

Launching an education intervention to increase awareness of alpha-1 antitrypsin deficiency (AATD) among primary care physicians in Southern Alberta

Andrew J. Cave¹, Boglarka Soos^{2,3}, Natalia Stavila¹, Tyler Williamson³, Stephanie Garies^{2,3}, Neil Drummond^{1,2,3}

¹Department of Family Medicine, University of Alberta; ²Department of Family Medicine, University of Calgary; ³Department of Community Health Sciences, University of Calgary

Introduction

- **Alpha-1 Antitrypsin Deficiency (AATD)** is a **genetic disorder** which predisposes individuals to premature onset of chronic obstructive pulmonary disease (COPD) and in some cases, liver damage, panniculitis, and vasculitis.
- Approximately **1 in every 2000-5000** individuals have this disorder¹, including **~1-5% of patients with COPD**.²
- **Less than 10%** of symptomatic individuals have been **properly diagnosed**. In general, there is a **delay of several years** between the onset of symptoms and diagnosis of disease.^{3,4}
- In Canada, primary care physicians provide care for 80-90% of patients with COPD.^{5,6} Guidelines recommend **screening** for AATD among patients with **COPD** who:
 - were diagnosed **before the age of 65** or
 - have **no history of smoking**.²
- In Alberta, screening is available through an inexpensive biochemical assay and is covered by all provincial fee schedules.

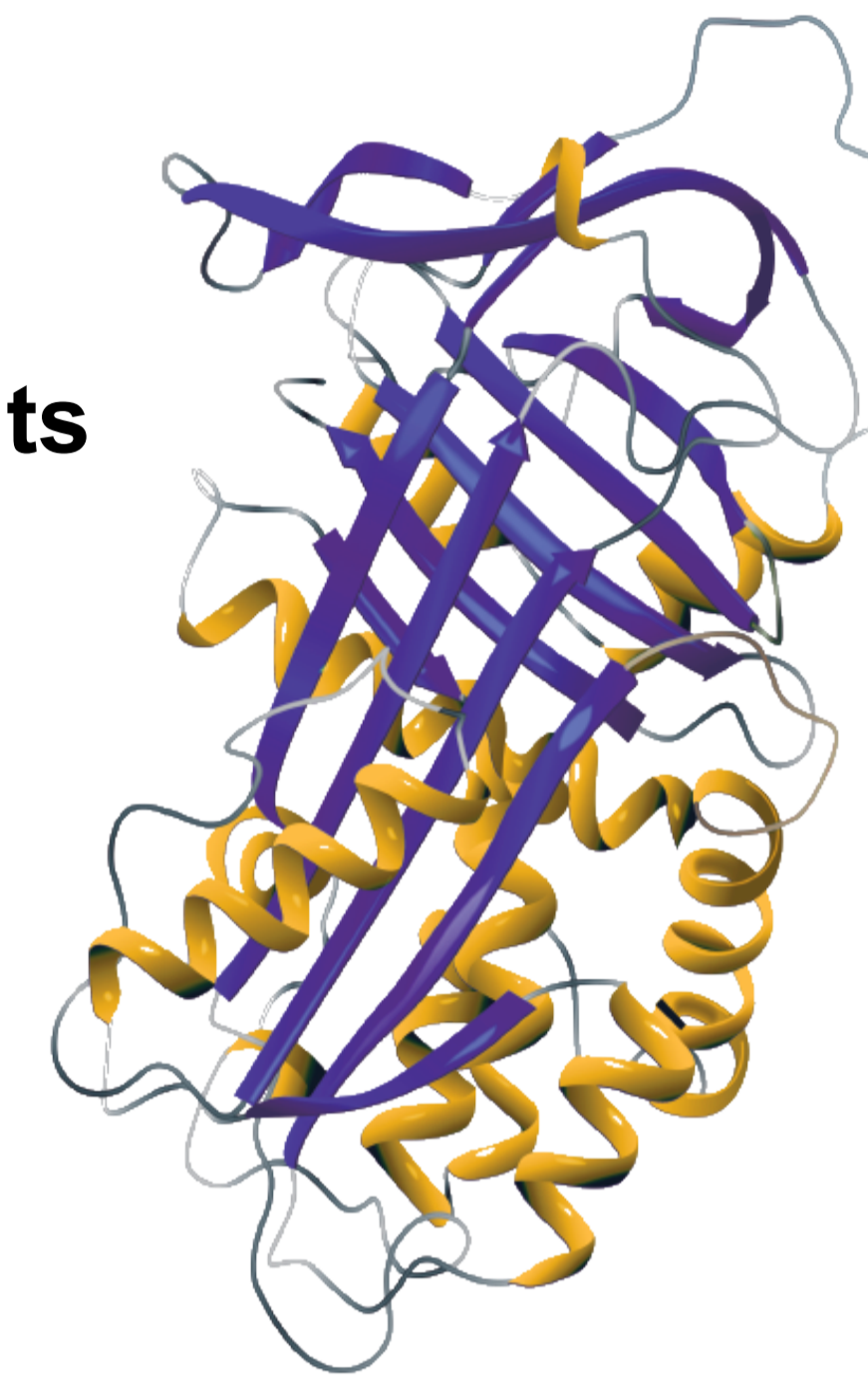


Figure 1: Alpha-1 molecule
Source: <https://www.alphanet.org/what-is-alpha-1/>

Methods

An **education intervention** was launched in southern Alberta to increase awareness and knowledge of AATD.

- Family physicians participating in the Southern Alberta Primary Care Research Network (SAPCRen) were provided information regarding:
 - **symptoms** of AATD
 - **testing** guidelines
 - recommendations for **disease management**
 - a **list of patients** in their panel **who have COPD and qualify for screening** based on guidelines²
- Patients identified as being at high-risk were followed to determine whether they received the recommended AATD screening.

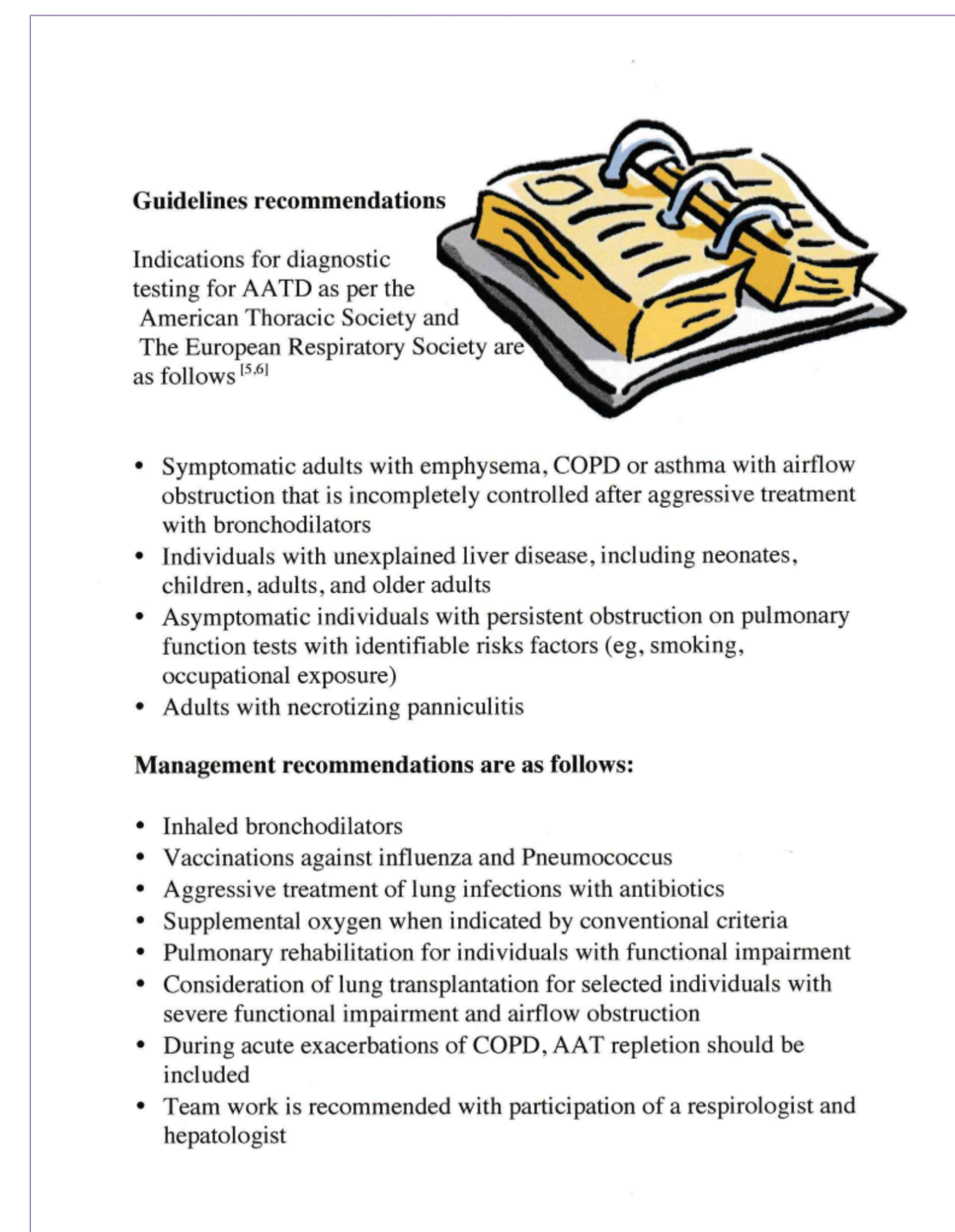


Figure 2: Selected page from educational package

Acknowledgements

The study was completed through the Department of Family Medicine at the University of Alberta and the University of Calgary. The study was supported by an unrestricted grant from Grifols Laboratories, Canada. None of the authors have any conflicts to declare.



Results

- 128 family physicians were emailed the information package and provided the EMR ID of **1071 patients** in total who qualified for AATD screening.
- Following the educational intervention, data from the EMR record of 918 (of 1071) patients were extracted.
 - Loss to follow-up occurred if a physician changed location of practice or a patient changed their provider.
- 90 of the 918 patients (9.8%) had an AATD lab test recorded. 69 patients were screened prior to the intervention (n=41 physicians) including:
 - 7 patients tested in the 365 days prior, and
 - 62 patients tested more than 1 year before the intervention.
- Seven (7) physicians screened 21 patients in the 6 months following the intervention.
- Change point detection was performed using pruned exact linear time (PELT) with various penalties. Changes in the mean and the mean and variance were considered. Statistically significant change points were observed at **3 and 5 weeks** after the distribution of the patient list.

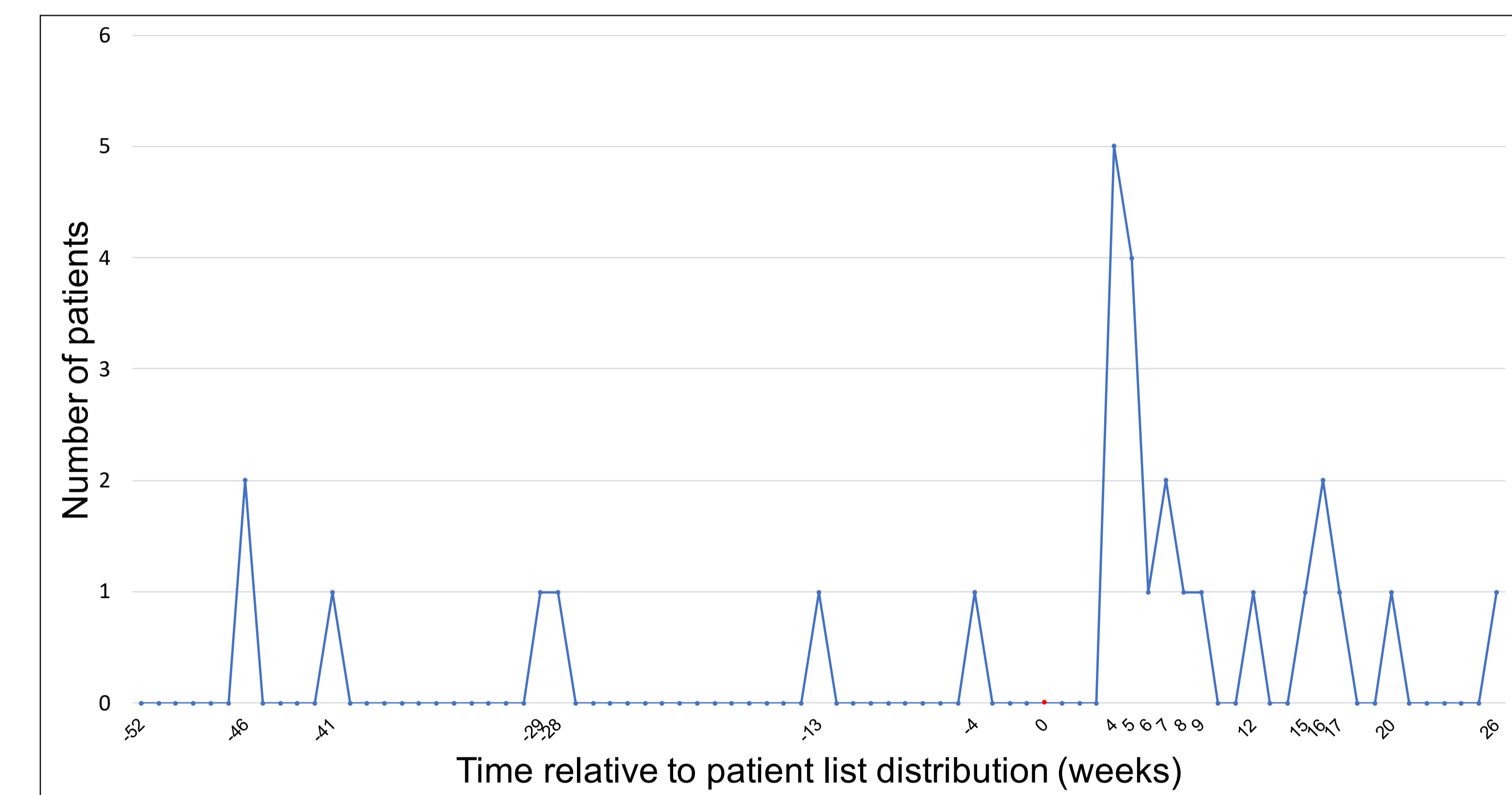


Figure 3: Distribution of AATD tests relative to patient list distribution, by week

Conclusions

This educational intervention aimed to increase knowledge of AATD & encourage physicians to screen appropriately by providing a list of patients at risk. It led to a short term but statistically significant increase in the screening rate among a small number of physicians (n=7). This study suggests a simple e-mail intervention may only impact the practice of a few early adopters and may be short lived.

Bibliography

1. Stroller JK, Lachbawan FL, Aboussouan LS. Alpha-1-Antitrypsin Deficiency. 2006 [Updated 2017]. In: Adam MP, Ardinger HH, Pagon RA, et al., editors. GeneReviews®. Seattle (WA): University of Washington, Seattle; 1993-2019. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK1519/>
2. Marciniuk DD, Hernandez P, Balter M, et al. Alpha-1 antitrypsin deficiency targeted testing and augmentation therapy: a Canadian Thoracic Society clinical practice guideline. *Can Respir J*. 2012;19(2):109-16.
3. Buttar BS & Bernstein M. The importance of early identification of alpha-1 antitrypsin deficiency
4. Lascano JE, Campos MA. The important role of primary care providers in the detection of alpha-1 antitrypsin deficiency. *Postgrad Med*. 2017;129:889-895.
5. Bourbeau J, Bhutal M, Hernandez P et al. CTS position statement: pharmacotherapy in patients with COPD - an update. *Can J Respir, Critical Care & Sleep Med*. 2017;1(4):222-241.
6. Labonte LE, Tan WC, Li PZ, et al. Undiagnosed chronic obstructive pulmonary disease contributes to the burden of health care use. Data from the CANCOLD study. *Am J Respir Critical Care Med*. 2016; 194(3):285-298.