Case Report and Review of Esophageal Lichen Planus Treated With Fluticasone

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Case report and review of esophageal lichen planus treated with fluticasone

Marie Lourdes Ynson, Faripour Forouhar, Haleh Vaziri

INTRODUCTION
Lichen planus is a well-recognized chronic idiopathic disorder involving the skin, nails and mucosal surfaces including the mouth, pharynx and perineum. It affects less than 1% of the general population. Mucosal surface involvement is found in about 30%-70% of patients diagnosed with lichen planus. Esophageal involvement, on the other hand, is considered to be rare with its true prevalence unknown. The study by Dickens et al demonstrated esophageal lichen planus in 26% of patients with mucocutaneous lesions while a larger study by Eisen found esophageal lesions in only 1% of patients. Here we describe a case of esophageal lichen planus who was treated with swallowed fluticasone with good response and present a review of this topic.

CASE REPORT
A 63-year-old female with history of gastric bypass surgery, coronary artery disease, and hyperlipidemia reported having intermittent solid food dysphagia for the past 3 years. She also had symptoms of gastroesophageal reflux disease that was initially controlled with omeprazole 20 mg daily. Later on, she developed multiple episodes of breakthrough and nocturnal reflux symptoms accompanied by occasional difficulty in swallowing large pills that happened 2-3 times a week. She denied food regurgitation, odynophagia, nausea, vomiting or weight loss. For further evaluation of her dysphagia, a barium swallow
was performed. During the procedure, the barium pill briefly got trapped in the upper esophagus at the level of C6-C7. This location correlated with the reported site of dysphagia (Figure 1). The rest of the exam was uneventful and no mass or stricture was noted except for a small sliding hiatal hernia. She subsequently underwent an upper endoscopy which showed areas of friable mucosa and dried blood in the upper esophagus with a mild to moderate stricture at the upper and mid-esophageal junction (Figure 2). Esophageal biopsies were obtained which showed hyperkeratosis, parakeratosis and spongiosis of the lower third of the epithelium with focal dyskeratosis. Chronic inflammatory cell infiltration of the epithelium was also present and was mainly composed of lymphocytes (Figure 3). Submucosa was not present for evaluation of lichenoid inflammation. Nonetheless, in the presence of the clinical history, the findings were interpreted to be consistent with lichen planus. On further questioning, patient reported having an oral lesion diagnosed to be lichen planus in 2009. Due to her history of previous lichen planus and endoscopy findings, patient was diagnosed with esophageal lichen planus and was started on swallowed fluticasone propionate 220 mcg twice daily for 6 wk. At her 4th wk follow-up she noted resolution of her symptoms. On follow-up endoscopy at 15 wk, only small light pink plaques were noted in the mid-esophagus while the rest of the esophagus showed normal looking mucosa (Figure 4). Pathology

Figure 1  Barium swallow showing barium pill (arrow) trapped in C6-C7.

Figure 2  Endoscopy image showing friable mucosa and dried blood in upper esophagus with a mild to moderate stricture at upper and mid-esophageal junction.

Figure 3  Chronic inflammatory cell infiltration of the epithelium was also present and was mainly composed of lymphocytes. A: Low power view of upper esophagus shows extensive severe keratinization, organizational disarray of cell arrangement and spongiosis of lower layer associated with inflammatory infiltration. (E, × 40); B: High power of parakeratotic cells and accumulation of keratohyaline granules in the cytoplasm of the mature keratinocytes. In this zone keratinization is mild. (HE, × 400); C: High power view of spongiosis of lower layers of squamous epithelium associated with sprinkling of lymphocytes typical of longstanding chronic inflammation. (HE, × 400).

Figure 4  Endoscopy image showing resolution of lesions in the upper esophagus post treatment.
showed normal to atrophic squamous mucosa with mild non-specific inflammation including rare eosinophils and no evidence of interface band-like lymphocytic infiltrate which was compatible with lichen planus post-therapy.

<table>
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<td>Age and gender distribution</td>
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<td>Histologic findings</td>
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<td>Risk of malignancy transformation</td>
<td>No increased risk of malignant transformation</td>
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*One report by Chryssostalis et al[8] showed esophageal lichen planus in a 22-yr-old male patient.

**DISCUSSION**

Lichen planus is a common disorder of squamous epithelium[8]. The exact cause of this disease is unknown; however infectious (viral), neurologic, genetic and immunologic causes have been proposed[6]. Lichen planus appears to be mediated by cytotoxic CD8+ T cells that attack an antigen in the basal epithelium in a manner resembling graft-versus-host disease[9]. There are no detailed studies published to specifically address the pathogenesis of esophageal lichen planus. Chandan et al[7] suggests that the pathogenesis may be similar to oral lichen planus. It has been suggested that oral lichen planus is due to an immune response to an exogenous or endogenous antigen found in the basal keratinocytes. This activates the Langerhans cells which then present the antigen to the CD4+ T lymphocytes and basal layer degeneration including characteristic Civatte bodies (i.e., apoptotic basal keratinocytes)[21].

Associated with hepatitis C

In our patient, esophageal lichen planus was favored over reflux esophagitis due to the extensive degree of hyperkeratosis and focal parakeratosis with accumulation of keratohyaline granules which are unlikely in reflux related injury. The presence of spongiosis in the lower third of the epithelium was also more consistent with esophageal lichen planus. The spongiosis in reflux diseases tend to occur in the upper third of the epithelium where acid is in contact with the mucosa. The other factor that can help to differentiate these 2 entities is the site of involvement in the esophagus. While reflux disease is usually more severe in the distal esophagus, lichen planus affects the middle and upper esophagus in most cases.
The most characteristic histologic finding in esophageal lichen planus is a bandlike or lichenoid lymphocytic infiltrate involving the superficial lamina propria and basal epithelium[2]. A predominance of mature T cells is present within this infiltrate. These are associated with basal keratinocyte degeneration which often include Civatte bodies (necrotic keratinocytes with anucleate remnants)[7]. Lymphocytic infiltration of the mucosa is not pathognomonic of this disease and medications such as gold, thiazide, and anti-malarials can induce lichen planus-like lesions and need to be excluded clinically[8,9].

No clear guidelines are present for the treatment of esophageal lichen planus and there is no specific way to treat this entity. Historically, systemic corticosteroids have been used as first-line treatment with a response rate of up to 74% based on multiple reports. However, relapse rate can be as high as 85% with steroid withdrawal[10]. Treatment response was also reported with adrenocorticotropic hormone injection, etretinate, topical tacrolimus, intralesional corticosteroids and cyclosporine[11,12]. Esophageal dilation is commonly used to treat strictures, although intralesional steroid injections and/or oral tacrolimus may decrease the frequency of strictures. It is important to note that koebner phenomenon which is defined as development of new lesions along the lines of trauma, can occur with dilation.

Although effective, systemic corticosteroid treatment can be associated with serious side effects and therefore there is a need for a safer alternative. In our review of literature, we have come upon case reports and a recent case series where treatment with swallowed fluticasone propionate has resulted in symptomatic improvement as well as endoscopic improvement in 4 out 6 treated patients[13,14]. Because of these promising results, we decided to try this novel approach for our patient which proved to be successful.

Oral lichen planus has a 1%-3% risk of malignant transformation to squamous cell carcinoma[15]. It is unknown however if the same is true for esophageal lichen planus. To date, 3 case reports have described squamous cell carcinoma and 1 described a verrucous carcinoma arising from these lesions[16,17,18]. In these reports, squamous cell carcinoma developed more than 20 years after the diagnosis of esophageal lichen planus[18]. Due to this risk, some advocated surveillance endoscopies in patients with esophageal lichen planus to detect early malignancy. In the study by Quispel et al.[18], high magnification indigo carmine chromoendoscopy was used to establish the prevalence of endoscopic and histopathologic esophageal abnormalities consistent with lichen planus and dysplasia in a cohort of patients with lichen planus. It was found that up to 50% of patients with orocutaneous lichen planus had esophageal involvement but no dysplasia was found. The authors of this study proposed to set a low threshold for performing endoscopy in patients with lichen planus and symptoms suggestive of esophageal involvement but recommended against routine screening[18]. Based on these data, we recommend that the frequency of screening by endoscopy be individualized, but the possibility of malignancy be kept in mind while evaluating these patients with a low threshold for further evaluation in patients with symptoms suggestive for esophageal cancer.

In conclusion, esophageal lichen planus should be suspected in middle-aged female patients who present with symptoms of dysphagia and odynophagia and found to have mucosal abnormalities involving the upper third of the esophagus. Once the diagnosis of esophageal lichen planus is made, these patients should be considered for treatment with swallowed fluticasone before other systemic and more toxic therapies. It is important to keep in mind that secondary to the risk for malignant transformation, these patients should be followed closely.

REFERENCES


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