Ventilatory function as a predictor of mortality in lifelong non-smokers: evidence from large British cohort studies

Ramyani P Gupta, David P Strachan

ABSTRACT

Background Reduced ventilatory function is an established predictor of all-cause mortality in general population cohorts. We sought to verify this in lifelong non-smokers, among whom confounding by active smoking can be excluded, and investigate associations with circulatory and cancer deaths.

Methods In UK Biobank, among 149,343 white never-smokers aged 40–69 years at entry, 2,401 deaths occurred over a mean of 6.5-year follow-up. In the Health Surveys for England (HSE) 1995, 1996, 2001 and Scottish Health Surveys (SHS) 1998 and 2003 combined, there were 500 deaths among 6579 white never-smokers aged 40–69 years at entry, followed for a mean of 13.9 years. SD (z) scores for forced expiratory volume in the first second (FEV1) and forced vital capacity (FVC) were derived using Global Lung Initiative 2012 reference equations. These z-scores were related to deaths from all causes, circulatory disease and cancers using proportional hazards models adjusted for age, sex, height, socioeconomic status, region and survey.

Results In the HSE–SHS data set, decreasing z-scores for FEV1 (zFEV1) and FVC (zFVC) were each associated to a similar degree with increased all-cause mortality (hazard ratios per unit decrement 1.17, 95% CI 1.09 to 1.25 for zFEV1 and 1.19, 95% CI 1.10 to 1.28 for zFVC). This was replicated in Biobank (HRs 1.21, 95% CI 1.17 to 1.26 and 1.24, 1.19 to 1.29, respectively). zFEV1 and zFVC were less strongly associated with mortality from circulatory diseases in HSE–SHS (HR 1.22, 95% CI 1.06 to 1.40 for zFVC) than in Biobank (HR 1.47, 95% CI 1.35 to 1.60 for zFVC). For cancer mortality, HRs were more consistent between cohorts (for zFVC: HRs 1.12, 95% CI 1.01 to 1.24 in HSE–SHS and 1.10, 1.05 to 1.15 in Biobank). The strongest associations were with respiratory mortality (for zFVC: HRs 1.61, 95% CI 1.25 to 2.08 in HSE–SHS and 2.15, 1.77 to 2.61 in Biobank).

Conclusions Spirometric indices predicted mortality more strongly than systolic blood pressure or body mass index, emphasising the importance of promoting lung health in the general population, even among lifelong non-smokers.

INTRODUCTION

Four decades of epidemiological research have consistently shown that reduced levels of ventilatory function, measured as one-second forced expiratory volume (FEV1) or forced vital capacity (FVC), are associated with higher all-cause mortality rates, and therefore shorter survival in the general population. Few studies have reported specifically on lifelong non-smokers, a group which form the minority of most populations surveyed hitherto, but are set to become more common in future as smoking becomes less prevalent in higher income countries.

Most publications have focused on FEV1, but a recent analysis of asymptomatic participants in the multietnic Atherosclerosis Risk in Communities (ARIC) Study reported that all-cause mortality was strongly associated
with diminished FVC, after adjustment for FEV1, but not the other way around, and there was no association between survival and the ratio of FEV1 to FVC. This conclusion was based on a combined analysis of smokers and non-smokers.

In this report, we compare the relationship of lung function measures (FEV1 and FVC) with subsequent mortality in UK Biobank (UKB) and in the Health Surveys for England (HSE) and Scottish Health Surveys (SHS). Biobank is the largest spirometric study ever performed in the UK and included a relatively high proportion of never-smokers. The national health surveys, although based on smaller numbers of subjects, recruited a wider age range, have a longer period of follow-up, higher response rates and are more representative of the general UK population than Biobank. The two sources are therefore complementary, in terms of precision and generalisability.

This paper focuses on findings for white lifelong non-smokers, among whom confounding by frequency or duration of active smoking can be excluded. We investigate associations of spirometric indices with total mortality and with major groups of causes of death, compare FEV1 and FVC as independent predictors of all-cause mortality and evaluate the possibility of reverse causation. We also compare results from UKB with those from the national health survey participants of a similar age at spirometric examination, to establish how widely generalisable are the findings from Biobank. Corresponding results for white former smokers and current smokers are included in the online supplement for completeness and described briefly in the text.

METHODS

HSE and SHS

Data were combined from the HSE 1995, 1996 and 2001 and the SHS 1998 and 2003, the years when spirometry was included in the protocol. These surveys aimed to recruit a representative sample of British adults through household sampling within selected parliamentary constituencies throughout England or Scotland. Participants were visited at home. Response rates ranged from 60% to 76% across the five surveys. The proportion of those visited who performed usable spirometry ranged from 63% to 84%.

Spirometry was performed using handheld pneumotachograph spirometers (Vitalograph Escort) with the best results of FEV1 and FVC recorded from three technically satisfactory blows. No flow-volume curves or reproducibility criteria were available for assessment. Valid lung function measurements were available for 6579 lifelong non-smokers aged 40–69 years and 1429 aged 70 years or more at the start of follow-up, all of white ethnicity.

Deaths occurring up to April 2013 were available for analysis in the HSE, and deaths up to December 2011 were linked in the SHS. Combining all five surveys, there were 500 deaths among white never-smokers aged 40–69 years at recruitment over a mean follow-up period of 13.9 years. Deaths from respiratory disease, circulatory disease, cancer and all other causes were coded using International Classification of Diseases (ICD9) (460–519, 390–459, 140–208 and all others, respectively) and ICD10 (chapters J, I, C and all others, respectively).

Smoking history was self reported. Socioeconomic status was measured at the level of the household, based on the social class of the head of the household, Nation (England or Scotland), region (within England) and survey year were included as additional covariates.

UK Biobank

This study recruited 502 682 volunteers aged 40–69 years in 22 recruitment centres throughout England, Wales and Scotland during 2006–2010, following invitations to 9 million people. Spirometry was performed using a handheld pneumotachograph spirometer (Pneumotrac 6800) from which volume–time arrays were stored for each blow.

Eighty-one per cent of the cohort performed two blows with acceptable start and measures of FEV1 reproducible within 250 mL. This was considered the most inclusive sample of ‘usable spiromats’. When end-blow quality was also considered, 58% of the cohort had evidence of a good plateau and both FEV1 and FVC reproducible within 150 mL, the criteria recommended by the American Thoracic Society-European Respiratory Society (ATS/ERS) Task Force on Standardisation of Spirometry. This subgroup of 58% was considered to be the ‘best quality’ spiromats, among which to evaluate the relative importance of FEV1 and FVC as predictors of mortality.

The present analysis is based on deaths occurring up to mid-August 2015, a mean follow-up period of 6.5 years. There were 2401 deaths among 149 343 lifelong non-smokers aged 40–69 years of white ethnicity who performed ‘usable spiromats’. Deaths from respiratory disease, circulatory disease, cancer and all other causes were coded using ICD10 (chapters J, I, C and all others, respectively).

Smoking history was self reported. Socioeconomic status was measured at the level of the household, using the Townsend Deprivation Index, grouped into quartiles for analysis. Biobank recruitment centre was used as an additional covariate to adjust for possible regional differences.

Adjustment of spirometric measures for gender, age and height

The Global Lung Initiative (GLI) 2012 reference equations for white ethnic groups were used in both sets of data to standardise FEV1 and FVC for age, sex and height. The GLI-2012 equations generate a ‘z-score’ which represents the relative position of an individual among the distribution predicted for lifelong non-smokers with no history of lung disease of the same gender, age and height. This allows for the spread of predicted values to differ by age, height and gender, expressing the relative ranking of an
individual in terms of a SD (z) score. For each individual in the analysis, there were three GLI-2012 z-scores, corresponding to their relative range for FEV1 (zFEV1), FVC (zFVC) and the ratio FEV1:FVC (zFEVFVC). Outlying observations were excluded by restricting all the analyses in both data sets to values of zFEV1 and zFVC within the range −5 to +5 z-score units. This exclusion removed 0.2% of UKB participants, 0.5% of participants in the national health surveys aged 40–69 years and 0.3% of national health survey participants aged 70 years or more.

Modelling of mortality
The relationship of spirometric indices with subsequent mortality was modelled by proportional hazards (Cox) regression, which estimates the relative increase in mortality rate (HR) for a unit change in each explanatory variable. The z-scores are expressed on a SD scale, so HRs for zFEV1 and zFVC are expressed per unit decrement (ie, an increase in risk for a decrease in lung function). A typical range of z-scores among lifelong non-smokers would be 4 units. A HR of 1.2 per unit decrement corresponds approximately to a twofold difference in mortality rate across this range.

Due to the high correlation between zFEV1 and zFVC among lifelong non-smokers (0.88 in Biobank, 0.80 in national health survey participants aged 40–69 years), we modelled the association of mortality with zFEV1 both alone and jointly with zFEVFVC, and similarly for zFVC. Among never-smokers, the correlations between zFEV1 and zFEVFVC (0.35 in Biobank, 0.36 in the national surveys) and between zFVC and zFEVFVC (~0.13 in Biobank, ~0.20 in the national surveys) were weak enough to avoid major collinearity in the joint models. The significance of the HR for zFEVFVC when modelled jointly with zFEV1 was used to assess whether zFVC predicted mortality independent of zFEV1, and vice versa when zFEVFVC was modelled jointly with zFEV1.

All proportional hazards models were restricted to white participants and adjusted for sex, age, standing height, socioeconomic status and region. Analyses of data from the five national health surveys were additionally adjusted for survey year as a categorical variable. Since each survey was conducted in a different year, inclusion of survey year in the model is closely equivalent to a fixed-effect meta-analysis of the results from each of the five surveys. In a more formal two-stage individual participant meta-analysis for all-cause mortality in lifelong non-smokers, there was no substantial or significant heterogeneity of HRs among the five national health surveys (I²=0.0%, p=0.775 for zFEV1; I²=10.5%, p=0.346 for zFVC). Therefore, for simplicity of presentation we report results for the five national surveys combined, but analyse UKB separately because one of our objectives is to investigate how closely these two sets of results correspond. Heterogeneity between HRs for the pooled national surveys and UKB was assessed by testing the significance of the difference between the corresponding log-HRs from these two data sets.

The assumption of proportionality of hazards was assessed by log-log plots and by fitting zFEV1 or zFVC as a time-dependent covariate in the model. No strong or statistically significant evidence of time dependence emerged for all-cause mortality among lifelong non-smokers. Nevertheless, results for all-cause mortality were partitioned at 5 years of follow-up for two reasons. First, because the minimum duration of follow-up in UKB was 4.87 years, so virtually all of that cohort had been followed for 5 years or more, allowing a more direct comparison with results from the national health surveys, all of which had been followed for more than 5 years. The second reason for partitioning at 5 years was to address the possibility of reverse causation (impaired spirometric performance due to pre-existing conditions which lead to early death). Reverse causation was also investigated by analysing mortality in subgroups with no prior history of respiratory disease, circulatory disease or cancer.

RESULTS
Participant characteristics
Online supplementary e-table 1 summarises the number of participants, duration of follow-up and deaths from all causes and subgroups of cause, in each data set, by sex, age and smoking history.

All-cause mortality
Table 1 compares the HRs for age–sex–height-adjusted FEV1 and FVC in relation to all-cause mortality in the combined HSE and SHS data set, and UKB, among participants aged 40–69 years at entry. All ‘usable spiromgrams’ from Biobank were included in this comparison, because no additional quality control had been applied in the national health surveys 1995–2003.

When each spirometric index was modelled singly, all associations were highly statistically significant (p<0.0001). Among lifelong non-smokers in both cohorts, FEV1 and FVC displayed similar strengths of association with all-cause mortality (table 1). Among former smokers, FEV1 was the stronger predictor in HSE–SHS but FEV1 and FVC showed similar strength of association with all-cause mortality in UKB. Among current smokers in both data sets, FEV1 was a stronger predictor of all-cause mortality than FVC (see online supplementary e-tables 2–4).

Table 1 also shows the most direct comparison between the two data sets, based on all-cause mortality within 5 years. Although the associations of spirometric indices with these earlier deaths were stronger in HSE–SHS than in UKB, the differences between the cohorts were not statistically significant (p=0.23 for FEV1, p=0.39 for FVC). Associations with deaths after 5 years are less comparable between the data sets, due to the shorter period of follow-up in UKB.

Stronger associations of FEV1 and FVC with earlier deaths than with later mortality from all causes were also evident among former smokers and current smokers.
Table 1: All-cause mortality in white lifelong non-smokers aged 40–69 years at entry in national health surveys (HSE and SHS) and in UK Biobank, in relation to GLI-2012 z-scores for FEV1 and FVC

<table>
<thead>
<tr>
<th>Timing of death</th>
<th>Cause of death</th>
<th>HSE–SHS white lifelong non-smokers, 40–69 years at entry</th>
<th>Biobank white lifelong non-smokers, 40–69 years at entry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Any time</td>
<td>All causes</td>
<td>6579 (500) 1.17 (1.09 to 1.25) 1.19 (1.10 to 1.28) 149343 (2401) 1.21 (1.17 to 1.26) 1.24 (1.19 to 1.29)</td>
<td></td>
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<tr>
<td>Within 5 years</td>
<td>All causes</td>
<td>6579 (103) 1.35 (1.17 to 1.56) 1.35 (1.16 to 1.57) 149343 (1599) 1.23 (1.17 to 1.28) 1.26 (1.20 to 1.32)</td>
<td></td>
</tr>
<tr>
<td>After 5 years</td>
<td>All causes</td>
<td>6476 (397) 1.12 (1.03 to 1.21) 1.14 (1.05 to 1.24) 147744 (802) 1.18 (1.11 to 1.26) 1.21 (1.13 to 1.29)</td>
<td></td>
</tr>
<tr>
<td>Any time</td>
<td>Respiratory</td>
<td>6579 (24) 1.72 (1.34 to 2.21) 1.61 (1.25 to 2.08) 149343 (69) 1.86 (1.53 to 2.27) 2.15 (1.77 to 2.61)</td>
<td></td>
</tr>
<tr>
<td>Any time</td>
<td>Circulatory</td>
<td>6579 (130) 1.21 (1.06 to 1.38) 1.22 (1.06 to 1.40) 149343 (431) 1.41 (1.30 to 1.53) 1.47 (1.35 to 1.60)</td>
<td></td>
</tr>
<tr>
<td>Any time</td>
<td>Cancer</td>
<td>6579 (241) 1.10 (1.00 to 1.22) 1.12 (1.01 to 1.24) 149343 (1535) 1.08 (1.03 to 1.13) 1.10 (1.05 to 1.15)</td>
<td></td>
</tr>
<tr>
<td>Any time</td>
<td>Other non-respiratory</td>
<td>6579 (105) 1.15 (0.98 to 1.34) 1.20 (1.03 to 1.41) 149343 (366) 1.46 (1.33 to 1.59) 1.45 (1.32 to 1.59)</td>
<td></td>
</tr>
</tbody>
</table>

Spirometric indices (FEV1 and FVC) are modelled as z-scores derived from the Global Lung Initiative (GLI) 2012 reference equations for whites. All HRs are expressed per unit decrement in z-score, adjusted for age, sex, height, socioeconomic status, geographical region and survey year. FEV1, one-second forced expiratory volume; FVC, forced vital capacity; HSE, Health Survey for England; SHS, Scottish Health Survey; z, z-score.

Influence of spirogram quality

Table 2 presents the results for all causes of death among Biobank participants with ‘best quality’ spirograms, for comparison with those obtained from the full set of UKB participants with ‘best quality’ spirograms. The pattern was evident in both data sets. Although HRs were generally consistent between the ‘best’ spirogram subgroup and the full Biobank, a comparison of results in the two cohorts is more similar to the former than the latter. Within UKB, results for best quality spirograms were generally consistent between the ‘best’ spirogram subgroup and the full Biobank.

Table 2: Cause-specific mortality

<table>
<thead>
<tr>
<th>Timing of death</th>
<th>Cause of death</th>
<th>HSE–SHS white lifelong non-smokers, 40–69 years at entry</th>
<th>Biobank white lifelong non-smokers, 40–69 years at entry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Any time</td>
<td>Respiratory</td>
<td>6579 (24) 1.72 (1.34 to 2.21) 1.61 (1.25 to 2.08) 149343 (69) 1.86 (1.53 to 2.27) 2.15 (1.77 to 2.61)</td>
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<td>Cancer</td>
<td>6579 (241) 1.10 (1.00 to 1.22) 1.12 (1.01 to 1.24) 149343 (1535) 1.08 (1.03 to 1.13) 1.10 (1.05 to 1.15)</td>
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<td>6579 (105) 1.15 (0.98 to 1.34) 1.20 (1.03 to 1.41) 149343 (366) 1.46 (1.33 to 1.59) 1.45 (1.32 to 1.59)</td>
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Spirometric indices (FEV1 and FVC) are modelled as z-scores derived from the Global Lung Initiative (GLI) 2012 reference equations for whites. All HRs are expressed per unit decrement in z-score, adjusted for age, sex, height, socioeconomic status, geographical region and survey year. FEV1, one-second forced expiratory volume; FVC, forced vital capacity; HSE, Health Survey for England; SHS, Scottish Health Survey; z, z-score.
<table>
<thead>
<tr>
<th>Subset analysed</th>
<th>z-score in model (plus covariates)</th>
<th>HSE–SHS white lifelong non-smokers, 40–69 years at entry</th>
<th>Biobank white lifelong non-smokers, 40–69 years at entry</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Total N (deaths) HR (95% CI) alone</td>
<td>Total N (deaths) HR (95% CI) joint</td>
<td>Total N (deaths) HR (95% CI) alone</td>
</tr>
<tr>
<td>All usable spiromgrams</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>FEV1 alone</td>
<td>6579 (500) 1.17 (1.09 to 1.25)</td>
<td>149343 (2401) 1.21 (1.17 to 1.26)</td>
<td></td>
</tr>
<tr>
<td>FEV/FVC adj FEV1</td>
<td>1.20 (1.11 to 1.29)</td>
<td>1.26 (1.21 to 1.31)</td>
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<tr>
<td>FEV/FVC adj FVC</td>
<td>0.94 (0.87 to 1.01)</td>
<td>0.90 (0.85 to 0.94)</td>
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<tr>
<td>All usable spiromgrams</td>
<td></td>
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</tr>
<tr>
<td>FVC alone</td>
<td>6579 (500) 1.19 (1.10 to 1.28)</td>
<td>149343 (2401) 1.24 (1.19 to 1.29)</td>
<td></td>
</tr>
<tr>
<td>FVC adj FEV/FVC</td>
<td>1.19 (1.11 to 1.28)</td>
<td>1.24 (1.19 to 1.29)</td>
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<tr>
<td>FVC adj FVC</td>
<td>1.02 (0.95 to 1.10)</td>
<td>1.02 (0.97 to 1.07)</td>
<td></td>
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<tr>
<td>Best quality spiromgrams</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FEV1 alone</td>
<td>(No data)</td>
<td>102945 (1583) 1.23 (1.18 to 1.29)</td>
<td></td>
</tr>
<tr>
<td>FEV1 adj FEV/FVC</td>
<td></td>
<td>1.29 (1.23 to 1.36)</td>
<td></td>
</tr>
<tr>
<td>FEV/FVC adj FEV1</td>
<td></td>
<td>0.86 (0.81 to 0.91)</td>
<td></td>
</tr>
<tr>
<td>Best quality spiromgrams</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FVC alone</td>
<td>(No data)</td>
<td>102945 (1583) 1.29 (1.23 to 1.35)</td>
<td></td>
</tr>
<tr>
<td>FVC adj FEV/FVC</td>
<td></td>
<td>1.29 (1.23 to 1.35)</td>
<td></td>
</tr>
<tr>
<td>FVC adj FVC</td>
<td></td>
<td>0.98 (0.93 to 1.04)</td>
<td></td>
</tr>
</tbody>
</table>

Spirometric indices (FEV1, FVC and FEV1:FVC ratio) are modelled as z-scores derived from the Global Lung Initiative 2012 reference equations for whites. All HRs are expressed per unit decrement in z-score, adjusted for age, sex, height, socioeconomic status, geographical region and survey year. FEV1, one-second forced expiratory volume; FVC, forced vital capacity; HSE, Health Survey for England; SHS, Scottish Health Survey.
data set in all smoking subgroups (see online supplementary e-table 4).

Comparisons by subgroups of age and sex
Online supplementary e-table 3 shows that within UKB the general pattern of results was similar in both sexes. There was no statistically significant effect modification by sex for either spirometric index in any smoking subgroup (p>0.10 for each interaction test).

Online supplementary e-table 3 also compares the results for all-cause mortality among younger (aged 40–59 years) and older (aged 60–69 years) UKB participants. The HRs in all smoking groups were consistent between these two age subgroups and there were no statistically significant age interactions for either spirometric index (p>0.40).

Online supplementary e-table 5 compares the spirometric associations with all-cause and cause-specific mortality among HSE and SHS participants aged 40–69 years and 70 years or more at entry. The pattern for all causes was consistent between these age groups in never-smokers and current smokers (p>0.2), but there were significant age interactions among ex-smokers (p<0.006).

Online supplementary e-table 3 compares the results for all-cause mortality among obese and non-obese Biobank participants. Although the pattern of results was generally consistent between these categories in all smoking subgroups, a statistically significant difference in the HR for FVC occurred among current smokers (p=0.022). Other interactions with obesity were non-significant (p>0.10).

Comparisons by prior disease history in UKB
Table 3 compares the spirometric associations with all-cause mortality in Biobank participants with and without a history of respiratory disease at the baseline spirometric examination. In the 18,545 (12%) white never-smokers with ‘usable spiromograms’ who had a history of diagnosed respiratory disease (of which 90% reported asthma), FVC was a stronger predictor of all-cause mortality than FEV1. Among the subgroup without a respiratory history, results were similar to the full cohort.

Lung function also emerged as a significant predictor of circulatory mortality among those with and without a prior history of heart attack, angina, stroke, thrombosis or hypertension (table 3). This pattern was confirmed among former smokers and current smokers (see online supplementary e-table 3).

Finally, the association of FEV1 and FVC with cancer mortality was shown to be stronger in those with a prior cancer diagnosis (table 3), but it was not statistically significant, despite the large sample size, among those with no cancer history at the spirometric examination. However, this finding is confined to the lifelong non-smokers: both FEV1 and FVC were more strongly and significantly associated with cancer death among former smokers and current smokers with no cancer history (see online supplementary e-table 6). Cancer mortality among participants with a history of cancer at entry was strongly and significantly associated with both FEV1 and FVC in all three smoking subgroups (see online supplementary e-table 6).

Comparison of spirometry with other predictors of mortality
Figure 1 (data in online supplementary e-table 7) compares the relative mortality across quartiles of body mass index, systolic blood pressure and FVC z-score, for all deaths and for deaths from circulatory disease, among white lifelong non-smokers in HSE–SHS and in UKB. A similar pattern emerged for all-cause mortality in both cohorts, with differentials in mortality across quartiles of zFVC being at least as great as those across quartiles of body mass index or systolic blood pressure.

DISCUSSION
Principal findings
A broadly coherent picture emerges from this comparison of UK national cohorts. Ventilatory function, even if measured imperfectly, consistently predicted both respiratory deaths and non-respiratory mortality from a range of causes. This was found even among lifelong non-smokers, so confounding by the amount or duration of active smoking is not the sole explanation. Both for all-cause mortality and more specifically for circulatory disease mortality, FEV1 and FVC were as strongly predictive as body mass index and more strongly predictive than systolic blood pressure.

Strengths and weaknesses of this study
UKB offers a spirometric study of lifelong non-smokers of unprecedented size, but its 5.5% participation rate may have compromised its generalisability. Assembling data from five UK national health surveys produced a cohort of never-smokers, larger than the combined number of participants in previous publications, in which the generalisability of Biobank results could be tested. The similar pattern of results in HSE–SHS and UKB suggests that the key findings are generalisable, at least to the British population.

Within Biobank, the large numbers permitted subgroup analyses by sex, age, obesity and pre-existing disease, of sufficient statistical power to exclude important interaction effects. These within-cohort comparisons provide further reassurance about the generalisability of the principal findings among lifelong non-smokers.

Although only 58% of the Biobank cohort performed spirometry which fulfilled internationally recommended criteria for acceptability and reproducibility, the results in this subgroup were very similar to those among the full set of Biobank participants who performed ‘usable’ spirometry. Those results were, in turn, consistent with the findings from national health surveys where the acceptability and reproducibility of spirometry was not formally assessed in the field. These within-cohort and cross-cohort comparisons suggest that the principal
Table 3 Comparison of spirometric prediction of mortality among subjects with and without prior disease history at entry in UK Biobank

<table>
<thead>
<tr>
<th>Condition at entry</th>
<th>Cause of death</th>
<th>Biobank age 40–69 years, white lifelong non-smokers without a history of the condition at entry</th>
<th>Biobank age 40–69 years, white lifelong non-smokers with a history of the condition at entry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Respiratory disease</td>
<td>All causes</td>
<td>130,798 (2081) 1.21 (1.16 to 1.26) 1.22 (1.17 to 1.27) 18,545 (320) 1.23 (1.12 to 1.35) 1.33 (1.21 to 1.47)</td>
<td></td>
</tr>
<tr>
<td>Circulatory disease</td>
<td>Circulatory disease</td>
<td>109,141 (189) 1.40 (1.23 to 1.59) 1.40 (1.23 to 1.60) 40,202 (242) 1.35 (1.20 to 1.51) 1.43 (1.27 to 1.60)</td>
<td></td>
</tr>
<tr>
<td>Cancer</td>
<td>Cancer</td>
<td>137,742 (978) 1.05 (0.99 to 1.11) 1.06 (1.00 to 1.12) 11,801 (557) 1.10 (1.02 to 1.18) 1.13 (1.04 to 1.22)</td>
<td></td>
</tr>
</tbody>
</table>

Spirometric indices (FEV1 and FVC) are modelled as z-scores derived from the Global Lung Initiative 2012 reference equations for whites. All HRs are expressed per unit decrement in z-score, adjusted for age, sex, height, socioeconomic status and geographical region (recruitment centre).

FEV1, one-second forced expiratory volume; FVC, forced vital capacity; z, z-score.

# Comparison with other studies

Previous studies of lifelong non-smokers have been of limited size: 662 males and 2048 females in the Copenhagen City Heart Study, of whom 195 died during 10 years of follow-up,3 and 3562 male London civil servants in the Whitehall Study, of whom 408 died over a period of 18 years.4 The Whitehall cohort was subsequently followed for 33–35 years, accumulating 1545 deaths among 3083 lifelong non-smokers.19 In the analyses we present here, there are almost twice this number of deaths among lifelong non-smokers, despite a shorter follow-up period, due to the much larger sample size at entry, particularly in UKB.

The two publications from the Whitehall cohort12,13 compared FEV1 and height as predictors of all-cause and cause-specific mortality among never-smokers. The first12 found that FEV1 predicted mortality independent of height, but height did not predict survival independent of FEV1. The second13 found that FEV1 and height were similarly related both to mortality and to a range of other risk factors, concluding that both FEV1 and height may be markers of early life exposures of relevance to longevity.

In our study of lifelong non-smokers, we confirmed that in determining survival in middle-aged never-smokers, FVC (rather than FEV1) is the index of greater importance, with FVC and height both significantly related to mortality. However, no formal comparison was made between the two indices. The Copenhagen cohort14 compared FEV1 and height as predictors of all-cause mortality among never-smokers, finding a stronger association with FEV1 than with height, but height did not predict survival independently of FEV1. The second15 found that both FEV1 and height were similarly related both to mortality and to a range of other risk factors, concluding that both FEV1 and height may be markers of early life exposures of relevance to longevity.

In contrast, among current smokers, FEV1 emerged as the more influential predictor. This may be because the recent linkage of hospital admissions and primary care consultations to the UKB cohort will allow associations with incidence and case fatality to be investigated more directly in future.

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FEV1:FVC ratio declines with both the dose and duration of smoking, and these also increase mortality risk. The ability to perform good quality spirometry is an integrated assessment of physical and cognitive function and therefore might be considered a predictor of mortality in its own right. In the US Six Cities Study, excessively variable spirometric performance was an indicator of poor health and associated with shorter survival.\(^20\) In contrast, the mortality experience of Biobank participants who produced ‘best quality’ spirograms did not differ greatly from that of their peers who produced “usable’ ‘but not’ ‘best quality’ blows.

**Possible implications**

In clinical practice, particularly in primary care, quality control of spirometry is unlikely to be much better than in the national health surveys where lung function was tested by a trained research nurse in the home setting. Therefore, while the results from the Biobank ‘best quality’ subgroup are of confirmatory interest, the more inclusive results for all ‘usable spirograms’ may be more generally relevant.

In both cohorts, age–sex–height-adjusted lung function emerged as a stronger predictor of all-cause mortality than either systolic blood pressure or body mass index, which are, respectively, the second and sixth most influential causes worldwide of loss of healthy lifespan, as measured by disability-adjusted life years.\(^21\) It is therefore puzzling to find U-shaped or J-shaped relationships of mortality with these two cardiovascular risk factors, but the similar patterns of results in Biobank and the national health surveys suggest that this is not a unique feature of either of these British cohorts.

Specifically for circulatory disease mortality, FEV1 and FVC were as strongly predictive as body mass index and more strongly predictive than systolic blood pressure. Therefore, spirometry may deserve consideration as an addition to cardiovascular risk scoring algorithms in future.

**CONCLUSION**

More generally, these results emphasise the potential importance of promoting and protecting lung health in the general population, even among lifelong non-smokers with no history of respiratory disease.

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**Competing interests** None declared.

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