

## COPD exacerbations · 1: Epidemiology

G C Donaldson, J A Wedzicha

Thorax 2006;61:164–168. doi: 10.1136/thx.2005.041806

The epidemiology of exacerbations of chronic obstructive pulmonary disease (COPD) is reviewed with particular reference to the definition, frequency, time course, natural history and seasonality, and their relationship with decline in lung function, disease severity and mortality. The importance of distinguishing between recurrent and relapsed exacerbations is discussed.

observed that patients who report a smaller proportion of their exacerbations tend to have a poorer health related quality of life.<sup>8</sup> This difference could be explained by the fact that these patients do not receive treatment and consequently their exacerbations take longer to recover and have a greater impact on patients' perception of their disease.

Patients can experience a wide variety of symptoms during an exacerbation. It is generally agreed that the most distinct or cardinal symptoms of an exacerbation are increased dyspnoea, sputum purulence, and sputum volume.<sup>9</sup> Anthonisen *et al*<sup>9</sup> graded these exacerbations into type 1 (all three cardinal symptoms), type 2 (two cardinal symptoms), and type 3 (one cardinal symptom plus one of the following: an upper respiratory tract infection in the past 5 days, fever without other cause, increased wheezing or cough, or an increase in heart rate or respiratory rate by 20% compared with baseline readings). However, patients may also present with other symptoms such as worsening exercise tolerance, fluid retention, increased fatigue, acute confusion, or general malaise.<sup>10</sup> There are small falls in peak expiratory flow (PEF), forced expiratory volume in 1 second (FEV<sub>1</sub>), and forced vital capacity (FVC) at exacerbation, indicating narrowing of the airways through a variety of mechanisms.<sup>4</sup> Arterial blood gases and oxygen saturation may also worsen, and severe exacerbations may progress to respiratory failure.

The prevalence of chronic obstructive pulmonary disease (COPD) is increasing worldwide, as is tobacco usage.<sup>1</sup> Periods of acute worsening of this disease termed exacerbations—which greatly affect the quality of life and health of patients with COPD<sup>2</sup>—will therefore place a greater burden on the health services. Exacerbations can require physician consultation in primary and secondary care, may result in admission to hospital, and eventually lead to death. The epidemiology of these events is therefore important for both the understanding of the disease process and in assessing therapeutic outcomes.

## DEFINITION OF AN EXACERBATION

The precise definition of an exacerbation is a controversial topic. Exacerbations have been defined using respiratory symptoms alone (symptom-based), or symptoms plus an event such as the prescription of medication by a general practitioner or hospital admission for an acute exacerbation.<sup>3</sup> Both approaches have merits and disadvantages and, as yet, there is no consensus. To date, no sputum or plasma biomarker has been found that, with good sensitivity and specificity, can identify either the presence or severity of an exacerbation. This may be due to heterogeneity in the aetiology of exacerbations, delay between onset and presentation of the exacerbation for sampling of the biomarker, or shifts in the baseline (stable) levels with age or disease severity.

Exacerbation rates reported for symptom-based studies tend to be higher than for event-based studies because previous studies have shown that about 50% of exacerbations will not be reported to the research team.<sup>4–5</sup> This is to be expected as patients with COPD may not understand their disease and the importance of seeking treatment,<sup>6–8</sup> they may be depressed, or lack mobility. It is not certain whether unreported exacerbations are sufficiently mild that they can be dismissed. Little is known about the nature of these particular exacerbations as, by definition, unreported exacerbations are not seen by healthcare professionals. However, we have

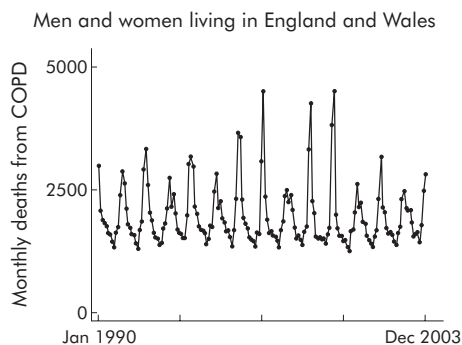
## MORTALITY

Figure 1 shows the monthly deaths from COPD in England and Wales for 1990–2003. It is projected that COPD will be the third leading cause of death worldwide in 2020.<sup>11</sup> There can be a number of causes recorded on death certificates due to the respiratory manifestations of COPD. Using the 9th International Classification of Disease (ICD) the codes were 490–492 and 496, and in the 10th ICD they are J40–J44. Most of these causes of death do not refer specifically to exacerbation. However, it is likely that patients at the terminal stage of COPD fulfil any definition of an exacerbation. Other exacerbation-related deaths may be attributed to related conditions (co-morbidities) such as arrhythmias, pulmonary embolism, or ischaemic heart disease. The use of death registration data may therefore not give the true picture on mortality due to exacerbation.

**Abbreviations:** BMI, body mass index; COPD, chronic obstructive pulmonary disease; FEV<sub>1</sub>, forced expiratory volume in 1 second; FVC, forced vital capacity; LRTI, lower respiratory tract infections; PEF, peak expiratory flow

See end of article for authors' affiliations

Correspondence to: Professor J A Wedzicha, Royal Free & University College Medical School, Hampstead Campus, London NW3 2PF, UK; j.a.wedzicha@medsch.ucl.ac.uk



**Figure 1** Monthly deaths from COPD (ICD9 490–492 and 496 for years before 2001 and ICD10 J40–J44 thereafter) at any age for men and women living in England and Wales. To be compatible with deaths in 1990–1992, data for the years from 1993 were divided by 0.920 to adjust for differences in coding instructions; deaths for 2001–2003 were then divided by 0.966 to allow for differences between ICD9 and ICD10. Data from the UK Office of National Statistics.

A number of studies have examined inpatient mortality from COPD. Estimates of inpatient mortality from exacerbation range from 4% to 30%. The wide variation in these estimates results from the fact that studies investigated different subgroups of patients.<sup>12</sup> Estimates for patients with COPD without respiratory failure vary from 5% to 11%, whereas COPD patients admitted in respiratory failure experience a higher rate (11–26%). Higher percentages are seen in those admitted to specialist respiratory or higher dependency wards (4–14%) or intensive care units (8–26%).<sup>12</sup> These differences undoubtedly reflect the admission of the more severely ill patients to these facilities. There are also variations attributable to the type of hospital in which the patient received care. Mortality appears to be higher in small district general hospitals (DGHs) (17.5%) than in teaching hospitals (11.9%) and large DGHs (11.2%), even after allowance for confounding factors.<sup>13</sup> Other factors predictive of higher inpatient mortality are fewer doctors (odds ratio (OR) 1.5) and fewer specialist physicians (OR 1.8).<sup>13</sup>

Soler-Cataluña *et al*<sup>14</sup> have recently shown that mortality is significantly related to the frequency of severe exacerbations requiring hospital care, independently of age, Charlson comorbidity index, body mass index (BMI), FEV<sub>1</sub> % predicted, or long term oxygen therapy in a cohort of 304 men studied over 5 years.

There are also data on mortality following discharge from hospital after treatment for an acute exacerbation. In the UK it has been reported that death occurs in 14% of cases (184/1342) within 3 months of admission.<sup>15</sup> Prognostic indicators of death during this follow up period were a low arterial pH, bilateral leg oedema, residence in a nursing home, age of 70 years or more, unrecordable PEF, oxygen saturation <86%, and assisted ventilation. In the USA, mortality rates following admission were 11% during hospitalisation, 20% after 2 months, 33% after 6 months, and 43% after 12 months.<sup>16</sup> In these patients survival was related to severity of illness, BMI, age, prior functional status, PaO<sub>2</sub>/Fio<sub>2</sub>, congestive heart failure, serum albumin, and the presence of cor pulmonale. In the Netherlands mortality during hospital stay was 8%, increasing to 23% after 1 year of follow up. Predictive factors were age, sex, FEV<sub>1</sub>, PaO<sub>2</sub>, Paco<sub>2</sub>, BMI, long term use of oral corticosteroids, a high co-morbidity index, and prior hospital admissions.<sup>17</sup>

There is an important caveat to all these data—they refer to deaths of patients whose exacerbation or underlying disease is sufficiently severe or life threatening to warrant admission to hospital. Most exacerbations are treated in the community and do not result in death. The statistics for exacerbation

related mortality should therefore always be qualified by the disease stage.

## ADMISSION TO HOSPITAL

One indication of the burden imposed by COPD exacerbations (defined here as ICD10 code J44) is that they were responsible for more than 0.9% of all 11.7 million hospital admissions and 2.4% of the 4.2 million acute medical admissions in England for 2003/2004. To place this in context, the equivalent figures for asthma (ICD10 code J45) are 0.56% and 1.3%, respectively.

There has been a steady rise since the early 1990s in the number of admissions to hospital for COPD. This is despite the fact that relatively few patients with an exacerbation are admitted, with most being treated on an outpatient basis. Table 1 shows recent data from the hospital episode statistics published by the Department of Health for the financial years from 1998 to 2003. The number of admissions for COPD has risen by 13.1% over this period, with the proportion of men admitted rising from 72% to 79%. Most of these admissions are on an emergency basis. Despite these increasing pressures on the health service, the mean length of stay has remained almost unchanged at about 10 days.

## PRIMARY CARE CONSULTATIONS

A number of studies have examined the impact of exacerbations on primary care consultations. Miravittles *et al*<sup>18</sup> reported data on 1001 patients seen in 201 Spanish practices. The median number of exacerbations seen in primary care was two per year, with 309 patients (31%) suffering three or more exacerbations in 1 year.

McGuire *et al*<sup>19</sup> estimated the number of GP consultations for chronic bronchitis and acute exacerbations in excess of the number of visits by the general population. They relied mainly on the General Practice Research Database (1994–5) in which the diagnosis and long term treatment of a chronic condition is recorded only once. These consultation data therefore refer to exacerbations rather than the stable disease. They reported an average of 4.01 GP visits per year in the 45–54 age range, 4.51 visits per year in the 55–64 age range, 3.6 per year in the 65–74 age range, 1.46 visits per year in the 75–84 age range, and –0.30 visits per year in those aged over 85 years. The decline in the number of visits with age probably reflects increased consultations for other causes in people without chronic bronchitis rather than a reduction in the number of consultations for acute exacerbations.

## EXACERBATION FREQUENCY

There is a subgroup of patients with COPD who experience an above normal number of exacerbations. A figure of 2.5 or 3 exacerbations per year is normal with a symptomatic definition of an exacerbation. These patients have a poorer quality of life,<sup>2</sup> higher inflammatory markers when stable,<sup>20</sup> and a faster decline in lung function (see below). They are more likely to become housebound,<sup>21</sup> to experience more colds,<sup>22</sup> and to be chronic sputum producers.<sup>23</sup> They are prone to readmission to hospital<sup>15–24</sup> and frequent severe exacerbations are associated with increased mortality.<sup>14</sup> This propensity towards frequent exacerbation appears for the most part to be independent of disease severity. The poor health and prognosis of this subgroup of patients suggests that they warrant targeting with further therapeutic interventions.

## EXACERBATIONS AND DECLINE IN FEV<sub>1</sub>

The seminal work of Fletcher and colleagues<sup>25</sup> on 792 smokers and non-smokers, with and without respiratory disease, suggested that lower respiratory tract infections played no role in the annual decline in FEV<sub>1</sub>. This conclusion was supported by Howard *et al*<sup>26</sup> who followed a similar group

**Table 1** Trends in hospital episode statistics (HES) from 1998/9 to 2003/4 for ICD10 code J44 (COPD with acute lower respiratory tract infection/exacerbation/unspecified)

| HES year (starting April) | Finished consultant episodes | Admissions | Men   | Emergency | Mean length of stay (days) | Elective | Bed days* |
|---------------------------|------------------------------|------------|-------|-----------|----------------------------|----------|-----------|
| 2003                      | 163480                       | 105600     | 83036 | 99590     | 10.0                       | 1683     | 1046371   |
| 2002                      | 140541                       | 93373      | 71848 | 87942     | 10.3                       | 1283     | 944835    |
| 2001                      | 135252                       | 92049      | 69968 | 86999     | 10.4                       | 1263     | 941216    |
| 2000                      | 127116                       | 89730      | 67231 | 84721     | 10.4                       | 1285     | 918174    |
| 1999                      | 129501                       | 95039      | 68487 | 89692     | 10.2                       | 1132     | 953393    |
| 1998                      | 125047                       | 93019      | 67263 | 87549     | 10.4                       | 1108     | 952495    |

\*Methodology changes make the number of bed days in the year starting 2003 inconsistent with previous years.

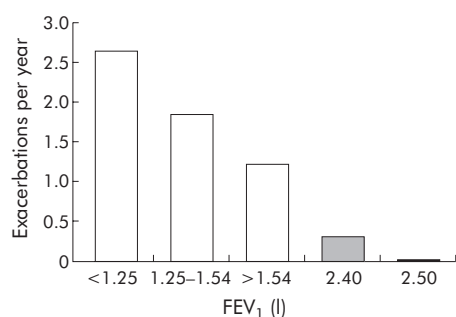
for 11 years and by Bates *et al*<sup>27</sup> in a 12 year study of male war veterans. However, indirect evidence that exacerbations contributed to the decline in FEV<sub>1</sub> continued to be reported even after these publications. Exacerbations were found to be more frequent in those with regular sputum production, and regular sputum production was reported to be associated with an accelerated decline in FEV<sub>1</sub><sup>28</sup> and with death.<sup>29</sup> Data from our group<sup>4</sup> also showed that PEF did not recover to pre-exacerbation levels within 91 days in 7.1% of exacerbations. This indicated that exacerbations could produce a permanent reduction in lung function, which suggested that patients who experienced more exacerbations would have a faster decline than those with fewer exacerbations.

Three studies have now shown that exacerbations play a role in the decline in FEV<sub>1</sub>. The Lung Health Study,<sup>30</sup> a 5 year prospective randomised multicentre study of smoking cessation, monitored 5887 smokers aged 35–60 years. Spirometric parameters and the number of lower respiratory tract infections resulting in a physician visit (LRTI) were recorded annually. The analysis showed no relationship between LRTI and FEV<sub>1</sub> decline in patients who stopped smoking completely but, in those who smoked intermittently or continuously, FEV<sub>1</sub> declined by 7.3 ml/year ( $p < 0.05$ ) and 7.1 ml/year ( $p < 0.001$ ), respectively, for each additional LRTI per year. Donaldson *et al*<sup>31</sup> examined 109 patients of median age 68.1 years with moderate to very severe COPD over a 4 year period. An additional decline in FEV<sub>1</sub> of 8.0 ml/year and in PEF of 2.22 l/min/year was seen in COPD patients who experienced frequent exacerbations ( $\geq 2.92$  exacerbations per

year) compared with those who had fewer exacerbations. Dowson *et al*,<sup>32</sup> investigating patients with emphysema due to  $\alpha_1$ -antitrypsin deficiency, found a positive correlation between exacerbation frequency and decline in FVC for the group as a whole ( $n = 43$ ) and in FEV<sub>1</sub> for 26 patients with a baseline post-bronchodilator FEV<sub>1</sub>  $> 35\%$  predicted.

### EXACERBATIONS AND DISEASE SEVERITY

There seems to be a general consensus that exacerbation frequency increases with disease severity. Figure 2 shows the number of exacerbations per year in relation to FEV<sub>1</sub>. Here, exacerbations were identified using an event-based definition and data were combined from three studies (the ISOLDE Study, the European Respiratory Society Study on COPD, and the Copenhagen City Lung Study).<sup>33</sup> Using a symptom-based definition, we reported that patients with severe COPD (GOLD category III,  $n = 38$ ) had an annual exacerbation frequency of 3.43 per year compared with 2.68 per year in those with moderate COPD (GOLD II,  $n = 94$ ;  $p = 0.029$ ).<sup>34</sup> Greenberg *et al*<sup>35</sup> found that respiratory illnesses—defined as increased cough, shortness of breath, or sputum production and/or a change in sputum colour—were more frequent in patients with moderate COPD (3 per year) than in those with mild COPD (1.8 per year). Miravittles *et al*<sup>18</sup> also reported that exacerbations, using a symptom-based definition, increased with disease severity. With an FEV<sub>1</sub> of  $> 60\%$  predicted the mean (SD) number of exacerbations per year was 1.6 (1.5), compared with 1.9 (1.8) with FEV<sub>1</sub> 59–40% and 2.3 (1.9) with FEV<sub>1</sub>  $> 40\%$ .



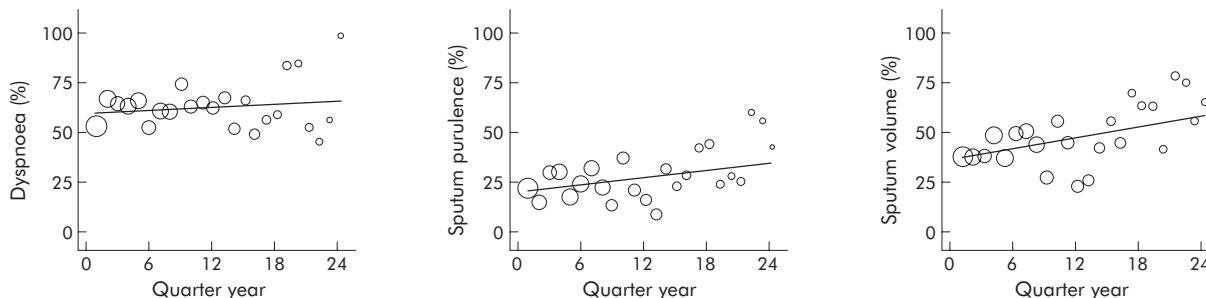
**Figure 2** Mean incidence of exacerbations per year in relation to forced expiratory volume in 1 second (FEV<sub>1</sub>) in the placebo arms of three 3 year studies. From left to right, the first three bars are from the Inhaled Steroids in Obstructive Lung Disease in Europe (ISOLDE) study in which exacerbations were defined as a worsening of respiratory symptoms that required treatment with oral corticosteroids or antibiotics, as judged by the general practitioner. The next bar represents data from the Copenhagen City Lung Study which defined exacerbations more broadly as an affirmative answer to the question: "Have you, since your last visit, experienced more cough and phlegm?" This probably explains the slightly higher exacerbation rate than expected from the FEV<sub>1</sub> values. The final bar on the right shows data from the European Society Study on COPD which used the same definition as the ISOLDE study. Reproduced with permission from Burge and Wedzicha.<sup>33</sup>

### TIME COURSE OF EXACERBATIONS

A number of variables have been monitored over time to investigate exacerbation severity and recovery. In our study of 504 exacerbations experienced by 91 patients,<sup>4</sup> respiratory symptoms of dyspnoea, sore throat, cough, and nasal congestion/discharge (cold) tended to worsen during the prodromal period 7 days before the onset of exacerbation ( $p < 0.05$ ). On the day of onset, symptoms increased sharply with 64% of exacerbations associated with symptoms of dyspnoea, 26% with increased sputum volume, 42% with sputum purulence, 35% with colds, 35% with wheeze, 12% with sore throat, and 20% with cough. The changes in lung function associated with these exacerbations were small: PEF fell by a median of 8.6 l/min, FEV<sub>1</sub> by 24 ml, and FVC by 76 ml. Recovery in PEF took a median of 6 days (IQR 1–14) and daily total symptom count took 7 days (IQR 4–14) to recover. Recovery of PEF was achieved in only 75.2% of exacerbations within 35 days, and 7.1% of exacerbations had still not returned to baseline after 91 days. The pattern of acute worsening at exacerbation and slow recovery was also seen in the improvements in sputum colour, neutrophil count, and isolation of bacterial species and bacterial load reported by Stockley *et al*.<sup>36</sup>

A similar pattern in recovery, but of a longer duration of up to several months, was seen in the time course of health





**Figure 3** Changes over time in the percentage of exacerbations involving increased dyspnoea, sputum volume, or sputum purulence on the day of onset. The x axis is time from the date patients enrolled in the study, thus standardising for differences in the time when the patients started and ended the study. Circle size indicates the number of exacerbations recorded in that quarter year. Adapted from Donaldson *et al*<sup>14</sup> with permission of European Respiratory Society Journals Ltd.

status. Data from the GLOBE study, a randomised double blind trial of antibiotic therapy, showed that health related quality of life assessed by the total score from the St George’s Respiratory Questionnaire fell (that is, patients felt better) by 11.8 units at 4 weeks and by a further 4.6 units at 12 weeks after an exacerbation.<sup>37</sup> More acute changes have been observed in the time spent outside the home during an exacerbation.<sup>21</sup> Typically, 32–35% of all patients stayed indoors on any day when stable, but this rose to 45% at exacerbation onset, peaked at 48% on the day after onset, and returned to baseline levels by 14 days. The longer duration of worsening health status compared with lung function suggests that exacerbations have a lasting impact. This could be anticipated as episodes of severe, possibly life threatening, dyspnoea would be difficult to overlook.

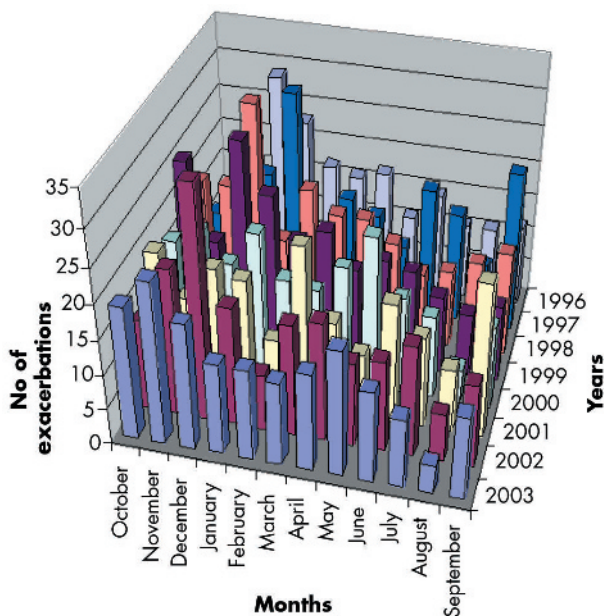
**NATURAL HISTORY OF EXACERBATIONS**

Little work has been done on how the severity and symptom composition of exacerbations changes over time. In our cohort study<sup>34</sup> we asked patients to record any worsening in their normal stable condition. Patients therefore recorded increased dyspnoea or onset of a new symptom such as a sore

throat. We found that the number of symptoms above normal at exacerbation increased by 0.05 per year ( $p = 0.047$ ), and that the number of days it took to recover from an exacerbation in terms of symptoms and FEV<sub>1</sub> took longer in successive years (0.32 and 0.55 days per year; both  $p < 0.05$ ). We also found that sputum purulence at exacerbation became more common over time, increasing by 4.1% per year (fig 3).

**SEASONALITY**

Exacerbations are about 50% more likely in winter (fig 4). This explains some of the increased pressures on hospital beds during the winter months. In an interesting study in Spain,<sup>5</sup> 441 (432 men) with a diagnosis of COPD were monitored for 2 years at 6 monthly intervals. The number of exacerbations for which the patients made an unscheduled visit to the investigator (reported exacerbations) were recorded, as were the number of exacerbations reported at follow up visits (unreported exacerbations), to give the total number of exacerbations. Of a total of 1015 exacerbations, 209 reported exacerbations occurred in the summer and 248 in winter, while 193 unreported exacerbations took place in the summer and 365 in winter. Unreported exacerbations were therefore more likely in winter (59.5%) than in summer (47.9%;  $p < 0.0001$ ). We reported that exacerbations were associated with colder outdoor temperature ( $p < 0.005$ ).<sup>38</sup> Increased exacerbations during winter are probably the result of improved survival of respiratory viruses in colder weather, the crowding together of people indoors during the winter, and possibly a reduced immunological response.<sup>39</sup>



**Figure 4** Number of exacerbations experienced by patients in the East London COPD study by calendar month and years 1996–2003. More exacerbations were seen in the winter months (November to February) than in the summer. Data from our COPD cohort.

**RECURRENT EXACERBATIONS**

It is now recognised that exacerbation frequency is an important outcome measure in COPD. To advance our understanding of frequent exacerbations we need to classify exacerbations further. It has been reported that approximately 30% of patients seen at hospital with an exacerbation will be seen again and possibly admitted with another exacerbation within 8 weeks.<sup>40–41</sup> The term “recurrent” has also been used to describe repeat admissions for acute hypercapnic respiratory failure.<sup>42</sup> It would be helpful to use this term to describe exacerbations that are new events, following close on from an earlier index exacerbation, but with an intervening period of usual (relatively good) health. Recurrent exacerbations would be distinct from relapsed exacerbations where, during the course of the exacerbation, the patient’s health worsens and an additional intervention is required. At no time during the relapsed exacerbation does the patient recover to his/her normal health. Further research is required to study both relapsed and recurrent exacerbations.

## CONCLUSIONS

Exacerbations are now recognised as an important outcome measure in COPD. In the last 10 years frequent exacerbations have been shown to be associated with poorer health outcomes and a greater burden on the health service. The future challenge lies in understanding the causes of frequent exacerbations, differences in exacerbation severity, and the failure of some exacerbations to resolve.

## Authors' affiliations

G C Donaldson, J A Wedzicha, Academic Unit of Respiratory Medicine, University College London, London, UK

Competing interests: JAW is Editor in Chief of *Thorax*.

## REFERENCES

- Lopez AD, Murray CC. The global burden of disease, 1990–2020. *Nat Med* 1998;**4**:1241–3.
- Seemungal TAR, Donaldson GC, Paul EA, et al. Effect of exacerbation on quality of life in patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 1998;**157**:1418–22.
- Pauwels R, Calverley P, Buist AS, et al. COPD exacerbations: the importance of a standard definition. *Respir Med* 2004;**98**:99–107.
- Seemungal TAR, Donaldson GC, Bhowmik A, et al. Time course and recovery of exacerbations in patients with chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2000;**161**:1608–13.
- Miravittles M, Ferrer M, Pont A, et al. Effect of exacerbations on quality of life in patients with chronic obstructive pulmonary disease: a 2 year follow up study. *Thorax* 2004;**59**:387–95.
- Rennard S, Decramer M, Calverley PMA, et al. Impact of COPD in North America and Europe in 2000: subjects' perspective of confronting COPD international survey. *Eur Respir J* 2002;**20**:799–805.
- Okubadejo AA, Jones PW, Wedzicha JA. Quality of life in patients with chronic obstructive pulmonary disease and severe hypoxaemia. *Thorax* 1996;**51**:44–7.
- Wilkinson TMA, Donaldson GC, Hurst JR, et al. Early therapy improves outcomes of exacerbations of chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2004;**169**:1298–303.
- Anthonisen NR, Manfreda J, Warren CP, et al. Antibiotic therapy in exacerbations of chronic obstructive pulmonary disease. *Ann Intern Med* 1987;**106**:196–204.
- National Institute for Clinical Excellence (NICE). Management of exacerbations of COPD. In: Chronic obstructive pulmonary disease: national clinical guideline on management of chronic obstructive pulmonary disease in adults in primary and secondary care, *Thorax* 2004;**59**(Suppl I):131–56.
- Murray CJ, Lopez AD. Alternative projections of mortality and disability by cause 1990–2020: Global Burden of Disease Study. *Lancet* 1997;**349**:1498–504.
- Patil SP, Krishnan JA, Lechtzin N, et al. In-hospital mortality following acute exacerbations of chronic obstructive pulmonary disease. *Arch Intern Med* 2003;**163**:1180–6.
- Roberts CM, Barnes S, Lowe D, et al. Evidence for a link between mortality in acute COPD and hospital type and resources. *Thorax* 2003;**58**:947–9.
- Soler-Cataluña JJ, Martínez-García MÁ, Román Sánchez P, et al. Severe acute exacerbations and mortality in patients with chronic obstructive pulmonary disease. *Thorax* 2005;**60**:925–31.
- Roberts CM, Lowe D, Bucknall CE, et al. Clinical audit indicators of outcome following admission to hospital with acute exacerbation of chronic obstructive pulmonary disease. *Thorax* 2002;**57**:137–41.
- Connors AF, Dawson NV, Thomas C, et al. Outcomes following acute exacerbation of severe chronic obstructive lung disease. *Am J Respir Crit Care Med* 1996;**154**:959–67.
- Groenewegen KH, Schols AM, Wouters EF. Mortality and mortality-related factors after hospitalization for acute exacerbation of COPD. *Chest* 2003;**124**:459–67.
- Miravittles M, Mayordomo C, Artés M, et al. Treatment of chronic obstructive pulmonary disease and its exacerbations in general practice. *Respir Med* 1999;**93**:173–9.
- McGuire A, Irwin DE, Fenn P, et al. The excess cost of acute exacerbations of chronic bronchitis in patients aged 45 and older in England and Wales. *Value Health* 2001;**4**:370–5.
- Bhowmik A, Seemungal TA, Sapsford RJ, et al. Relation of sputum inflammatory markers to symptoms and lung function changes in COPD exacerbations. *Thorax* 2000;**55**:114–20.
- Donaldson GC, Wilkinson TM, Hurst JR, et al. Exacerbations and time spent outdoors in chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2005;**171**:446–52.
- Hurst JR, Donaldson GC, Wilkinson TMA, et al. Epidemiological relationships between the common cold and exacerbation frequency in COPD. *Eur Respir J* 2005;**26**:846–52.
- Powrie DJ, Wilkinson TM, Donaldson GC, et al. Chronic sputum production in COPD is associated with increased airway inflammation and exacerbation frequency. *American Thoracic Society Meeting*. 2005;140: F88 (abstract).
- García-Aymerich J, Monso E, Marrades RM, et al. Risk factors for hospitalization for a chronic obstructive pulmonary disease exacerbation: EFRAM study. *Am J Respir Crit Care Med* 2001;**164**:1002–7.
- Fletcher C, Peto R, Tinker C, et al. *The natural history of chronic bronchitis and emphysema*. Oxford: Oxford University Press, 1976.
- Howard P. A long-term follow-up of respiratory symptoms and ventilatory function in a group of working men. *Br J Ind Med* 1970;**27**:326–33.
- Bates D. The fate of the chronic bronchitic: a report of the 10-year follow-up in the Canadian Department of Veteran's Affairs coordinated study of chronic bronchitis. *Am Rev Respir Dis* 1973;**108**:1043–65.
- Vestbo J, Prescott E, Lange P. Association of chronic mucus hypersecretion with FEV<sub>1</sub> decline and chronic obstructive pulmonary disease morbidity. Copenhagen City Heart Study Group. *Am J Respir Crit Care Med* 1996;**153**:1530–5.
- Prescott E, Lange P, Vestbo J. Chronic mucus hypersecretion in COPD and death from pulmonary infection. *Eur Respir J* 1995;**8**:1333–8.
- Kanner RE, Anthonisen NR, Connett JE. Lower respiratory illnesses promote FEV<sub>1</sub> decline in current smokers but not ex-smokers with mild chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2001;**164**:358–64.
- Donaldson GC, Seemungal TAR, Bhowmik A, et al. Relationship between exacerbation frequency and lung function decline in chronic obstructive pulmonary disease. *Thorax* 2002;**57**:847–52.
- Dowson LJ, Guest PJ, Stockley RA. Longitudinal changes in physiological, radiological, and health status measurements in alpha(1)-antitrypsin deficiency and factors associated with decline. *Am J Respir Crit Care Med* 2001;**164**:1805–9.
- Burge S, Wedzicha JA. COPD exacerbations: definitions and classifications. *Eur Respir J* 2003;**21**(Suppl 41):46–53.
- Donaldson GC, Seemungal TA, Patel IS, et al. Longitudinal changes in the nature, severity and frequency of COPD exacerbations. *Eur Respir J* 2003;**22**:931–6.
- Greenberg SB, Allen MA, Wilson J, et al. Respiratory viral infections in adults with and without chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2000;**162**:167–73.
- Stockley RA, O'Brien C, Pye A, et al. Relationship of sputum color to nature and outpatient management of acute exacerbations of COPD. *Chest* 2000;**117**:1638–45.
- Spencer S, Jones PW, GLOBE Study Group. Time course of recovery of health status following an infective exacerbation of chronic bronchitis. *Thorax* 2003;**58**:589–93.
- Donaldson GC, Seemungal T, Jeffries DJ, et al. Effect of temperature on lung function and symptoms in chronic obstructive pulmonary disease. *Eur Respir J* 1999;**13**:844–9.
- Eccles R. An explanation for the seasonality of acute upper respiratory tract viral infections. *Acta Otolaryngol* 2002;**122**:183–91.
- Sethi S, File TM. Managing patients with recurrent acute exacerbations of chronic bronchitis: a common clinical problem. *Curr Med Res Opin* 2004;**20**:1511–21.
- Skwarska E, Cohen G, Skwarski KM, et al. Randomised controlled trial of supported discharge in patients with exacerbations of chronic obstructive pulmonary disease. *Thorax* 2000;**55**:907–12.
- Chu CM, Chan VL, Lin Awn, et al. Readmission rates and life threatening events in COPD survivors treated with non-invasive ventilation for acute hypercapnic respiratory failure. *Thorax* 2004;**59**:1020–5.