Contributions of cardiovascular risk and smoking to chronic obstructive pulmonary disease (COPD)-related changes in brain structure and function

Background: Brain damage and cardiovascular disease are extra-pulmonary manifestations of chronic obstructive pulmonary disease (COPD). Cardiovascular risk factors and smoking are contributors to neurodegeneration. This study investigates whether there is a specific, COPD-related deterioration in brain structure and function independent of cardiovascular risk factors and smoking.

Materials and methods: Neuroimaging and clinical markers of brain structure (micro- and macro-) and function (cognitive function and mood) were compared between 27 stable COPD patients (age: 63.0±9.1 years, 59.3% male, forced expiratory volume in 1 second [FEV₁]: 58.1±18.0% pred.) and 23 non-COPD controls with >10 pack years smoking (age: 66.6±7.5 years, 52.2% male, FEV₁: 100.6±19.1% pred.). Clinical relationships and group interactions with brain structure were also tested. All statistical analyses included correction for cardiovascular risk factors, smoking, and aortic stiffness.

Results: COPD patients had significantly worse cognitive function (p=0.011), lower mood (p=0.046), and greater gray matter atrophy (p=0.020). In COPD patients, lower mood was associated with markers of white matter (WM) microstructural damage (p<0.001), and lower lung function (FEV₁/forced vital capacity and FEV₁) with markers of both WM macro (p=0.047) and microstructural damage (p=0.028).

Conclusion: COPD is associated with both structural (gray matter atrophy) and functional (worse cognitive function and mood) brain changes that cannot be explained by measures of cardiovascular risk, aortic stiffness, or smoking history alone. These results have important implications to guide the development of new interventions to prevent or delay progression of neuropsychiatric comorbidities in COPD. Relationships found between mood and microstructural abnormalities suggest that in COPD, anxiety, and depression may occur secondary to WM damage. This could be used to better understand disabling symptoms such as breathlessness, improve health status, and reduce hospital admissions.

Keywords: chronic lung disease, cigarette smoke, cognition, depression, MRI, neuroimaging

Introduction

Chronic obstructive pulmonary disease (COPD) is associated with a number of extra-pulmonary, manifestations including cardiovascular disease, diabetes, arthritis, osteoporosis, obesity, metabolic syndrome, muscle weakness, sleep disturbance, and anemia.1,2 These occur at a higher rate than in smokers and never-smokers,3 have a deleterious effect on patient outcomes and wellbeing,4,5 and contribute...
substantially to the financial burden of the disease. However, the disease presentation is highly heterogeneous and it is currently unclear whether these comorbidities are pathogenically linked to the disease or reflect the co-existence of numerous age-related risk factors and conditions.

Cognitive dysfunction, anxiety, and depression are key comorbidities of COPD where they are associated with greater disability, poorer medical compliance, increased risk of exacerbation, and mortality. However, their pathophysiology in relation to COPD remains poorly understood. Neuroimaging findings suggest that there are underlying changes to brain structure and function associated with both reduced lung function in the general population and with established COPD. The pattern of structural changes is consistent with cerebral small-vessel disease (SVD). However, the majority of these studies have failed to adequately control for individual differences in cardiovascular risk and smoking history; factors known to accelerate age-related SVD. Consequently, it cannot be established whether this is a COPD-specific effect per se, or the cumulative effect of greater cardiovascular risk in COPD.

The objectives of this study are to determine whether COPD is associated with specific differences in brain structure (macro- and micro-) and function (cognitive function and mood) beyond those which can be attributed to cardiovascular risk factors or smoking. This will be achieved by cross-sectional comparison of a very well-characterized cohort of COPD patients with age-matched non-COPD control subjects with a history of smoking. Additionally, the relationships between clinical measures and brain structure and whether these relationships differ between groups will be investigated in order to establish whether the same disease processes are active in both COPD patients and non-COPD smokers.

It is hypothesized that when compared to non-COPD smokers, COPD patients will show evidence of differences in brain structure and function that cannot be explained by cardiovascular risk factors and smoking. It is predicted that the pattern of differences in brain structure will be consistent with age-related SVD.

Materials and methods
Study participants
Twenty-seven stable COPD patients (defined as forced expiratory volume in 1 second (FEV1)/forced vital capacity (FVC) <70% and/or clinical/radiological diagnosis of emphysema who had not experienced an exacerbation within the preceding 4 weeks) were recruited from inpatient and outpatient clinics at North Bristol NHS Trust, and 23 non-COPD smoker controls were recruited from the Bristol Primary Care Research North Hub between 2014 and 2015. All subjects provided written informed consent for participation in the study. Inclusion criteria required all subjects to be current or ex-smokers with greater than 10 pack years smoking history. Subjects were excluded from participation if their resting oxygen saturations (SaO2) were below 92% on room air, were on long-term oxygen therapy, they had known alpha-1-anti-trypsin deficiency or co-morbid neurological, cardiovascular, or psychiatric conditions likely to affect neuroimaging findings. Subjects with incidental findings on neuroimaging, visual, or hearing impairment that precluded neuropsychological assessment or contraindications for magnetic resonance imaging (MRI) were also excluded. For a full list of inclusion and exclusion criteria see Table S1. Groups were well matched for age, sex, educational attainment category (defined as, none, GCSE, A-level, degree or other), body mass index, blood pressure, and smoking status. However, COPD patients had smoked for a numerically greater number of pack years (see Table 1).

Ethics statement
Ethical approval was obtained from the NRES Committee South West – Frenchay (13/SW/0319). The study was conducted in accordance with the Declaration of Helsinki.

Procedure
The data used in this study formed part of a larger protocol, therefore, data acquisition took place across three study visits. Standard demographic information (age, sex, educational attainment category, smoking status, pack year history, body mass index) and clinical measures of disease severity (post-bronchodilator spirometry performed as per the recommendations of the American Thoracic Society and European Respiratory Society Consensus Statement, disease status (COPD Assessment Test, CAT), SaO2 (by pulse oximetry), mood (Hospital Anxiety and Depression Scale, HADS) and cognitive function (Montreal Cognitive Assessment, MoCA) were collected on visit one. A brief description of the HADS and MoCA can be found in the Supplementary Material. Aortic stiffness (aortic pulse-wave velocity and central augmentation index measured using SphygmoCor system), blood pressure and earlobe capillary blood gasses (PO2, PCO2, and pH) were collected on
visit two at the Respiratory Research Unit at the North Bristol Lung Centre. MRI scans were acquired on visit three at the Clinical Research Imaging Centre (CRICBristol), Bristol University.

### Image acquisition and processing

3-Tesla T₁-weighted (T1W), fluid-attenuated inversion recovery (FLAIR) and diffusion tensor images (DTI) were acquired for all subjects allowing measures of tissue...
macrostructure and tissue microstructure to be calculated. Representative images for each group can be found in Figure 1. This procedure is summarized below. For detailed description of image acquisition and processing see the Supplementary Material.

**Tissue macrostructure**
Supratentorial gray matter, white matter, and cerebrospinal fluid (CSF) tissue volumes were calculated from the T1W images and white matter hyperintensities (WMHs) volumes from the T1W and FLAIR images using a semi-automated procedure described in Spilling et al 2017 and Lambert et al 2015. All volumes were normalized for head size (% of total intracranial volume, TIV).

**Tissue microstructure**
Fractional anisotropy (FA) (local tissue directionality) and mean diffusivity (MD) (local magnitude of diffusion) were calculated for each voxel of the DTI. The median and normalized peak height of the distribution of FA and MD values within the normal-appearing white matter (NAWM) (white matter excluding WMHs) were calculated.

**Statistical analysis**
Statistical analysis was performed using IBM SPSS Statistics (IBM SPSS, version 24). Data residuals were checked for Gaussian distribution using Shapiro–Wilk’s tests, histograms, and quartile-quartile plots. Non-Gaussian data were log10-transformed (or reflected and log10-transformed). Group differences in clinical measures (including demographic information, disease severity, aortic stiffness, blood pressure, blood gases, mood, and cognitive function) and brain macrostructure (normalized gray matter, white matter, CSF, and WMH volumes) and microstructure (median FA, FA peak height, median MD, MD peak height) were tested using analysis of covariance and chi-squared tests. Relationships between clinical measures and brain structure, and group interactions with these measures were tested using multiple linear regression models. Post-hoc multiple linear regression models were used to probe significant interactions. Results were considered significant at \( p < 0.05 \). Unless indicated otherwise, all statistical analyses were performed using models with demographic and cardiovascular risk factors entered as covariates of no interest (age, sex, smoking status, pack year history, mean arterial pressure, and aortic pulse-wave velocity). Additionally, educational attainment category was included in any model testing group differences or relationships with cognitive function.

**Results**
**Between-group differences**

**Clinical measures**
Group comparisons of clinical measures can be found in Table 1. As expected, COPD patients had significantly
lower lung function (p<0.001) and worse disease status (p<0.001) than non-COPD smoker controls, with unstandardized regression coefficients indicating that COPD was associated with a decrease in FEV₁ of 44% pred. and an increase in CAT score of 10. Following correction for cardiovascular risk, COPD patients had significantly worse mood (p=0.046) and following additional correction for educational attainment, worse cognitive function (p=0.011) than non-COPD smoker controls. Unstandardized regression coefficients indicated that COPD was associated with an increase in HADS total score of 3 points and decrease in MoCA – total score of 2 points. There were no group differences in SaO₂ or in aortic pulse-wave velocity and central augmentation index (see Table 1).

Brain structure
Following correction for cardiovascular risk, COPD patients were found to have significantly lower normalized gray matter volume than non-COPD smoker controls (p=0.020). Unstandardized regression coefficients indicated that the presence of COPD was associated with a 1.1% decrement in normalized gray matter volume. No other significant differences were found for measures of brain macro- or microstructure (see Table 2).

Clinical relationships and group interactions
Lung function, disease status, and blood gases
Multiple linear regression models assessing the main effect and group interactions of lung function (FEV₁ and FEV₁/FVC) on brain structure showed that the FEV₁ model explained 37% of the variance in median MD within the NAWM (r²=0.37, p=0.029) and that the interaction between group and FEV₁ significantly predicted median MD (p=0.032). Post-hoc analysis indicated that for COPD patients lower FEV₁ was related to greater median MD (p=0.028), but not for non-COPD smoker controls (p=0.133). Similarly, multiple linear regression models assessing the main effects and group interactions of FEV₁/FVC explained 43% of the variance in normalized white matter volume (r²=0.43, p=0.005) with the group interaction significantly predicting normalized white matter volume (p=0.039). Post-hoc analysis indicated that lower FEV₁/FVC was related to lower normalized white matter volume in COPD patients (p=0.047) but not in non-COPD smoker controls (p=0.461). These interactions are shown in Figure 2A, B. No relationships were found between CAT and SaO₂ and brain structure.

Aortic stiffness
Multiple linear regression models found no significant main effects or group interactions of aortic stiffness (aortic pulse-wave velocity and central augmentation index) on brain structure.

Cognitive function
Multiple linear regression models assessing the main effect and group interactions of MoCA – total score did not find any significant relationships with brain structure.

Mood
The multiple linear regression model assessing the main effects and group interactions of mood (HADS – total score) on brain structure explained 38% of the variance in median FA (r²=0.38, p=0.022), 34% of the variance in median MD (r²=0.34, p=0.048) and 57% of the variance in

**Table 2** Group differences in brain structure

<table>
<thead>
<tr>
<th></th>
<th>Controls</th>
<th>COPD patients</th>
<th>F</th>
<th>B</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grey Matter Volume (% TIV)</td>
<td>42.6±1.4</td>
<td>42.0±1.7</td>
<td>5.82±1</td>
<td>−1.09±</td>
<td>0.020±*</td>
</tr>
<tr>
<td>White Matter Volume (% TIV)</td>
<td>27.9±2.6</td>
<td>28.1±1.7</td>
<td>0.031±</td>
<td>0.117±</td>
<td>0.861±</td>
</tr>
<tr>
<td>CSF Volume (% TIV)</td>
<td>29.5±2.5</td>
<td>29.8±2.6</td>
<td>1.71±4*</td>
<td>0.973±</td>
<td>0.198±</td>
</tr>
<tr>
<td>White Matter Hyperintensity Volume (% TIV)</td>
<td>0.22 (0.14)</td>
<td>0.30 (0.32)</td>
<td>0.16±3*</td>
<td>−0.004±</td>
<td>0.989±</td>
</tr>
<tr>
<td>Median FA</td>
<td>0.43±0.01</td>
<td>0.42±0.02</td>
<td>0.74±8*</td>
<td>−0.005±</td>
<td>0.392±</td>
</tr>
<tr>
<td>FA peak height (%) (×10⁻²)</td>
<td>2.88±0.16</td>
<td>2.82±0.11</td>
<td>0.57±9*</td>
<td>0.000±</td>
<td>0.451±</td>
</tr>
<tr>
<td>Median MD (×10⁻⁴ mm²/s)</td>
<td>7.44±0.17</td>
<td>7.52±0.37</td>
<td>0.79±9*</td>
<td>0.000±</td>
<td>0.377±</td>
</tr>
<tr>
<td>MD peak height (%)</td>
<td>0.18±0.02</td>
<td>0.18±0.02</td>
<td>1.00±8*</td>
<td>−0.006±</td>
<td>0.323±</td>
</tr>
</tbody>
</table>

Notes: Group comparison of brain macro- and microstructure. For Gaussian raw unadjusted data means ± standard deviations are presented, for non-gaussian raw unadjusted data medians (interquartile ranges) are presented. *Parametric ANCOVAs; †ANCOVAs performed on transformed data, F-statistics, unstandardized regression coefficients (B) and p-values are presented. ‡Model included age, sex, smoking status, pack years smoked, mean arterial pressure, and aortic pulse-wave velocity as covariates of no interest. *p<0.05.

Abbreviations: TIV, total intracranial volume; CSF, cerebrospinal fluid; FA, fractional anisotropy; MD, mean diffusivity.
MD peak height ($r^2=0.57, p<0.001$). It showed that there were significant main effects of greater HADS – total score on lower median FA ($p=0.009$), lower MD peak height ($p=0.005$) and higher median MD ($p=0.038$) and group by total HADS score interactions for median FA ($p=0.011$), median MD ($p=0.012$), and MD peak height ($p=0.020$). Post-hoc analysis indicated that for COPD patients higher HADS – total score was related to lower median FA ($p<0.001$), lower MD peak height ($p<0.001$), and higher median MD ($p<0.001$) whereas there were no relationships for non-COPD smoker controls: median FA: $p=0.616$; MD peak height: $p=0.125$ and median MD: $p=0.334$. Graphs showing these interactions can be found in Figure 2C, D.

**Discussion**

This study was designed to test whether COPD is an additional independent risk factor for deterioration in brain structure and function beyond that attributable to age, traditional vascular risk factors, and smoking. COPD patients showed evidence of greater cerebral atrophy, lower cognitive function, and worse mood. Additionally, exploratory tests of group interactions showed that lower lung function (FEV$_1$ and FEV$_1$/FVC) was associated with a deterioration in white matter macro- and microstructure, and worse mood state was associated with a deterioration in white matter microstructure in COPD patients but not in non-COPD smokers.

Previous studies have shown that COPD and reduced lung function are associated with a number of neuroimaging features of small-vessel cerebrovascular disease, including small subcortical infarcts, WMHs, white matter microstructural abnormalities, cerebral microbleeds, and cerebral atrophy.\textsuperscript{15-29} Consequently, it has been suggested that COPD-related deteriorations in brain structure and function occur secondary to SVD\textsuperscript{15,42} either due to the high prevalence of age-related vascular risk factors or because COPD is itself a risk factor. Our study could not replicate these white matter findings\textsuperscript{15,17,19,22,25} and we suggest that they are likely to have been caused by group differences in cardiovascular risk and smoking exposure. However, in our study, COPD patients did have greater cerebral atrophy, lower cognitive function and worse mood state than non-COPD smokers and these could not be explained by differences in cardiovascular risk factors or smoking. The 1.1% decrement in normalized gray matter volume associated with COPD in this study is greater than the annual rate of gray matter volume and brain volume decline reported in normal aging (ranging approximately between 0.3% and 0.8%) eg,\textsuperscript{43-45} and SVD (0.9%).\textsuperscript{46} It is
also similar in magnitude to the decrease in gray matter volume cross-sectionally associated with diabetes mellitus when compared to normal controls eg, 1.2%. Consistent with Cleutjens et al,48 no relationships were found between brain structure and the lower cognitive function in COPD patients. However, significant relationships were found between reduced lung function and greater macro- and microstructural white matter abnormalities in the COPD patients which suggests that other mechanisms are contributing to neurodegeneration and impaired cognition and mood in COPD.

The generalized decrease in gray matter volume found in this study has not been shown previously in COPD, although other studies have reported localized reductions in gray matter density, cortical thickness, and hippocampal volume19–24 notably in regions associated with dyspnea and fear of physical activity.19,20,24 We used a segmentation technique optimized for use in elderly cohorts with WMHs,25,41 and use of this technique may have improved sensitivity to detecting subtle gray matter alterations not detectable in other studies. Alternatively, the inconsistency in gray matter findings may reflect heterogeneity between COPD cohorts.3

Like other chronic diseases, COPD is associated with a higher prevalence of anxiety and depression than the general population,49 with reported rates ranging from 7% to 50% and from 10% to 57%, respectively.10 Using the HADS, the present study found some degree of anxiety or depression in 39% and 22% of COPD patients, respectively, with COPD patients having significantly lower overall mood than non-COPD smokers. Group interactions were found such that worse mood was associated with greater white matter microstructural abnormalities in COPD patients but not in non-COPD smokers. Similar cross-sectional relationships have been reported in normal elderly and SVD cohorts where white matter alterations (both in terms of white matter microstructural change and severity of WMHs) were associated with higher incidence of depressive disorder, greater disability, and lower depression remission rates.50–54 A recent meta-analysis found that multiple markers of both cerebral (WMHs, microbleeds, and microinfarctions) and peripheral forms (plasma markers of endothelial dysfunction) of microvascular dysfunction were associated with increased odds of incident late-life depression.55 Furthermore, longitudinal studies have suggested that markers of SVD may precede the onset of depressive symptom.54,55 In these circumstances, it has been hypothesized that depression results from localized disruption of frontostriatal white matter networks involved in affective regulation.56 This is consistent with functional MRI findings in COPD, which show that COPD patients have an enhanced neural responses in gray matter regions involved in emotional processing and memory (including the medial prefrontal cortex,31 anterior cingulate cortex,31 amygdala30 and hippocampus30), to anticipation of dyspnea30 and dyspnea-related word-cues.31 These enhanced responses are associated with worse perception of dyspnea,30 greater anxiety30, and depression31 and worse disease status (health-related quality of life and exercise tolerance). Furthermore, it has been demonstrated to respond to pulmonary rehabilitation.32 However, recent studies suggest that frontostriatal network disruption leads to apathy rather than depression in SVD.57,58

**Advantages and limitations**

This study benefits from a well-defined stable COPD cohort and successful recruitment of a non-COPD cohort with a history of smoking, allowing the effects of smoking and COPD to be differentiated. The sample size is comparable to other similar MRI studies of the brain in COPD,18–20,22,23,25 however, modest sample size may limit the generalisability of these results in COPD patients with other levels of disease severity. The secondary analyses testing for clinical relationships and group interaction with brain measures were exploratory. The covariates were grouped into pre-defined clinically meaningful domains to reduce multiple comparisons and linear regression models were adjusted for demographic and cardiovascular risk factors, however, we are unable to completely exclude the possibility of type-I statistical error due to multiple comparisons. This study used the HADS to evaluate participants’ overall mood. This questionnaire was originally developed as a clinical screening tool with two separate sub-scales measuring anxiety and depression37 meaning that the present study has extended its use beyond its original intent. The HADS has known limitations in terms of the stability of this underlying factor structure (particularly in disease)59 and ceiling effects on individual items.60 A number of other authors have supported the validity of using the total HADS score (as in the present study) as a measure of overall psychological distress (eg,61–63). Serum cholesterol, a vascular risk factor, was unavailable for this dataset and could not be controlled for in statistical analyses.
Conclusion

COPD is associated with a specific pattern of structural and functional brain abnormalities that could not be explained by conventional measures of cardiovascular risk, smoking history or aortic stiffness. Worse lung function is associated with deterioration in white matter macro- and microstructure, and deterioration in white matter microstructure is associated with lower mood. This suggests that mechanisms other than cardiovascular risk and smoking contribute to brain changes in COPD. Cognitive dysfunction, anxiety, and depression are key comorbidities of COPD and are associated with greater disability risk of exacerbation and mortality. COPD-related anxiety and depression may occur secondary to white matter damage. These findings have important implications for the prevention and management of neuropsychiatric comorbidities in COPD.

Abbreviations

ANCOVA(s), analysis of covariance; BET, brain extraction tool; BMI, body mass index; CAT, COPD Assessment Test; COPD, chronic obstructive pulmonary disease; CRIC, Clinical Research Imaging Centre; CSF, cerebrospinal fluid; DTI, diffusion tensor imaging; FA, fractional anisotropy; FEV₁, forced expiratory volume in 1 Second; FLAIR, fluid attenuated inversion recovery; FVC, forced vital capacity; HADS, Hospital Anxiety and Depression Scale; IQR, interquartile range; MD, mean diffusivity; MoCA, Montreal Cognitive Assessment Test; MRI, magnetic resonance imaging; NAWM, normal-appearing white matter; PO₂, partial pressure of oxygen; PCO₂, partial pressure of carbon dioxide; SD, standard deviation; SaO₂, oxygen saturation; SVD, small vessel disease; TE, echo time; TI, inversion time; TIV, total intracranial volume; TR, repetition time; T₁W, T₁-weighted; WMHs, white matter hyperintensities.

Acknowledgment

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Author contributions

JWD and PWJ designed the study, JWD recruited the participants and acquired the clinical data. JWD and NJT acquired the MRI. CAS performed the analysis and drafted the manuscript. All authors contributed to data analysis, drafting and revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

PWJ is employed as a Global Medical Expert for GlaxoSmithKline. JWD reports grants from the British Lung Foundation, during the conduct of the study and has received personal fees and travel support unrelated to the content of this manuscript from Chiesi, Boehringer Ingelheim & NAPP pharmaceutical. DRB reports grants from National Institute for Health Research, during the conduct of the study. The authors report no other conflicts of interest in this work.

References


27. Murray AD, Staff RT, Shenkin SD, Deary IJ, Starr JM, Whalley LJ. Large vessel disease and cerebral small vessel disease: a prospective follow-up study. *Stroke* 2011;42:133–138. doi:10.1161/STROKEAHA.110.594267


40. Lillenthal JL, Riley RL. On the determination of arterial oxygen saturations from samples of “capillary” blood. *J Clin Invest* 1944;23(6):904–906. doi:10.1172/JCI11565


Supplementary materials
Inclusion/Exclusion criteria

Clinical measures
Hospital Anxiety and Depression Scale (HADS)
The HADS is a 14-item self-report questionnaire comprising two 7-item subscales measuring anxiety and depression. It was originally developed as a clinical screening tool for use in a general medical outpatient setting and so explicitly excludes items that might be confounded by somatic aspects of illness or serious mental disorders.¹

Montreal Cognitive Assessment (MoCA)
The MoCA is a brief 30-item cognitive assessment tool designed to be sensitive to mild cognitive impairment in individuals presenting with subjective cognitive complaints.² The MoCA has previously been applied to chronic obstructive pulmonary disease (COPD) cohorts where it has been shown to be sufficiently sensitive to detect mild cognitive impairment in COPD patients with moderate-severe disease.³

Image acquisition
All images were acquired with a 3-Tesla Siemens Magnetron Skyra MRI scanner equipped with a 32-channel head coil with a maximum gradient strength of 45 mT/m. Sagittal T1-weighted 3D volume (T1W) images were acquired using a magnetization prepared rapid gradient echo sequence (TE=2.25 ms, TR=1800 ms, TI=800 ms, flip angle 9°, 169 contiguous sagittal slices with a 0.9 mm³ isotropic voxel dimension and field-of-view of 225 mm×240 mm×180 mm). Axial fluid-attenuated inversion recovery (FLAIR) was acquired using an inversion recovery sequence (TE=126 ms, TR=11,000 ms, TI=2690 ms, flip angle=150°, with 60 contiguous slices, voxel dimension of 0.7 mm×0.7 mm×3 mm and field-of-view of 201.25 mm×230 mm×180 mm). Diffusion tensor images (DTI) were acquired using an echo-planar imaging sequence with opposite phase-encode polarities (TE=76 ms, TR=6000 ms, flip angle =90°, 55 contiguous axial slices with a voxel dimension of 2.0 mm×2.0 mm×2.5 mm and field-of-view of 192 mm×192 mm×137.5 mm). For each phase-encode polarity, 8 volumes were acquired without diffusion sensitization and 60 with non-collinear diffusion gradients applied.

Image processing
Tissue macrostructure
The T1W images were re-sampled to 1 mm³ isotropic. The FLAIR was affine-registered to the T1W images using Advanced Normalisation Tools⁴ and a semi-automatic procedure used to segment the T1W images into supra-tentorial gray matter, white matter and cerebrospinal fluid tissue probability maps. White matter hyperintensities (WMHs) were segmented using the combined image intensities from the T1W and FLAIR images, then binarised at a manually determined threshold (i.e., dichotomized so that 1=WMH and 0=non-WMH). This process is described in full in Spilling et al, 2017 and Lambert et al, 2015.⁵,⁶ Tissue volumes were quantified by integrating the values within each tissue segmentation and normalizing for head size – calculated as a percentage of total intracranial volume (gray matter + white matter + cerebrospinal fluid). Additionally, these tissue segmentations were used to define regions of normal-appearing white matter (NAWM) on the DTI (see below).

Table S1 Inclusion and exclusion criteria

<table>
<thead>
<tr>
<th>Inclusion criteria</th>
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<tbody>
<tr>
<td>Informed consent</td>
<td>Resting SaO₂&lt;92% on room air</td>
</tr>
<tr>
<td>Aged 40–85 years</td>
<td>Known alpha-1 anti-trypsin deficiency</td>
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<tr>
<td>&gt;10 pack year history</td>
<td>Pregnancy or lactating</td>
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<tr>
<td>FEV₁/FVC&lt;70% (COPD only)</td>
<td>Ischaemic heart disease</td>
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<tr>
<td>Exacerbation free for 4 weeks (COPD only)</td>
<td>Cerebrovascular disease</td>
</tr>
<tr>
<td>FEV₁/FVC&gt;70% (Non-COPD smoking controls only)</td>
<td>Uncontrolled hypertension</td>
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<tr>
<td>Diabetes mellitus</td>
<td>Hepatic failure</td>
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<td>Hepatic failure</td>
<td>Obstructive sleep apnoea</td>
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<td>Non-cured tumors</td>
<td>Non-cured tumors</td>
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<td>Psychiatric disease that would impact on consent or compliance</td>
<td>Psychiatric disease that would impact on consent or compliance</td>
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<td>Neurological disease</td>
<td>Neurological disease</td>
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<td>Known history of dementia</td>
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<td>Current or past alcohol/drug abuse</td>
<td>Current or past alcohol/drug abuse</td>
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<td>Visual or hearing impairment that precluded neuropsychological assessment</td>
<td>Visual or hearing impairment that precluded neuropsychological assessment</td>
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<td>Contraindications for MRI</td>
<td>Contraindications for MRI</td>
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<td>Incidental findings on MRI</td>
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Abbreviations: FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity; COPD, chronic obstructive pulmonary disease; MRI, magnetic resonance imaging.


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Tissue microstructure

The DTI data were corrected for movement artifacts, eddy-current distortions, and susceptibility-induced local gradients using FSL’s (FMRIB Software Library, version 5.0.6) “eddy”.

The diffusion tensor model was fitted at every voxel within the DTI using FSL’s (FMRIB Software Library, version 5.0.6) “dtifit”, the skull removed using FSL’s (FMRIB Software Library, version 5.0.6) Brain Extraction Toolbox and mean diffusivity (MD) and fractional anisotropy (FA) maps calculated from the DTI, indicating the local magnitude and directionality of diffusion, respectively.

T1W images were aligned to the DTI data using the boundary-based registration procedure implemented in FSL’s (FMRIB Software Library, version 5.0.6) “epi-reg” script.

This transformation was applied to the T1W tissue segmentations and binary WMH map (using trilinear interpolation) to align them with the DTI. The WMH map was re-binarised at 0.5 creating a WMH mask. These segmentations were used to define the probability of the DTI voxels belonging to each tissue-type. Voxels were considered to belong to the supra-tentorial NAWM where the probability of belonging to the white matter was higher than for any other tissue-type providing that they were not included within the WMH mask.

Normalized histograms (i.e. probability density functions) of FA and MD values within the NAWM were constructed using 100 equal bins ranging in value from 0 to 1 for FA and 0 to 2×10⁻⁴ mm²/s for MD. The median and peak height values were used to characterize the distribution of these histograms.

References