Pregnancy in Alpha 1 Antitrypsin (AAT) Deficiency and the role of intravenous AAT therapy

Alpha 1 Antitrypsin Deficiency (AATD) is a genetic disorder that results in reduced plasma levels and/or functionality of Alpha 1 Antitrypsin (AAT), a serine protease inhibitor. Deficiency in AAT predisposes patients to greater risk of early-onset chronic obstructive pulmonary disease/emphysema due to excessive proteolysis of lung parenchyma, in addition to liver disease as a result of polymerisation of certain mutant AAT proteins. Weekly intravenous (i.v.) infusion of human AAT (AAT therapy) is currently the only condition-specific treatment for AATD-associated disease, and clinical data suggest that AAT therapy may slow the progression of emphysema in patients with AATD.

There is evidence indicating a role for AAT in conception and pregnancy. Indeed, AAT has been suggested as important to angiogenesis and vascularisation of the endometrium, as well as trophoblast invasion and embryo implantation. Associations between low AAT levels and pregnancy-related complications, such as preeclampsia, spontaneous abortion, and preterm labour, have also been described. Nevertheless, evidence is still limited regarding the clinical consequences of AATD on pregnancy, and less so regarding the use of AAT therapy in pregnant women. Here, we present a case detailing the clinical course of a pregnant woman with AATD, including the initiation of AAT therapy during pregnancy. The patient signed informed consent for use of her de-identified clinical data for research, analysis, and reporting.

The patient, a 31-year-old, underweight (BMI: 24.3), non-smoking female, had been diagnosed with AATD (Pi*SZ genotype) at the age of 21. She was tested for AATD following a smoking female, had been diagnosed with AATD (Pi*SZ genotype) at the age of 21. She was tested for AATD following a

During pregnancy, pulmonology consultation, serum AAT testing and PFTs were performed on a regular basis. The patient experienced one moderate exacerbation at Week 17 of pregnancy, and at Week 18; moreover, a significant drop in her forced expiratory volume in the 1st second (FEV1) (-0.25 L) compared to Week 9 value was recorded at Week 18. Serum AAT level was markedly reduced. So, intravenous (i.v.) augmentation therapy was initiated and was regularly continued throughout pregnancy at a dose regimen of 60 mg/kg/week aimed at maintaining serum AAT level consistently above the protective threshold of 80 mg/dL (11 μM/L). No adverse event associated with augmentation therapy was reported. The patient experienced a severe exacerbation at Week 28 of pregnancy; however, there were relevant improvements to respiratory function parameters (Table 1). Following consultation with a multidisciplinary care team on the risks (in particular severe respiratory tract infection) and benefits of continuing her pregnancy, the patient consented to undergo a caesarean section at 32 weeks, delivering a healthy baby.

At a follow-up visit conducted 4 weeks post-partum, she reported mild exertional dyspnea. A mild exacerbation with uncomplicated clinical course was reported at 3-month follow-up. Over the next 2 years, exacerbations had reduced relative to pre-pregnancy/AAT therapy to three moderate and one severe per year on average. At the patient’s latest consultation (30 months post-partum), PFT showed mild obstructive impairment with recovery of FEV1 to her pre-pregnancy values (Table 1). A lung CT scan was performed, showing a diffuse low attenuation area in the right upper lobe (Fig. 1).

Data regarding AATD and pregnancy are limited, possibly due to the underdiagnosis of women of childbearing age; indeed, patients with AATD are usually identified in their 40s or 50s. Complications associated with preterm labour, preeclampsia and spontaneous abortion and the risk of rapid decline in PFTs highlight a need for close monitoring of the patient throughout pregnancy. Care of moderate exacerbations a year requiring oral antibiotics and/or corticosteroids, and four severe exacerbations a year requiring hospitalization. The patient became pregnant at 29 years of age. Pulmonary Function Tests (PFTs) performed at her last pulmonology consultation prior to pregnancy (6 months pre-pregnancy) showed a severe restrictive ventilatory impairment. (Table 1). Serum AAT level was slightly reduced.

Abbreviations: AAT, Alpha 1 Antitrypsin; AATD, Alpha 1 antitrypsin deficiency; FEV1, forced expiratory volume in the 1st second; FVC, forced vital capacity.

https://doi.org/10.1016/j.pulmoe.2022.01.014
2531-0437/© 2022 Sociedade Portuguesa de Pneumologia. Published by Elsevier España, S.L.U. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

Please cite this article in press as: G. Guarnieri, A. Achille, S. Lococo et al., Pregnancy in Alpha 1 Antitrypsin (AAT) Deficiency and the role of intravenous AAT therapy, Pulmonology (2022), https://doi.org/10.1016/j.pulmoe.2022.01.014
pregnant women with AATD currently follows guidelines for general lung disease, with emphasis on the management of respiratory symptoms and prevention of exacerbations. Whether to initiate AAT therapy relies on expert opinion and clinical experience; however, there is currently little evidence to guide this decision. Although there are no known pregnancy-specific safety concerns with augmentation therapy, data for the use of AAT therapy in pregnancy are limited to a single recent case report by Gauckel et al., which describes a patient who continued with AAT therapy throughout pregnancy and delivered a healthy baby at term. This case adds to the very limited data regarding AAT therapy during pregnancy, supporting the argument that it can be safely initiated in response to severe impairment of respiratory function. The case also adds to the evidence that it is possible for patients with AATD to experience no pregnancy-related complications and deliver a healthy baby; however, close patient monitoring is essential to a positive outcome.

Funding sources

Writing support for this publication was funded by CSL Behring. No study sponsor was involved in designing or conceptualising the study, in collecting, analysing/interpreting the data, in drafting the manuscript or in the decision to submit the manuscript for publication.

Author contributions

Conceptualization: GG; data collection: AA and SL; data interpretation: AV; Writing – Review and editing: all authors.

Conflicts of interest

AV received research grants from CSL Behring. The other Authors have no conflicts of interest to declare.

Acknowledgements

Medical writing assistance was provided by Amy Adlard of Meridian HealthComms Ltd.

References


---

**Table 1** Time course of Alpha 1 Antitrypsin level and lung function. ERS Task Force Global Lung Initiative 2012 reference values [5] were used for lung volumes. (AAT= Alpha 1 Antitrypsin; FEV1= forced expiratory volume in the 1st second; FVC= forced vital capacity; NA= not available; TLC= total lung capacity).

<table>
<thead>
<tr>
<th>Time Point</th>
<th>AAT Level (mg/dL)</th>
<th>FEV1, L (% predicted)</th>
<th>FVC, L (% predicted)</th>
<th>TLC, L (% predicted)</th>
</tr>
</thead>
<tbody>
<tr>
<td>6 months pre-pregnancy*</td>
<td>74</td>
<td>0.79 (31)</td>
<td>1.13 (31)</td>
<td>2.77 (54%)</td>
</tr>
<tr>
<td>9 weeks pregnant</td>
<td>NA</td>
<td>1.90 (59)</td>
<td>2.18 (59)</td>
<td>NA</td>
</tr>
<tr>
<td>18 weeks pregnant**</td>
<td>38</td>
<td>1.65 (51)</td>
<td>2.06 (56)</td>
<td>NA</td>
</tr>
<tr>
<td>23 weeks pregnant</td>
<td>NA</td>
<td>1.46 (45)</td>
<td>1.72 (46)</td>
<td>NA</td>
</tr>
<tr>
<td>26 weeks pregnant</td>
<td>108</td>
<td>1.7 (53)</td>
<td>1.89 (51)</td>
<td>NA</td>
</tr>
<tr>
<td>30 months post-partum</td>
<td>68</td>
<td>1.90 (61)</td>
<td>2.22 (62)</td>
<td>4.87 (96)</td>
</tr>
</tbody>
</table>

* Acute asthma exacerbation.
** Initiation of Alpha 1 Antitrypsin therapy.
G. Guarnieri, A. Achille, S. Lococo, A. Vianello*

Respiratory Pathophysiology Division, Department of Cardio-Thoracic and Vascular Sciences, University of Padova, Via Giustiniani 2, 35128 Padova, Italy

*Corresponding author.

E-mail addresses: gabriella.guarnieri@unipd.it (G. Guarnieri), alessia.achille@aopd.veneto.it (A. Achille), sara.lococo@aopd.veneto.it (S. Lococo), andrea.vianello@aopd.veneto.it (A. Vianello).

Received 18 January 2022; Accepted 27 January 2022

Available online xxx