PiSZ alpha-1 antitrypsin deficiency (AATD)
Green, Clara; Vayalapra, Sushanth; Hampson, Judith; Mukherjee, Deyashini; Stockley, Robert; Turner, Alice

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PiSZ alpha-1 antitrypsin deficiency (AATD): pulmonary phenotype and prognosis relative to PiZZ AATD and PiMM COPD

C E Green,1 S Vayalapra,1 J A Hampson,1 D Mukherjee,2 R A Stockley,1 A M Turner1,3

ABSTRACT

Introduction The PiSZ genotype results in less severe deficiency of alpha-1 antitrypsin (AAT) than PiZZ. Less is known about phenotypic and prognostic features.

Methods We studied 699 PiZZ, 126 PiSZ and 316 PiMM patients. All AAT deficiency (AATD) patients were augmentation naive. PiSZ were compared with PiZZ patients for clinical phenotype at baseline including CT findings, smoke exposure, progression of lung disease and survival. Similarly, PiSZ patients diagnosed as a result of investigation for possible lung disease (lung index cases) were compared with PiMM. Multivariable analytical techniques and matching (PiSZ to PiZZ) were employed to account for demographic differences.

Results Pack-years smoked and FEV1 exhibited a negative correlation in PiZZ and ZZ patients (both r=-0.43), with emphysema and COPD occurring more commonly in PiZZ patients at <20 pack-year exposure. In multivariable analyses, PiSZ patients were less likely to have emphysema (p<0.01) and had better survival than PiZZ (p=0.017), but lung function decline did not differ significantly. 42% of PiSZ patients had upper-zone-dominant emphysema on CT scan. Analyses of AAT level confirmed a critical threshold at 11 μM, particularly with regard to phenotypes classical of PiZZ AATD.

Significant baseline differences suggested that PiSZ had presented earlier to health services than PiMM. Once this was accounted for, risk of emphysema did not differ between PiSZ and PiMM although survival was lower in PiMM patients (p<0.01).

Conclusions PiSZ patients are less susceptible to cigarette smoke than PiZZ. The pattern of emphysema may be similar at diagnosis to usual COPD.

INTRODUCTION

Alpha-1 antitrypsin deficiency (AATD) results from several deficiency alleles, with severe deficiency occurring in Z allele homozygotes (PiZZ), or carriers of null alleles leading to absence of AAT1 and is associated with early-onset emphysema.2 The S allele leads to a milder deficiency, and there has been considerable debate about whether heterozygous PiSZ patients are at increased risk of lung disease, largely due to the confounding factor of acquisition bias in some prior work. A meta-analysis concluded that PiSZ patients exhibit a risk of COPD over three times that of a normal individual;5 however, this result was influenced by one study showing a large increase in risk.6

Previous studies of PiSZ patients are summarised in online supplementary table 1. Epidemiological studies suggest that there are >500 000 PiSZ individuals in Europe,7 who typically exhibit an AAT level less than half that of a normal (PiMM) person.8 An interaction between AAT and cigarette smoke exposure occurs in murine models, where emphysema develops more rapidly.9 This, together with the available epidemiology, suggests that PiSZ patients should have a lung disease risk lying between PiMM and PiZZ patients, given the same level of smoke exposure. Consistent with this, we have shown in a relatively small cohort (n=63), matched for smoking status, that PiSZ patients exhibit less severe lung disease than PiZZ.8 Even low levels of smoke exposure may be deleterious to lung function in PiZZ patients; however, an important threshold may occur at 20 pack-years, above which associations between FEV1 and pack-years are less apparent.10 Whether this differs in PiSZ patients is unknown, and little is known about their prognosis. We sought to confirm the pattern of lung disease exhibited by PiSZ patients, explore the relationship between cumulative smoke exposure, AAT level and severity of disease, and compare prognosis between PiSZ, PiZZ and PiMM patients.
METHODS

Subjects

The UK AATD registry (ADAPT) was established in 1996 and
has been described in detail elsewhere.10 Briefly, patients
undergo annual lung physiology, biological sample collection
and clinical assessment. At enrolment most patients have a chest
CT scan; quantitative CT analysis was undertaken by density
mask analysis of 5 mm CT scan slices; voxel indices (VI) at
−910 (the proportion of lung voxels below −910 Hounsfield
Units (HU)) and the ratio between the upper and lower zones
calculated.11 Lung function decline was determined on all
patients with ≥3 years’ follow-up; methods for calculation of
decline have been described in our previous work.12 All patients
with PiSZ and PiZZ genotypes were selected. Those diagnosed
due to investigation for symptoms/signs of lung disease were
termed lung index cases.

PiMM patients with usual COPD comprised all patients in
the West Midlands COPD cohort (WMCC; described in our
previous published work13) and all patients from the Chronic
Diseases Resource Centre (CDRC; a University of Birmingham
COPD patient registry, identified from local clinics). The CDRC
has superseded the WMCC and has assessment procedures
similar to the UK AATD registry. Data on CT density and
decline in lung function were not available for PiMM patients,
as referring centres used a variety of CT protocols, and had
either not been followed up for sufficient time or had insuffi-
cient lung function data to calculate decline.

Statistical analysis

All analyses were carried out in SPSS (SPSS V20; IBM, USA).
Univariable analyses compared PiSZ and PiZZ patients for
demographics, smoking status, index status, occupational expos-
ure (assessment methods described in our previous work12),
comorbidities, lung function, CT densitometry (where available)
survival and lung function decline. Analyses were performed
using χ2 tests for categorical variables and Mann-Whitney U
tests for continuous variables, substratifying by index status,
smoking status and presence of COPD for analyses pertaining to
decline. Co-variables for logistic, linear and Cox regression ana-
lyses were selected if p<0.1 in univariable analysis and no sig-
nificant collinearity with another included variable. Prior
literature on influences on mortality14 and decline,12 15 16 in
AATD were used to aid prioritisation of co-variables where
necessary. All results from the multivariable analysis are reported
as two-tailed; significance was taken at p<0.05. Since there
were demographic differences between PiSZ and PiZZ patients,
for which subgroup analyses and statistical adjustment in regres-
sion analysis might be imperfect, an attempt at a matched ana-
lysis was also made (see online supplementary data). Within the
whole AATD group, the relationship of AAT level to clinical
phenotype and decline was conducted using similar techniques.
Finally, univariable analyses were undertaken comparing lung
index PiSZ and PiMM patients for demographics, smoking
status, clinical phenotype and survival. Results guided logistic
and linear regressions similar to the PiSZ versus PiZZ
comparisons.

RESULTS

Comparison of PiSZ and PiZZ patients

Table 1 shows patients’ characteristics and univariable analyses.
PiZZ patients were more likely to be lung index cases and ex
smokers, had higher rates of emphysema, bronchiectasis and
chronic bronchitis, worse lung function (in all parameters), lower
AAT levels and worse survival. Almost all (99.8%) PiZZ patients
and 11.3% PiSZ patients had AAT<11 μM. CT scans were available
on 37 non-index and 64 index PiSZ patients (index vs non-
index characteristics; see online supplementary table 2). Smoking
appeared to be a greater influence on clinical phenotype than
PiSZ versus PiZZ genotype (smoke exposed vs never smokers’
characteristics; see online supplementary table 3).

Direct correlation between FEV1% predicted and pack-years
occurred up to a break point, which appeared similar to the
published 20 pack-year threshold in PiZZ patients, though possi-
bly higher in PiSZ (figure 1A, B). The proportion of emphy-
sema and COPD cases occurring below 20 pack-years was greater
in PiZZ patients (figure 1C, D). There was no difference in
prevalence of common comorbidities (IHD, osteoporosis,
anxiety, depression), age, pack-years smoked or occupational
exposure risks (all p>0.2). Multivariable logistic regressions,
stratified for method of ascertainment of AATD, including
smoke exposure and age as covariates, showed that emphysema
was less common in lung index and family screened PiSZ
patients (OR 0.15 and 0.03; p=0.003 and <0.001, respect-
ively), while rates of chronic bronchitis and bronchiectasis were
similar between groups (all p>0.2). In linear regressions strati-
ﬁed for method of ascertainment of AATD and adjusting for
smoking status and age, lung index PiSZ patients had signifi-
cantly better lung function compared with PiZZ (all parameters
shown in table 1, p<0.05), while non-index PiSZ patients were
better in most parameters (residual volume and total lung cap-
acity no difference, other parameters p<0.05). Matched ana-
lyses were no different (see online supplementary results).
Differences in emphysema between PiSZ and PiZZ patients
were more marked in the lower zones (figure 2); 42.9% of PiSZ
patients scanned had upper-zone-dominant emphysema com-
pared with 14.1% of PiZZ (both p<0.01).

Sufficient physiological data to calculate decline were available
in 68 PiSZ and 514 PiZZ patients, with a mean of five measures
per PiZZ and four per PiSZ patient. 22.5% of PiZZ patients
had COPD with FEV1 <30% predicted (n=116), 33.5% had
FEV1 30–50% (n=172), 25% (n=129) had COPD with FEV1
>50% predicted and the remainder did not have COPD at the
start of the decline calculation period (n=97). In the PiSZ
patients, these ﬁgures were 11.3%, 8.1%, 17.7%, 29% and
33.9% (n=8, 5, 12, 20 and 23), respectively. Decline in FEV1
in some subgroups of interest is shown in table 2. Median
decay was greatest when baseline FEV1 was 50–80% predicted
(PiZZ: −56.3 mL/year, PiSZ: −65.3 mL/year, p=0.707). There
was no difference in annual decline of FEV1 or gas transfer
(DLCO and KCO) %predicted when analysed as categories
(ﬁgure 3) or continuous variables. Regression analyses stratiﬁed
for smoking status, presence of COPD and index status, adjust-
ing for baseline lung function and age did not inﬂuence this
similarity (all p>0.18 for smoking status, p>0.33 for COPD,
p>0.77 for index status), nor did matching (see online supple-
imentary results). Cox regression analyses, including age,
smoking status, index status and baseline FEV1 as co-variables,
showed significantly better survival in PiSZ patients (ﬁgure 4A).

We then explored the importance of AAT level with regard to
susceptibility to smoke, subsequent development of lung disease
and decline. When considered as a continuous variable, AAT
level had no signiﬁcant impact on logistic regressions seeking
associations of emphysema, chronic bronchitis and bronchiec-
tasis. However, a level of ≥11 μM associated with development
of emphysema (p<0.001), an UZ/LZV1 suggestive of lower-
zone-dominant disease (p<0.001) and chronic bronchitis
(p<0.001). There was a strong trend toward interaction between
AAT ≥11 μM and pack-years smoked with regard to risk of

Table 1  Clinical characteristics of patients

<table>
<thead>
<tr>
<th>Feature</th>
<th>PIZZ n=699</th>
<th>PISZ n=126</th>
<th>PISZ lung index cases N=64</th>
<th>PIMM n=316</th>
<th>p Value</th>
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<tbody>
<tr>
<td>Male</td>
<td>404 (57.8)</td>
<td>71 (56.3)</td>
<td>38 (59.4)</td>
<td>182 (57.7)</td>
<td>0.717</td>
</tr>
<tr>
<td>Age</td>
<td>50.8 (16.4)</td>
<td>53.5 (20.0)</td>
<td>55.4 (11.5)</td>
<td>68.4 (13.6)</td>
<td>0.021</td>
</tr>
<tr>
<td>Smoking status</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ex smoker</td>
<td>510 (73.0)</td>
<td>56 (44.4)</td>
<td>37 (57.8)</td>
<td>192 (60.0)</td>
<td>0.004</td>
</tr>
<tr>
<td>Never smoked</td>
<td>161 (23.0)</td>
<td>43 (34.1)</td>
<td>15 (23.4)</td>
<td>3 (0.9)</td>
<td></td>
</tr>
<tr>
<td>Current smoker</td>
<td>28 (4.0)</td>
<td>27 (21.4)</td>
<td>12 (18.8)</td>
<td>121 (39.0)</td>
<td></td>
</tr>
<tr>
<td>Pack-years smoked</td>
<td>13.5 (22.5)</td>
<td>13.0 (35.4)</td>
<td>26.2 (3.0)</td>
<td>44.2 (32.5)</td>
<td>0.402</td>
</tr>
<tr>
<td>Reason for diagnosis</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Lung disease</td>
<td>532 (76.2)</td>
<td>77 (61.0)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Family screening</td>
<td>118 (16.9)</td>
<td>38 (30.5)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Liver disease</td>
<td>11 (1.6)</td>
<td>1 (0.8)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other reason</td>
<td>38 (5.4)</td>
<td>10 (7.9)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>COPD</td>
<td>587 (84.0)</td>
<td>66 (52.4)</td>
<td>43 (67.2)</td>
<td>316 (100)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Emphysema</td>
<td>528 (75.5)</td>
<td>46 (35.6)</td>
<td>35 (54.7)</td>
<td>257 (81.3)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Chronic bronchitis</td>
<td>249 (35.6)</td>
<td>31 (24.6)</td>
<td>18 (28.1)</td>
<td>198 (62.8)</td>
<td>0.034</td>
</tr>
<tr>
<td>Bronchiectasis</td>
<td>189 (27.0)</td>
<td>22 (17.4)</td>
<td>11 (17.2)</td>
<td>96 (30.3)</td>
<td>0.041</td>
</tr>
<tr>
<td>AAT level</td>
<td>4.0 (2.1)</td>
<td>14.4 (4.4)</td>
<td>14.9 (3.7)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FEV1/pp</td>
<td>47.9 (45.6)</td>
<td>94.5 (51.3)</td>
<td>76.5 (2.6)</td>
<td>46.4 (26.0)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>FEV1/FVC (%)</td>
<td>38.2 (26.4)</td>
<td>67.5 (37.5)</td>
<td>55.0 (2.7)</td>
<td>43.3 (20.0)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TLCpp</td>
<td>115.3 (21.9)</td>
<td>107.4 (18.3)</td>
<td>109.2 (1.9)</td>
<td>103.0 (20.2)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RVpp</td>
<td>119.3 (54.9)</td>
<td>96.9 (46.3)</td>
<td>107.9 (4.6)</td>
<td>119.8 (53.8)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>DLOpp</td>
<td>63.1 (26.0)</td>
<td>86.3 (34.2)</td>
<td>76.5 (3.1)</td>
<td>48.6 (23.9)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>KCOpp</td>
<td>64.3 (28.8)</td>
<td>89.0 (29.5)</td>
<td>78.1 (3.0)</td>
<td>59.0 (29.6)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Follow-up time (years)</td>
<td>10.4 (5.3)</td>
<td>7.0 (8.5)</td>
<td>7.8 (6.6)</td>
<td>3.2 (3.0)</td>
<td>0.021</td>
</tr>
<tr>
<td>Deceased</td>
<td>172 (24.6)</td>
<td>10 (7.9)</td>
<td>7 (10.9)</td>
<td>91 (28.8)</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Data are shown as n (%) or median (IQR). In PISZ index cases, data are shown as mean (SE) due to normal distribution.

AAT, alpha-1 antitrypsin; DLCO, carbon monoxide diffusing capacity; KCO, carbon monoxide transfer coefficient; RV, residual volume; TLC, total lung capacity.

Figure 1  Relationship between pack-years smoked and lung function in alpha-1 antitrypsin deficiency. There was a linear relationship between baseline FEV1 and pack-years up to a threshold of 30 pack-years in PiSZ patients, which then disappeared (A; r=−0.43 at <30 pack-years, r=0.02 at ≥30 pack-years). The relationship was similar in PiZZ patients, but with a lower pack-year threshold (B; r=−0.43 at <20 pack-years, r=0.04 at ≥20 pack-years). Illustrative smoothed regression lines are shown on the graphs. Both emphysema (C) and COPD (D) were more common at exposures <20 pack-years in PiZZ patients.

chronic bronchitis (p=0.09), but not other phenotypes. Furthermore, interaction between AAT level and pack-years smoked was significant in a multivariable general linear model for FEV1 decline (one-tailed test p=0.026), whereas AAT level alone and PiSZ’ pack-years interaction were not (both one-tailed p=0.150).

**Comparison with usual COPD**

The primary reason for inclusion of PiMM patients was to see whether their phenotype was similar to PiSZ, hence univariable analyses were restricted to PiSZ index cases, as this eliminated ascertainment differences. Logistic regressions adjusting for age, smoke exposure and baseline FEV1 (different in univariable analyses; [table 1](#)) demonstrated that PiSZ patients had a similar risk of emphysema (p=0.274) and bronchiectasis (p=0.344), but lower risk of chronic bronchitis (OR 0.36 (0.18–0.72); p=0.016). Cox regression confirmed better survival in PiSZ patients ([figure 4B](#)).

**Discussion**

We have confirmed that PiSZ patients have a lower risk of lung disease and are less susceptible to effects of cigarette smoke than PiZZ individuals, exhibiting a clinical phenotype, which is in many cases similar to usual PiMM COPD. Interpreting the prognostic data is more difficult due to issues of power and baseline differences between groups.

**Pulmonary phenotype**

This study is the largest to date on PiSZ AATD. In our earlier study, lung function and CT scan appearances concurred with those reported here; many PiSZ patients exhibited no emphysema, and many who did had upper-zone-dominant disease.8 This agrees with data from an earlier, small, British PiSZ cohort17 and with much of the past PiSZ literature. The expected number of UK PiSZ patients exceeds PiZZ, yet there are far fewer PiSZ known to the registry. This may reflect ascertainment bias since AAT testing possibly occurs more frequently in patients with COPD exhibiting classical AATD lung disease (ie, lower-zone-dominant emphysema). A missed diagnosis could have consequences since there is specific treatment for AATD lung disease in the form of AAT augmentation (available in the USA and many European countries, albeit not yet in the UK), which undiagnosed patients would not receive. Most international AATD guidelines specify that their scope is PiZZ patients or those with a circulating AAT level typical of PiZZ, and recommend the use of augmentation in the presence of emphysema and a specified level of FEV1.18 19 This would exclude almost 90% of our PiSZ patients on the basis of AAT level.6 The main effect of augmentation is on progression of emphysema, best measured by CT densitometry,20 although effects on FEV1 decline have been reported in non-randomised studies.21 Progression of emphysema on augmentation varies according to lung zone,11 hence augmentation might benefit PiSZ patients who exhibit lower-zone-dominant disease and low AAT levels (equivalent to PiZZ). Our data examining the threshold AAT level of 11 μM support this, although specific studies in PiSZ patients would be required to determine this with confidence.

We also confirmed that PiSZ patients are less susceptible to smoke. A direct relationship between FEV1 and smoke exposure occurred below 30 pack-years in PiSZ and 20 pack-years in PiZZ patients. The reasons for this are discussed elsewhere,22 one of which is a ‘floor effect’, in which lung function has dropped sufficiently by the threshold that in many patients either no further decline occurs or they die. A smaller amount of ‘resistant’ smokers is also seen, adding to the lack of correlation at higher smoke exposures. While the data for the specific threshold were weak, the proportion of PiSZ patients developing emphysema or COPD at exposures <20 pack-years was also lower, supporting its existence. Moreover, the American NHLBI cohort reported the importance of smoking in disease development in PiSZ patients with AAT <11 μM, alongside less lung function impairment than in PiZZ, thus supporting our data.23 A difference in the threshold at which smoke susceptibility occurs between PiSZ and ZZ patients is consistent with current understanding of AAT–neutrophil elastase (NE) interactions, whereby the area of obligate damage is exponentially related to AAT level, rising markedly at <11 μM,23 and therefore leading to a critical area of lung damage more quickly in PiZZ. Interestingly, among the PiSZ never smokers clinically significant
emphysema was uncommon, implying that in the absence of smoke it is not a risk factor—similar to PiMZ.24

We then compared PiSZ and MM patients to test our hypothesis that there would be a degree of phenotypic overlap, perhaps sufficient to make the two groups indistinguishable, thus accounting for missed diagnoses (relative to PiZZ). The potential impact of individuals with AATD diagnosed through family screening, who were not present in the usual COPD

Figure 3  Decline in lung function in alpha-1 antitrypsin deficiency. The graphs show the proportion of patients with lung function that did not deteriorate, decreased by up to 1% of predicted (ie, fell from 50% predicted in year 1 to ≥49% predicted in year 2), between 1% and 2% predicted and >2% predicted per year. (A) FEV1, (B) DCLO and (C) carbon monoxide transfer coefficient (KCO). There were no differences in the proportion of patients in each group (FEV1, p=0.67; DCLO, p=0.22; KCO=0.60).

Figure 4  Cox regression demonstrating survival. (A) PiSZ (dashed line) versus PiZZ (solid line); analysis adjusted for baseline FEV1, age, index status and smoking. PiSZ patients had better survival (p=0.017; log rank). (B) Lung index PiSZ (dashed line) versus PiMM (solid line); analysis adjusted for baseline FEV1, age and pack-years smoked. PiSZ patients had better survival (p<0.001; log rank).
group, meant that statistical analyses compared lung index cases—risks of emphysema were similar, after adjustment for smoke exposure and age. This is consistent with PiSZ patients outside our cohorts being diagnosed less readily.

Prognosis

Our data implied that disease progression in PiSZ patients might be similar to PiZZ. This result was unexpected; PiSZ patients were generally better at baseline, which we have shown to relate to more rapid FEV1 decline.\(^{10,16}\) While PiZZ patients were worse, which we have shown to relate to more rapid gas transfer decline.\(^{10,16}\) We therefore expected differences to occur in raw decline values, which would be less apparent after adjustment for baseline lung function and smoke exposure. However, raw values were similar and remained so regardless of the method by which we attempted to control for baseline differences. This may be explicable if AAT level is more critical than PiSZ genotype; consistent with this our data implied that AAT level, rather than PiSZ genotype, interacted with smoke exposure in the lung function decline models. AAT level does vary among PiSZ patients, presumably due to modifying effects of other genes or environmental factors, which is one reason why current guidelines for treatment take this feature into account.

No prior PiSZ studies have reported FEV1 decline (see online supplementary table 1); however, there are data available on decline in PiZZ, where loss of FEV1 was generally 60–80 mL/year.\(^{13} 23–28\) This is somewhat higher than the median in our PiZZ patients. Notably, the largest prior PiZZ study\(^{23}\) (see online supplementary table 4) exhibited both starting FEV1 (49% predicted) and FEV1 decline similar to many of our PiZZ patients (54 mL/year vs 56.3 mL/year in our FEV1 50–80% subgroup). Our lower median is explicable by cohort characteristics. There are two groups in whom FEV1 decline is known to be lower—those with FEV1 <30% predicted\(^{10}\) or with no emphysema on their CT, who comprise 22.5% and 26.3% of our AATD cohort, respectively. High rates of patients with FEV1 <30% account for the low decline seen in our PiZZ current smokers, whose minimum FEV1 was just 12%, thus had no lung function to lose. Furthermore, our method of determining decline may be more accurate compared with previous studies, which calculated decline from just two data points,\(^{23}\) or over only 2 years in some patients.\(^{28}\) Thus we feel our decline data are accurate and comparable to other published data, once our cohorts’ characteristics are taken into account. However, we acknowledge that a survivor bias, and relatively small numbers, in the decline analyses may have limited power to detect true differences between PiSZ and ZZ patients. While we tried to adjust for baseline differences in lung function (eg,11.3% of patients had FEV1 <30% predicted in the PiZZ decline group compared with 22.5% of the PiZZ), statistical adjustment could have been incapable of fully accounting for differences. To overcome this, we attempted matching (see online supplementary data), which again showed no difference in decline. However, high smoke exposures in some PiSZ made matching imperfect also. Further studies of decline in PiSZ patients are therefore indicated.

Despite apparently similar lung function decline, subsequent survival was better in PiSZ compared with PiZZ. There are several possible reasons for these apparently inconsistent results. First, declining lung function may not be a good predictor of survival; we have recently shown that declining CT densitometry may be a better measure.\(^{29}\) Alternatively, it may be that neither statistical adjustments nor matching were capable of fully compensating for baseline differences, such that better survival reflects their higher starting point. One factor unlikely to have influenced the result is comorbidity, which did not differ between PiSZ and ZZ patients. Survival was worse in usual COPD than PiSZ patients, probably due to higher comorbidity burden attendant on greater smoke exposure, although we had insufficient data to test this hypothesis. The proportion of patients surviving 5 years was about 70% in usual COPD and between 90% and 95% in both PiSZ and ZZ individuals (figure 4). Since our AATD cohort is a national centre, and many patients travel some distance to be seen, there may be a survivor bias, which could have inflated the difference seen between PiSZ and MM. If present, it would have affected PiSZ and ZZ patients equally, hence their comparative survival result is more robust.

Strengths and limitations

The main strengths of our study are the large number of AATD patients, depth of clinical phenotyping and length of follow-up. Ideally we would have compared emphysema zone and lung function decline between PiSZ and MM, as we did for PiSZ versus ZZ, but were limited by lack of data. However, its utility for clinical management would be small since optimal management of COPD should occur in both usual COPD and COPD due to PiSZ AATD. The main difference would emerge if augmentation were available, for which the more informative longitudinal PiSZ comparison is with PiZZ patients. Further work to clarify whether differences in neutrophil function occur between PiSZ and MM patients, as they do between PiZZ and MM,\(^{30}\) whether NE activity differs\(^{31,32}\) between groups and the utility of augmentation in PiSZ patients may be valuable follow-up studies.

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