Progress in Alpha-1 Antitrypsin Deficiency: Collaboration as the Foundation of New Knowledge

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Abbreviations: alpha-1 antitrypsin deficiency, AATD; alpha-1 antitrypsin, AAT; chronic obstructive pulmonary disease, COPD; computed tomography, CT; total lung capacity, TLC; functional residual capacity, FRC


Background

By way of background, AATD is a relatively common but under-recognized autosomal co-dominant condition which predisposes to emphysema, panniculitis, and cirrhosis.4,5 Of the >200 alleles described, the Z allele is the most common severe deficiency type and PI*ZZ homozygotes are at risk for liver and lung disease. Estimates suggest that ~100,000 Americans are PI*ZZ but - highlighting the ongoing challenge of detection - fewer than 10,000 have been diagnosed presently.4 Worldwide, there are an estimated 181,894 PI*ZZ individuals.7 Similarly, most are undetected.8 As manifestations of this continuing under-recognition, affected individuals with severe deficiency of AAT often experience long diagnostic delays and see multiple health care providers between their initial symptom and first diagnosis.9,10 These delays are associated with adverse clinical effects.11

First described in 1963 by Laurell and Eriksson in Malmo, Sweden,12 AATD has been extensively studied since, with substantial strides made in understanding the detection, pathobiology, molecular biology, and treatment of AATD. Selected milestones in this understanding of AATD include: the first description of AATD in 1963,12 the first report of liver disease associated with AATD by Sharp et al in 1969,13 formation of the National Heart Lung and Blood Institute Registry for Individuals with AATD in 198814,15 and the Alpha-1 International Registry to characterize the population and the natural history of AATD,16 elucidation of the misfolding and
progress in Alpha-1

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Intrahepatocyte polymerization that characterizes the PI*ZZ state,\textsuperscript{17} initiation of large population-based screening studies in Sweden and the United States,\textsuperscript{18,19} the formation of the Alpha-1 Foundation to broker interactions between the stakeholder communities in 1995,\textsuperscript{6} the description of and initial United States Food and Drug Administration approval of augmentation therapy in 1987, and, more recently, investigation of a number of novel potential therapies (including gene therapy, corrector molecules, and interfering mRNA, among others). On the basis of these many strides, in the current era of precision medicine and focus on distinctive endotypes in chronic obstructive pulmonary disease (COPD), AATD can be regarded as the first well-characterized COPD endotype.

**Team Science**

Importantly, as demonstrated by the papers in this volume and by the very existence of this journal (which began after the COPD Foundation formed as an expansion of its predecessor, the Alpha-1 Foundation), the strides in understanding AATD over its nearly 56-year history represent the outcomes of exemplary collaborations among various communities, all in service of helping patients and pursuing a cure for AATD. Key participants in this collaboration have been the AATD patient community – so-called “alphas” among whom visionary “impatient patient” leaders like the late John Walsh have demonstrated simply inspirational leadership,\textsuperscript{6} patient advocacy and support organizations like the Alpha-1 Foundation which have brokered relationships between the patient community, clinicians, scientists, government, and pharma and funded research,\textsuperscript{6} the clinical and scientific communities of physicians who care for alphas and scientists who investigate AATD; governmental funding agencies like the National Heart, Lung, and Blood Institute which have helped shape and fund key studies,\textsuperscript{14,15} and the pharmaceutical industry which has sponsored research and scholarship,\textsuperscript{3} manufactures and has driven approval of augmentation therapy for AATD, and is advancing new therapeutic ideas. As evidence of the power of this collaborative milieu around AATD, 2 of the 3 papers in this volume were funded by AlphaNet,\textsuperscript{1,2} a patient support organization that emerged from the Alpha-1 Foundation,\textsuperscript{6} and that provides infusion services and support to the alpha-1 community; AlphaNet provides funding to support the mission of the Alpha-1 Foundation, which is “finding a cure for alpha-1 antitrypsin deficiency and improving the lives of people affected by alpha-1 worldwide.”\textsuperscript{20} Through an active grants program and committed volunteers from the scientific, clinical, and patient communities, the Alpha-1 Foundation in turn has issued $71 million in grants to support AATD research since its inception in 1995.\textsuperscript{20}

How then do these 3 papers reflect and contribute to the ever-growing knowledge base about AATD and what are the takeaways from these papers for the clinician? First, the paper by Choate et al\textsuperscript{1} examines self-reported differences between individuals with the PI*ZZ versus PI*SZ genotype. Notwithstanding the risks of bias related to self-reported data from a cohort ascertained by their participating in a disease management program, this paper extends prior reports from smaller cohorts of PI*SZ individuals\textsuperscript{21-23} by newly describing lifestyle and quality of life-related features of having PI*SZ AATD. As PI*SZ individuals differ from PI*ZZ individuals in important ways (e.g., PI*SZ individuals reported more exacerbations and hospitalizations than their PI*ZZ counterparts), this paper emphasizes the critical importance of genotyping at-risk individuals for AATD, as has been advocated in recent guidelines.\textsuperscript{24} Because having AATD has consequences for the proband and also potentially for his/her family, establishing the diagnosis of AATD and the specific genotype is critical. Furthermore, because clinical features among PI*SZ and PI*ZZ individuals differ with different interventional opportunities suggested by Choate et al, testing for AATD must include not only a serum level but also a genotype.

Recognizing how impactful the development and registration of augmentation therapy has been for individuals with AATD, the papers by Brantly et al\textsuperscript{3} and Sieluk et al\textsuperscript{2} remind us that many questions and challenges persist in understanding the effectiveness of augmentation therapy and whether augmentation therapy can be widely available to all eligible patients. Taken together, the weight of evidence from 3 randomized placebo-controlled trials and many observational studies of augmentation therapy clearly supports the efficacy of augmentation therapy to slow the progression of emphysema due to AATD. At the same time, because of the challenges of studying an uncommon disease, none of the 3 randomized trials of augmentation therapy is individually definitive.
For example, as Brantly et al\textsuperscript{3} point out, the primary endpoint in the RAPID trial using computed tomography (CT) density based on both total lung capacity (TLC) and functional residual capacity (FRC) actually failed to achieve statistical significance,\textsuperscript{3,25} though multiple other endpoints in the trial clearly support efficacy, including the significantly lower rate of density loss using TLC-based CT lung density change, and the evident slowing of density decline when individuals initially randomized to placebo crossed over to active drug in the open label extension of the RAPID trial.\textsuperscript{26}

In a similar vein, Sieluk et al\textsuperscript{2} present an analysis of the actual costs associated with having AATD. A key finding is that the estimated total annual cost of treatment for AATD in augmentation therapy users is $127,537, of which 75.3\% is ascribed to the cost of augmentation therapy itself. Fortunately, for those with insurance, the vast majority of the expense is born by insurance, with the average annual out-of-pocket expense for augmentation therapy much less at $2084. Yet, clinical experience clearly shows that the cost of augmentation therapy remains an impediment to full access to the drug for all eligible patients. As novel therapies emerge, whether gene therapy, corrector molecules, or RNA interference to suppress Z protein production, the affordability of therapy must be a paramount consideration.\textsuperscript{27-29}

Taken together, the findings in these 3 papers address a spectrum of issues that surround AATD – how to detect affected individuals, how to treat patients with established emphysema, and the costs of therapy that impact access and affordability of care. These studies, like the journal in which they are published, tangibly exemplify ways in which the pharmaceutical, patient, and clinical/scientific communities have joined to study AATD in service of improving the lives of affected individuals and curing AATD. Having drawn the attention of AATD investigators from multiple disciplines, including pulmonary medicine, adult and pediatric hepatology, pharmacoconomics, structural biology, and epidemiology, among others, AATD is not only the first well-characterized COPD endotype but also an early and powerful example of “team science.”\textsuperscript{30} The model invites emulation.
References


