Case report

Acute eosinophilic pneumonia following electronic cigarette use

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ABSTRACT

Electronic cigarette (e-cigarette) use, or vaping, is gaining widespread popularity among adults aged 18–35. E-cigarettes are battery-operated devices that heat liquid nicotine, producing an aerosol or vapor, which the user then inhales [1]. E-cigarettes were first developed in 2003, entered the United States in 2006 [2], and have been promoted as a safe and effective alternative to traditional cigarettes [3]. In spite of their rapid rise in popularity and worldwide sales, the effects of e-cigarettes use on short and long-term health are poorly understood. This case describes a young, healthy female who developed hypoxic respiratory failure from acute eosinophilic pneumonia (AEP) after using e-cigarettes.

1. Introduction

E-cigarettes are battery-operated devices that heat liquid nicotine, producing an aerosol or vapor, which the user then inhales [1]. E-cigarettes were first developed in 2003, entered the United States in 2006 [2], and have been promoted as a safe and effective alternative to traditional cigarettes [3]. In spite of their rapid rise in popularity and worldwide sales, the effects of e-cigarettes use on short and long-term health are poorly understood. This case describes a young, healthy female who developed hypoxic respiratory failure from acute eosinophilic pneumonia (AEP) after using e-cigarettes.

2. Case report

A previously healthy 18-year-old female presented to the Emergency Room after one day of fever, nonproductive cough, difficulty breathing, and pleuritic chest pain. Two months prior to presentation she started vaping using a “Baby Smok Beast Mod” device with 6% nicotine fluid 5 times per day for 30 minutes. She denied traditional cigarette smoking, drug use, exposure to pulmonary irritants, recent respiratory illness or history of deployment to the Middle East. Initial vitals were remarkable for oxygen saturation of 88% on room air, temperature of 102.4 °F, heart rate of 122 beats/min, respiratory rate of 22 breaths/min, and blood pressure of 104/68 mm Hg. On physical exam, she was found to be in mild distress with tachycardia, tachypnea and facial flushing. There was neither accessory muscle use nor chest wall tenderness, and her lungs were clear to auscultation bilaterally. There was no lower extremity edema or calf tenderness. Laboratory tests revealed significant leukocytosis of 19.6 × 10⁹/L with 91.2% granulocytes and 0.5% eosinophils. Initial chest x-ray demonstrated right lower lung airspace consolidation consistent with a pneumonia. Intravenous azithromycin was initiated and she was admitted to the hospital for further monitoring. Overnight, she developed moderate distress with worsening tachycardia, dyspnea, and hypoxemia. Increasing levels of oxygen by nasal cannula were required to maintain her oxygen saturation above 92%, meanwhile her respiratory rate increased to 30 breaths/min. Repeat examination revealed bibasilar inspiratory crackles. D-dimer was elevated to 0.79 mcg/ml. Repeat chest radiograph demonstrated increasing airspace opacities. A computed tomography pulmonary angiogram (CTPA) was completed due to concern for pulmonary embolism (PE). The CTPA excluded PE, but found diffuse ground-glass patchy airspace disease and coalescing nodules (Fig. 1).

Due to worsening respiratory failure overnight, the patient was transferred to the Intensive Care Unit (ICU) for closer monitoring and respiratory support. Bronchoscopy with bronchial alveolar lavage (BAL) was performed, revealing 26% eosinophils in the lavage fluid. Sputum and BAL cultures were negative for viral, fungal, and bacterial pathogens. No other infectious etiologies to include TB, legionella, strongyloides, coccidioiodes, or histoplasma were found. Given her acute onset of symptoms, negative alternative workup, and significant BAL eosinophilia the diagnosis of AEP was made. She was started on methylprednisolone 125mg intravenously every 6 hours. After two days she showed significant improvement and was switched to prednisone 60mg
oral once daily. She fully recovered 6 days after starting steroids, and was discharged home on oral prednisone with subsequent tapering. On discharge, her vital signs and physical exam were within normal limits. She fully recovered 6 days after starting steroids, and was discharged home on oral prednisone with subsequent tapering. On discharge, her vital signs and physical exam were within normal limits.

3. Discussion

Since its original description in 1989 [4], fewer than two hundred AEP cases have been reported; there is a 2:1 male predominance. AEP is a challenging diagnosis to make since patients frequently appear to have a rapidly progressive infectious process with chest radiographs mimicking bacterial pneumonia. A diagnosis of AEP is based upon identification of characteristic symptoms, a detailed patient history, a thorough clinical evaluation, and eosinophilia on BAL [5,6]. Previous studies have reported a close relationship between conventional cigarette smoking and AEP [7–9]. The mechanism by which cigarette smoking induces AEP is suspected to be a strong inflammatory stimulus that recruits macrophages and neutrophils to lung tissue. This induces pro-inflammatory cytokines, such as interleukin (IL)-5, IL-6, IL-7, and tumor necrosis factor, which may be the inciting event causing eosinophil-rich exudate within the alveoli [10,11].

E-cigarettes are gaining widespread popularity over the past few years with 10.8 million adult users in the United States as of 2016 [12]. Recent studies have shown that the mechanism of inflammation and cytokine stimulation in e-cigarette users is similar to cigarette smokers, including elevation of IL-6 and IL-8 [13,14]. This raises concerns that e-cigarettes may induce AEP and other lung diseases, as their use promotes pulmonary inflammation by a mechanism similar to traditional cigarette smoking. Due to the wide variety of vaping devices and fluid brands (and constituent chemical compounds), more research is required to determine the exact cause of e-cigarette induced AEP.

Physicians should consider AEP in previously healthy patients with hypoxic respiratory failure who have a history of recent e-cigarette use. Previous AEP cases have been associated with traditional cigarettes [7], pulmonary irritants [15], or military service in the Middle East [16]. As E-cigarette use becomes more prevalent, it may become a more common trigger of AEP. In this case of clear association, the patient had no other traditional exposures which could cause AEP, leaving e-cigarettes as the most likely causative irritant. One prior case of e-cigarette-associated AEP was reported in a male patient in 2014 [17]; this is the first case of AEP involving a female after use of e-cigarettes. Although most cases of AEP are diagnosed in the male population, further research may suggest whether AEP is more common in males due to confounding factors such as job preference and smoking prevalence, or organic factors.

References

[1] M.A. Orellana-Barrios, D. Payne, Z. Mulkey, K. Nugent, Electronic cigarettes: A narrative review for clinicians, Am. J. Med. 128 (7) (2015) 674–681.
[2] P. Hajek, J.F. Etter, N. Benowitz, T. Eissenberg, H. McRobbie, Electronic cigarettes: review of use, content, safety, effects on smokers and potential for harm and benefit, Addiction 109 (11) (2014) 1801–1810.
[3] O. Rom, A. Pecorelli, G. Valacchi, A.Z. Renzick, Are E-cigarettes a safe and good alternative to cigarette smoking? Ann. N. Y. Acad. Sci. 1340 (2015) 65–74.
[4] J.N. Allen, E.R. Pacht, J.E. Gadek, W.B. Davis, Acute eosinophilic pneumonia as a reversible cause of noninfectious respiratory failure, N. Engl. J. Med. 321 (9) (1989) 569–574.
[5] J.N. Allen, W.B. Davis, Eosinophilic lung diseases, Am. J. Respir. Crit. Care Med. 150 (5 Pt 1) (1994) 1423–1436.
[6] F. Philt, B. Etienne-Mastroianni, A. Parrot, C. Guerin, D. Robert, J.F. Cordier, Idiopathic acute eosinophilic pneumonia: a study of 22 patients, Am. J. Respir. Crit. Care Med. 166 (9) (2002) 1235–1239.
[7] H. Shintani, M. Fujimura, M. Yasui, et al., Acute eosinophilic pneumonia caused by cigarette smoking, Intern. Med. 39 (1) (2000) 66–68.
[8] H. Uchiyama, T. Suda, Y. Nakamura, et al., Alterations in smoking habits are associated with acute eosinophilic pneumonia, Chest 133 (5) (2008) 1174–1180.
[9] E. Grossi, G. Poletti, V. Poletti, Acute eosinophilic pneumonia with respiratory failure: a case likely triggered by cigarette smoking, Monaldi Arch. Chest Dis. 61 (1) (2004) 58–61.
[10] J.N. Allen, Z. Liao, M.D. Wevers, E.A. Allenberger, S.A. Moore, E.D. Allen, Detection of IL-5 and IL-1 receptor antagonist in bronchoalveolar lavage fluid in acute eosinophilic pneumonia, J. Allergy Clin. Immunol. 97 (6) (1996) 1366–1374.
[11] W.G. Kuschnier, A. D'Alessandro, H. Wong, P.D. Blan, Dose-dependent cigarette smoking-related inflammatory responses in healthy adults, Eur. Respir. J. 9 (10) (1996) 1969–1994.
[12] M. Mirbolouk, P. Charkhchi, S. Kianoush, et al., Prevalence and distribution of E-cigarette use among U.S. Adults: behavioral risk factor surveillance system, 2016, Ann. Intern. Med. 169 (7) (2018) 429–438.
[13] C.A. Lerner, I.K. Sundar, H. Yan, et al., Vapors produced by electronic cigarettes and e-juices with flavorings induce toxicity, oxidative stress, and inflammatory response in lung epithelial cells and in mouse lung, PLoS One 10 (2) (2015) e0116732.
[14] Q. Wu, D. Jiang, M. Minor, H.W. Cha, Electronic cigarette liquid increases inflammation and virus infection in primary human airway epithelial cells, PLoS One
[15] W.N. Rom, M. Weiden, R. Garcia, et al., Acute eosinophilic pneumonia in a New York City firefighter exposed to World Trade Center dust, Am. J. Respir. Crit. Care Med. 166 (2002) 797.

[16] A.F. Shorr, S.L. Scoville, S.B. Cersovsky, et al., Acute eosinophilic pneumonia among US Military personnel deployed in or near Iraq, J. Am. Med. Assoc. 292 (2004) 2997.

[17] D. Thota, E. Latham, Case report of electronic cigarettes possibly associated with eosinophilic pneumonitis in a previously healthy active-duty sailor, J. Emerg. Med. 47 (1) (2014) 15–17.