers value the time alone to voice their feelings and concerns, with patients expressing more cues that reveal emotion (surpassing the desire for information) and carers contributing more to conversation in private clinics.³⁸ Therefore, in joint interviews, it is possible patients and carers may have been more reluctant to express feelings of anxiety or depression. Nevertheless, the large majority of interviews were conducted separately, so we do not expect this to have significantly affected our results, but any effect would be an under- (rather than an over-) reporting of psychological morbidity in this population.

Conclusion

Symptoms of anxiety and depression of patients with advanced COPD and carers are significantly associated, and associated with younger age, female sex, being separated/divorced/widowed, higher educational level, poor physical health, greater disease and subjective caring burden, unmet carer-support needs, and poor patient self-efficacy. Dyad symptoms of anxiety or depression were associated with greater patient fatigue. Given their high prevalence, considerable impact on mortality, quality of life, health care service use, and their associations with each other, it is necessary

to screen for and address anxiety and depression in patients and carers.

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Author contributions

MF conceived and designed the study. GE, RM, SB, and HHB participated in conducting the study. MF and ACG acquired the data. Ella M and Emma M analyzed and interpreted the data. Ella M, Emma M, and MF drafted the manuscript. All authors contributed to revision of the manuscript for important intellectual content. All authors approve the final version to be published, and agree to be accountable for all aspects of the work. Ella M and Emma M are siblings.

Disclosure

The authors report no conflicts of interest in this work.

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