



## Impact of the UK lockdown on people at risk of COPD

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*To the Editor:*

On 23 March 2020, the UK Government announced a nationwide lockdown in response to the COVID-19 epidemic, with people banned from leaving their homes except for essential journeys and exercise.

From the beginning, public health advice issued by the UK National Health Service (NHS) encouraged smokers to quit. This advice was based on studies that showed a five times greater risk infection of the respiratory tract by the influenza virus in smokers and that bacterial infection/pneumonia is twice as likely in those who smoke [1, 2]. Early data during the pandemic were inconclusive as to whether tobacco smoking or nicotine conferred increased susceptibility to COVID-19 infection or not, but infected smokers had poorer clinical outcomes [3].

Whether people living in the UK took the public health advice is unclear as mass survey data are contradictory. One early cross-sectional survey suggested that 12.2% of those who tried to quit in the 3 months prior to March 2020 were motivated by COVID-19 but this was not statistically tested [4]. Data from another survey during April 2020 showed no significant change in smoking behaviour, with 15.9% of people sampled smoking in the year prior to February 2020 compared to 17.0% of people surveyed in April 2020 [5], whilst another survey suggested that smokers were stress-smoking more since the pandemic [6]. Our aim was to describe smoking and other behaviours during a nationwide lockdown.

Participants enrolled in the British Lung Foundation Early COPD cohort were aged 30–45 years, current tobacco smokers with >10-pack-year smoking history, forced expiratory volume in 1 s (FEV<sub>1</sub>) ≥80% predicted and a body mass index <35 kg·m<sup>-2</sup>. They were scheduled to be seen face-to-face in clinic every 6 months for measurement of full lung function, completion of various questionnaires, and physical collection of blood and sputum samples, but during lockdown, telephone interviews were performed and written questionnaires returned as photographs attached to e-mails. Remote interviews took place between 16 April and 28 September 2020, and paired with the closest pre-pandemic visit.

By telephone, participants were asked about smoking habits, upper respiratory tract infection (URTIs) and lower respiratory tract infections (LRTIs). They were also sent the COPD Assessment Test (CAT), Leicester Cough Questionnaire, Hospital Anxiety and Depression Scale questionnaire, and questions 1 and 2 of the St George's Respiratory Questionnaire (SGRQ) for diagnoses of chronic bronchitis [7, 8]. 27 participants did not return the questionnaires. Statistical comparisons were made by paired t-test or Wilcoxon matched pairs test; comparisons between groups were made by Mann–Whitney and Chi-squared tests as appropriate. Remote telephone visits were approved by a nonsubstantial amendment to the ethics approval (London-Riverside Research Ethics Committee, 16LO2041).

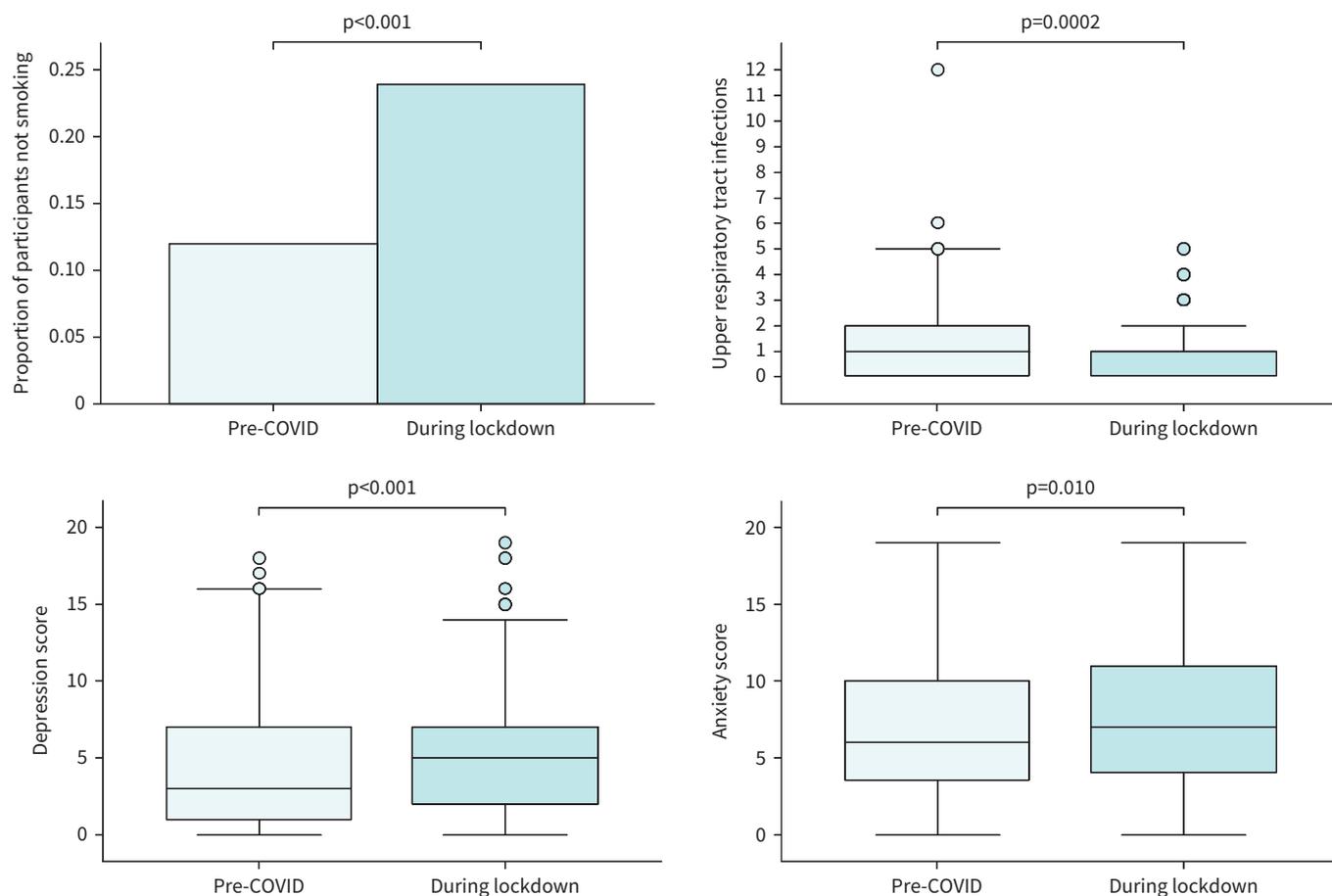
Data were obtained during lockdown between the 16 April and 25 September 2020 from 260 individuals (25 participants in Belfast, 38 in Birmingham, 25 in Edinburgh, 101 in London, 27 in Manchester and 44 in Nottingham). The mean±SD age was 38.1±4.6 years (n=259), 155 (59.6%) were male and the post-bronchodilator FEV<sub>1</sub> at enrolment was 3.81±0.8 L (101±11% of Global Lung Function Initiative predicted). The interval between the two visits was 196 days (interquartile range (IQR) 80–238 days).

Figure 1 demonstrates that during lockdown, the proportion of participants recorded as having stopped smoking for at least a week doubled from 31 (12.0%) out of 259 prior to lockdown to 62 (23.9%) out of 259 (Chi-squared  $p<0.001$ ) during lockdown. Smoking habit data were available at both visits for 258 participants; 40 had stopped, nine restarted and 209 participant's habits were unchanged. Of these 209,



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**FIGURE 1** The proportion of nonsmoking participants, frequency of upper respiratory tract infections, and depression and anxiety scores before and during lockdown in the UK during the COVID-19 pandemic.

187 continued to smoke and 22 had already quit. In those who smoked before and during lockdown, assessed over the 7 days preceding each visit, tobacco consumption (cigarettes per day; Wilcoxon  $p = 0.587$ ) and rolling tobacco (grams per week; Wilcoxon  $p = 0.924$ ) was unchanged.

During lockdown, the anxiety component scores increased from  $6.7 \pm 4.4$  to  $7.4 \pm 4.7$  ( $n = 233$ ; t-test  $p = 0.010$ ) and depression scores increased from  $4.30 \pm 3.9$  to  $5.14 \pm 4.1$  ( $n = 233$ ; t-test  $p < 0.001$ ). Anxiety increased in 78 out of 233 and depression in 89 out of 233 participants by  $\geq 2$  units, the minimally important difference in COPD [9].

Respiratory symptoms improved during lockdown, with CAT scores falling from 9 (IQR 6–15) to 8 (IQR 5–13) ( $n = 233$ ; Wilcoxon  $p = 0.008$ ) and the total Leicester Cough Questionnaire score indicating improvement, as it increased from 19.7 (IQR 17.6–20.6) to 20.0 (IQR 18.7–20.8) ( $n = 204$ ; Wilcoxon  $p = 0.045$ ).

A comparison of the 40 patients who quit during lockdown with those 187 who smoked at both visits showed no difference prior to lockdown in CAT scores (Mann–Whitney  $p = 0.814$ ) or in the prevalence of chronic bronchitis defined by the SGRQ (Chi-squared  $p = 0.607$ ). However, after quitting, CAT scores were lower (5 (IQR 3–13) compared with 9 (IQR 6–13)) in smokers (Mann–Whitney  $p = 0.0307$ ) and chronic bronchitis less prevalent (four (11.8%) out of 34) than in those who continued to smoke (47 (28.1%) out of 167) (Chi-squared  $p = 0.045$ ).

The number of URTIs recalled by the participants fell from a median of 1 (IQR 0–2; group mean 1.24) when asked pre-pandemic to a median 1 (IQR 0–1; group mean 0.84) (Wilcoxon  $p = 0.0002$ ) when asked during lockdown. There was no difference in the number of LRTIs (Wilcoxon  $p = 0.192$ ).

In summary, during the UK national lockdown and a period of enforced social distancing, there was a significant increase in the proportion of 30–45-year-old smokers who quit smoking despite increased levels of depression and anxiety. It has been reported that depressed individuals are more likely to be nicotine dependent but paradoxically, older depressed adults coupled with health concerns are more motivated to quit [10]. Stress may have enhanced lockdown as an important moment at which to quit. One limitation in our study is the absence of objective measures of nicotine use. However, participants who self-reported that they had quit smoking had a reduced prevalence of chronic bronchitis and lower CAT scores, which suggests they were truthful about their habits. There was also a reduced number of URTIs, again suggestive that participants had fewer social interactions and opportunities for cross-infection.

Our findings suggest that any evaluation of the health response to the pandemic needs to balance deterioration in mental health in specific age groups with improvements in respiratory health.

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## References

- 1 Arcavi L, Benowitz NL. Cigarette smoking and infection. *Arch Intern Med* 2004; 164: 2206–2216.
- 2 Stämpfli MR, Anderson GP. How cigarette smoke skews immune responses to promote infection, lung disease and cancer. *Nat Rev Immunol* 2009; 9: 377–384.
- 3 Vardavas CI, Nikitara K. Covid-19 and smoking: a systematic review of the evidence. *Tob Induc Dis* 2020; 18: 20.

- 4 Tattan-Birch H, Perski O, Jackson S, *et al.* Covid-19, smoking, vaping and quitting: a representative population survey in England. *Addiction* 2021; 116: 1186–1195.
- 5 Jackson SE, Garnett C, Shahab L, *et al.* Association of the COVID-19 lockdown with smoking, drinking and attempts to quit in England: an analysis of 2019–20 data. *Addiction* 2021; 116: 1233–1244.
- 6 Mintel Press Office. Panic puffs: half of smokers are stress-smoking more since the pandemic started. <https://www.mintel.com/press-centre/social-and-lifestyle/panic-puffs-half-of-smokers-are-stress-smoking-more-since-the-pandemic-started>. Date last updated: 15 April 2021.
- 7 Kim V, Zhao H, Regan E, *et al.* The St. George’s respiratory questionnaire definition of chronic bronchitis may be a better predictor of COPD exacerbations compared with the classic definition. *Chest* 2019; 156: 685–695.
- 8 Kim V, Crapo J, Zhao H, *et al.* Comparison between an alternative and the classic definition of chronic bronchitis in COPDgene. *Ann Am Thorac Soc* 2015; 12: 332–339.
- 9 Wynne SC, Patel S, Barker RE, *et al.* Anxiety and depression in bronchiectasis: response to pulmonary rehabilitation and minimal clinically important difference of the hospital anxiety and depression scale. *Chron Respir Dis* 2020; 17: 1479973120933292.
- 10 Sachs-Ericsson N, Schmidt NB, Zvolensky MJ, *et al.* Smoking cessation behavior in older adults by race and gender: The role of health problems and psychological distress. *Nicotine Tob Res* 2009; 11: 433–443.