Prevalence and phenotype of subjects carrying rare variants in the Italian registry for alpha1-antitrypsin deficiency

I Ferrarotti, J Baccheschi, M Zorzetto, C Tinelli, L Corda, B Balbi, I Campo, E Pozzi, G Faa, P Coni, G Massi, G Stella, M Luisetti

LETTER TO JMG

A
lpha1-antitrypsin deficiency (AATD) is a genetic condition associated with an increased risk of developing chronic obstructive pulmonary disease (COPD) early in life and, to a lesser extent, liver disease.1 Significant advances have been made during the last decades in understanding its epidemiology and it has been recently suggested that AATD is one of the commonest inherited disorders not only in Caucasians but also among other ethnic groups worldwide.2, 3 Although AAT is a highly pleomorphic glycoprotein, with approximately 100 variants having being identified,4 two major deficient variants, namely Z and S, account for most cases of AATD, since the vast majority of such individuals carry the PI*ZZ or PI*SZ genotype, coding for approximately 15% and 25%, respectively, of normal AAT plasma levels. The establishment of international registries, including large series of AATD individuals,4-6 has allowed not only better definition of the epidemiology of AATD, but also more precise definition of the associated clinical phenotypes.4-8

Nevertheless, there are at least 30 AAT alleles other than the PI*Z and the PI*S alleles which are associated with significantly reduced or absent plasma AAT levels.4 Given the extreme rarity of such variants, often described in the literature as single case reports, little is known about their epidemiology and even less is known about the associated clinical phenotypes.

The Italian Registry for Severe AATD was established in 1996 as a result of a nationwide screening programme sponsored by the two major Italian scientific respiratory societies.9-11 Although Italy is considered a country with a medium-low prevalence of AATD (mean PI*Z gene frequency: 0.0013),11 the programme succeeded in identifying a relatively large cohort of AATD individuals. During the development of the screening program, we noticed that, in addition to the groups of AATD individuals carrying the PI*ZZ and PI*SZ genotypes, there was an unexpectedly large group of subjects carrying at least one rare AATD allele. We therefore decided to study this group of subjects, focusing particularly on characterising their clinical phenotypes.

METHODS

Screening programme

The targeted screening programme, based on dried blood spots, has already been described in detail.11 Briefly, paper filters and questionnaires were distributed to respiratory physicians throughout the country. Recommendations for AATD screening were the presence of the following: early-onset COPD, familial clustering of COPD, reduced levels of α1-globulins on electrophoresis, serum levels of AAT <80 mg/dl (nephelometry) or <150 mg/dl (immunodiffusion), or a family history of AATD. Paper filters containing the blood spots were shipped to the Central Phenotyping Laboratory in Rome and submitted to isoelectric focusing. In the case of an abnormal isoelectric focusing pattern, the referring physician was asked to ship a serum sample and a frozen whole blood sample. If the abnormal isoelectric focusing pattern was confirmed, the sample was then investigated at a molecular level. The subject’s demographic and clinical data were retrieved from questionnaires filled in by the referring physicians and shipped together with the specimens. It was recommended that the diagnosis of COPD follow international guidelines.12 All subjects gave their consent to undergo the genetic investigation, which was approved by the ethical committees of the institutions involved.

Key points

- Most subjects affected by α1-antitrypsin deficiency (AATD) carry the PI*ZZ or PI*SZ genotype. Nevertheless, there are at least 30 AAT alleles other than PI*Z or PI*S associated with reduced or absent plasma AAT levels. Little is known about their epidemiology or associated clinical phenotypes.
- Over 98 months, 2922 subjects enrolled by the Italian Registry for AATD, which conducts a screening programme based on dried blood spots, were screened. A total of 155 subjects with severe AATD were identified (132 index cases), together with 152 individuals with intermediate AATD (84 cases).
- Among subjects with severe AATD, we recorded an 11% prevalence of deficient genotypes other than the common PI*ZZ or PI*SZ (15 out of 132 deficient index subjects identified). Among subjects with intermediate AATD, we recorded a 15% prevalence of deficient genotypes other than the common PI*MZ (13 out of the 84 index cases).
- As the cohort of subjects carrying rare AATD variants was relatively large, we were able investigate their characteristics in terms of associated clinical phenotypes, pulmonary lung function, smoking habit, and geographic distribution. We found that these rare variants had a special position within the spectrum of genotype-phenotype correlations in α1-antitrypsin deficiency.
- The prevalence of rare α1-antitrypsin variants found in Italy is the highest so far recorded worldwide and, interestingly, occurs in a country with a medium-low prevalence of the common PI*ZZ genotype.

Abbreviations: AATD, alpha1-antitrypsin deficiency; COPD, chronic obstructive pulmonary disease; OR, odds ratio; 95% CI, 95% confidence interval
Detection of AATD variants

Genomic DNA was extracted from whole blood cells using the standard technique. The S and Z variants were genotyped by a commercially available amplification-reverse hybridisation test kit (Symbiosis, Cocconato, Italy) and by PCR-RFLP using TaqI as the restriction enzyme. The genomic DNA was sequenced after PCR amplification of all coding exons (II-V). All sequencing products were obtained using the ABI Prism BigDye Terminator Cycle Sequencing Ready Reaction kit (Applied Biosystems, Arrington, UK) and were analysed in an automatic ABI 377 DNA sequencer.

Statistical analysis

Separate files were created for the different genotypes. Subjects homozygous or compound heterozygous for deficiency variants were assigned to the severe AATD group, whereas individuals heterozygous for a deficiency allele and the normal M allele were assigned to the intermediate AATD group. Shapiro-Wilk’s test was used to test the normal distribution for quantitative variables and data values are presented as mean (SD) values. Comparisons between means were performed with analysis of variance, using the Scheffé test for post-hoc comparison. Associations between genotypes and disease status were compared with Fisher’s exact test. To detect associations between FEV1/FVC (forced expiratory volume in 1 s/forced vital capacity), smoke, and genotype, univariate analyses were carried out using logistic regression, followed by multivariate analyses; results are presented as unadjusted and adjusted odds ratios (OR) with 95% confidence intervals (95% CI). A p value <0.05 was considered to indicate statistical significance; all tests were two-sided. Analyses were performed with STATA, release 7.0 (Stata, College Station, TX, USA) and with STATISTICA for Windows (StatSoft, Bedford, UK, 2002).

RESULTS

During the 98 months from February 1996 to April 2004, 2922 subjects were screened, thanks to more than 250 physicians throughout the country who shipped at least one paper filter. A total of 155 individuals (5.3%) with severe AATD were detected (table 1). Of these, 132 were index cases: index cases: 84 (29%) and 21 subjects with intermediate deficiency, resulting from the combination of a rare allele with the normal M variant (0.7% of all those screened; index cases: 13; 0.4%). The rare allele frequency detected was 0.077, taking into account subjects with both severe and intermediate AATD.

The genotyping details of the 37 individuals carrying at least one rare AATD variant are given in table 2. In this series we found 16 individuals carrying the rare variant in a homozygous fashion or in compound heterozygosity with the common Z deficient variant. The M/R (M/rare) subset consists of subjects with intermediate AATD (one rare AAT variant in combination with the normal M allele). Non-index individuals were identified during family screening.

### Table 1

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Number of subjects</th>
<th>Genotype</th>
<th>Number of subjects</th>
</tr>
</thead>
<tbody>
<tr>
<td>PI*ZZ</td>
<td>114</td>
<td>PI*SZ</td>
<td>25</td>
</tr>
<tr>
<td>R/D</td>
<td>16</td>
<td>M/R</td>
<td>21</td>
</tr>
<tr>
<td>Z/Q procida</td>
<td>1</td>
<td>M/Q varallo</td>
<td>1</td>
</tr>
<tr>
<td>Z/I</td>
<td>2</td>
<td>M/M malton</td>
<td>1</td>
</tr>
<tr>
<td>Z/Powell</td>
<td>2</td>
<td>M/Q proclia</td>
<td>1</td>
</tr>
<tr>
<td>Z/SZ</td>
<td>2</td>
<td>M/Plowell</td>
<td>1</td>
</tr>
<tr>
<td>M/Plowell</td>
<td>2</td>
<td>M/Q proclia</td>
<td>1</td>
</tr>
<tr>
<td>M/Q proclia</td>
<td>1</td>
<td>M/Q proclia</td>
<td>1</td>
</tr>
<tr>
<td>M/Q varallo</td>
<td>1</td>
<td>M/Q varallo</td>
<td>1</td>
</tr>
<tr>
<td>M/M malton</td>
<td>1</td>
<td>M/M malton</td>
<td>1</td>
</tr>
<tr>
<td>M/Q proclia</td>
<td>1</td>
<td>M/Q proclia</td>
<td>1</td>
</tr>
</tbody>
</table>

Index cases are in brackets.
Among subjects with severe AATD, COPD (alone or in combination with chronic liver disease) was diagnosed more frequently in the R/D group (100%), than in the two other related groups, PI*ZZ (79%), and PI*SZ (38%, p = 0.004). In seven cases out of 96 PI*ZZ (7%) and in two out of 15 R/D (13%), COPD and liver disease were diagnosed simultaneously. Prevalence of chronic liver disease did not differ among subjects carrying severe AATD; it was absent in M/R subjects, but it was detected in 11% of PI*MZ individuals. A number of other associated conditions were reported: with the exception of one PI*MZ subject diagnosed with Wegener’s granulomatosis, which has been repeatedly associated with AATD, the other conditions are most likely to be chance associations. The percentage of healthy subjects increased significantly among the two groups with intermediate AATD (42% in the PI*MZ group, and 46% in the M/R group).

With reference to the functional phenotype (table 5), the mean basal FEV1 value (% predicted) was significantly lower in the R/D group as a whole than in the related PI*SZ group (p = 0.00095), but not significantly lower than in the PI*ZZ group. The mean FEV1/FVC among the five genotype groups was consistently different (table 5). However, when individuals with normal lung function (healthy, liver disease without COPD, other conditions) were disaggregated from the whole group, the FEV1 and FEV1/FVC in the remaining patients with COPD no longer differed among the R/D, PI*ZZ, and PI*SZ subgroups. The difference between PI*MZ and M/R subjects for FEV1 was significant, but it may have been influenced by the low number of M/R subjects with COPD.

The effect of smoking habit and genotype on FEV1/FVC is summarised in table 6. Logistic regression and multivariate methods were used; values obtained by the latter method are higher than those obtained by nephelometry.

## DISCUSSION

Our finding of a 11% prevalence of subjects with severe AATD carrying genotypes other than PI*ZZ and PI*SZ is, to our knowledge, the highest reported so far. The NHLBI Registry for severe AATD with 1021 subjects includes 1.7% with genotypes other than PI*ZZ or PI*SZ. The Alpha One Foundation Research Network Registry also includes subjects with intermediate AATD: individuals with rare AATD variants accounted for 5.7% of the total. However, in our series including subjects with both severe and intermediate AATD increased the prevalence to 13%. Thus, our large series of rare AATD variants, detected within the relatively small sized Italian registry, prompted us to investigate the characteristics of these subjects more closely.

Molecular characterisation showed that the majority of subjects with rare AATD variants, including both index and non-index cases, carried at least one Mmalton allele (16/37 individuals, 21/74 alleles). This mutation is raised on the M2 base allele and consists of the deletion of an entire TTC codon in exon II, and subsequent deletion of the Phe51 or Phe52 residue of the mature protein. Ten of the 44 subjects carried at least one Mprocida allele, based on M1(Val213), a T→C point mutation at codon 41 exon II, leading to a proline for leucine substitution.

The I allele, found in two individuals, and the Plowell allele (also referred to as Q0 Cardiff), found in four individuals, are both raised on the M1(Val213) base allele. The I allele is characterised by a C→T point mutation at codon 39 exon II, leading to a cysteine for arginine substitution, whereas the Plowell allele is characterised by an A→T transversion at codon 256 exon III, leading to a valine for asparagine substitution.

The Mmolen variant, found in one subject, is raised on the M1(Ala213) base allele and it is characterised by a C→T point mutation at codon 369 exon V, leading to a leucine for proline substitution. The Q0precida allele (also referred to as Nullprecida or Nullsola di precida) is a null variant characterised by a 17 kb deletion encompassing exons II–V. The Q0lnovo, a null variant raised on the M1(Val213) base allele, more recently identified, is characterised by a C insertion in exon V, causing a 3′ frameshift mutation, in turn resulting in a stop codon at residue 376. Finally, two AATD variants were recently found in our series of subjects: the Mvarallo allele was first described in a family from a Northern Italian village.

## Table 3

<table>
<thead>
<tr>
<th>Characteristics of the 132 index subjects with severe AATD (PI<em>ZZ, PI</em>SZ, and PI<em>RD) and the 84 index subjects with intermediate AATD (PI</em>MZ and PI*MZ)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>PI*ZZ</strong> (n = 96)</td>
</tr>
<tr>
<td>Male gender, n (%)</td>
</tr>
<tr>
<td>Mean (SD) age, years</td>
</tr>
<tr>
<td>Smoking habit</td>
</tr>
<tr>
<td>Current, n (%)</td>
</tr>
<tr>
<td>Former, n (%)</td>
</tr>
<tr>
<td>Never, n (%)</td>
</tr>
<tr>
<td>Mean (SD) AAT plasma level, mg/dl</td>
</tr>
</tbody>
</table>

## Table 4

<table>
<thead>
<tr>
<th>Characteristics of the clinical phenotype</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Genotype</strong></td>
</tr>
<tr>
<td><strong>PI*ZZ</strong> (n = 96)</td>
</tr>
<tr>
<td><strong>PI*SZ</strong> (n = 21)</td>
</tr>
<tr>
<td><strong>R/D</strong> (n = 15)</td>
</tr>
<tr>
<td><strong>PI*MZ</strong> (n = 71)</td>
</tr>
<tr>
<td><strong>M/R</strong> (n = 13)</td>
</tr>
</tbody>
</table>

*Category includes: psoriasis, chronic sinusitis, pulmonary fibrosis, coronary artery disease, Wilson’s disease, Wegener’s granulomatosis, idiopathic dilated cardiomyopathy, and pulmonary alveolar proteinosis.

---

www.jmedgenet.com
This allele is characterised by a 30 bp deletion accompanied by a 22 bp fragment insertion at the 41–51 codon region in exon II,25 whereas the Q0cairo is a null variant raised on the by a 22 bp fragment insertion at the 41–51 codon region in

This allele is characterised by a 30 bp deletion accompanied by a 22 bp fragment insertion at the 41–51 codon region in exon II,25 whereas the Q0cairo is a null variant raised on the by a 22 bp fragment insertion at the 41–51 codon region in

This allele is characterised by a 30 bp deletion accompanied by a 22 bp fragment insertion at the 41–51 codon region in exon II,25 whereas the Q0cairo is a null variant raised on the by a 22 bp fragment insertion at the 41–51 codon region in

Thus, within the spectrum of prevalence of lung disease associated with AATD, carrying a rare AATD variant(s) places R/D individuals at the highest risk, in the same position as PI*ZZ individuals, whereas M/R individuals are in an intermediate position, with PI*SZ and PI*MZ individuals at the end with lower risk.28 Since it is widely accepted that phenotypes result from the interaction between genetic determinants (in this case, the plasma levels of AAT) and environmental determinants (in this case, cigarette smoking), then the phenotypic ranking seems to result from the interaction between the AAT plasma levels reported in our series, with the R/D and PI*ZZ groups at the lowest end (29 and 28 mg/dl, respectively), followed by PI*SZ (62 mg/dl) and M/R (61 mg/dl), and with PI*MZ individuals at the highest end (93 mg/dl), and the smoking prevalence, ranking R/D (100%); M/R (85%); PI*ZZ (67%); PI*SZ (52%), and PI*MZ (52%). These findings further support the concept that the genetic risk factor for COPD is significantly related to the AAT level,1 and that cigarette smoking may influence the risk rate27 28 (table 6).

Chronic liver disease, which is the second most common feature associated with AATD, was detected in 13% of R/D subjects, 13% of PI*ZZ subjects, 19% of PI*SZ subjects, and 11% of PI*MZ subjects (table 4). Diagnosis of chronic liver


Prevalence and phenotype of subjects carrying rare variants in the Italian registry for alpha 1-antitrypsin deficiency

I Ferrarotti, J Baccheschi, M Zorzetto, C Tinelli, L Corda, B Balbi, I Campo, E Pozzi, G Faa, P Coni, G Massi, G Stella and M Luisetti

doi: 10.1136/jmg.2004.023903

Updated information and services can be found at:
http://jmg.bmj.com/content/42/3/282

These include:

References
This article cites 25 articles, 5 of which you can access for free at:
http://jmg.bmj.com/content/42/3/282#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

- Epidemiology (630)
- Molecular genetics (1254)
- Liver disease (51)
- Clinical genetics (256)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/