Oral Lichen Planus: Diagnosis and Management

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LITERATURE REVIEW

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ABSTRACT

Oral lichen planus (OLP) is a rather common oral disease, mainly affecting adults, occurring more often in women than in men. The etiopathogenesis is still unclear. The diagnosis may be cumbersome, even in the presence of a biopsy. In addition, there are several lesions that may resemble lichen planus (lichenoid lesions) both clinically and histopathologically. Treatment of OLP can only be symptomatic and usually consists of topical application of corticosteroids. The disease is characterized by remissions and exacerbations and may persist in some patients lifelong. There is an ongoing debate in the literature as whether OLP is a potentially malignant disease. Because of this uncertainty, annual follow-up is advised.

Key words: oral lichen planus; oral mucosa

INTRODUCTION

Lichen planus (LP) is a chronic mucocutaneous disorder which may involve the oral mucosa only (“isolated OLP”) or in combination with involvement of other mucosal surfaces, e.g. vagina (“vulvovaginal-gingival syndrome”), oesophagus, pharynx, and nails. It has been suggested that extra-oral involvement in oral lichen planus is more common than suspected.1,2 A few cases have been reported of transformation of OLP into mucous membrane pemphigoid, being referred to as lichen planus pemphigoides.3 Attention will be paid at the diagnostic aspects of OLP and lichenoid lesions, as well at the various treatment modalities. An important question whether OLP is a benign lesion or a potentially malignant one.

Etiology

Lichen planus is most likely caused by T-cell autoimmunity. Apparently, Langerhans cells play an important role in this process.8 Controversial views have been reported about the role of human papilloma virus (HPV), particularly HPV 16.9,10 There is no evidence that psychological factors are a direct etiological factor. In some populations there is a significant, yet unexplained, association between lichen planus and hepatitis C virus.11 Yet, in other populations an association with thyroid diseases, especially hypothyroidism, has been reported.12

Lichen planus or lichen planus-like lesions, usually referred to as lichenoid lesions or lichenoid reactions, may occasionally be caused by certain drugs.13 Also prolonged, direct anatomic contact with large amalgam restorations (“contact lesion”) may produce a lichenoid appearance of the oral mucosa. Furthermore, a chronic graft versus host reaction after an allogenic stem cell transplantation may strongly resemble OLP. Interestingly, the lower lip is a site of predilection in this particular patient population.
Clinical aspects

Signs and symptoms

Oral lichen planus has various clinical manifestations, such as reticular (characterized by white striae, also referred to as “Wickhams striae”), annular, papular, erythematous, plaque-type, ulcerative and bullous type. The reticular, the erythematous and the plaque types are the most common ones. The various types may occur simultaneously in an individual patient and may also change in time. (Figure 1 and 2). It has been suggested that the plaque type is more common among smokers. There is nearly always a bilateral, more or less symmetrical distribution. The buccal mucosa, the gingiva and the dorsal surface of the tongue are the sites of predilection. Occurrence in the floor of the mouth and the palate is rare. Superficial mucoceles may develop within or adjacent to mucosal areas that are involved by lichen planus, while in dark-skinned people post-inflammatory melanosis has been reported to occur in the affected areas.

Particularly the erythematous type may cause symptoms, such as pain and, in case of gingival involvement, severe bleeding during tooth brushing. Patients affected by the erythematous or ulcerative type of OLP usually do not tolerate spicy food. When the gums are involved patients may complain about the esthetic aspect of their gums. In involvement of the gingiva the presence of the “vulvovaginal gingiva syndrome” should be considered.

The course of oral lichen planus is characterized by remissions and exacerbations with intervals of several weeks or months of both the clinical signs and the symptoms. There are various scoring systems, mainly designed for scientific studies, to quantify the clinical signs and symptoms of OLP.14

Clinical differential diagnosis, including lichenoid lesions

Lichenoid lesions may be caused by direct anatomic contact with a large (amalgam) restoration. Replacement of the restoration should result in healing of the lesion within several weeks, but this should be checked by the clinician. One may consider to take a biopsy before advising to have the amalgam restoration removed in order to exclude the possibility of a dysplastic or even a malignant lesion. The predictive value of epicutaneous patch testing for amalgam and mercury allergy is rather questionable.15,16 In contrast to drug-induced cutaneous lichenoid lesions drug-induced oral lichenoid lesions are extremely rare. Nevertheless, one may consider to adjust the medication (Box 1), if applicable. Of course, such adjustment requires communication with the prescribing specialist.

As has been mentioned already, lichenoid lesions may occur in chronic graft-versus-host disease, being clinically sometimes indistinguishable from “idiopathic” OLP. At times, it may be difficult, if not impossible, to clinically distinguish oral lichen planus not only from lichenoid lesions but also from leukoplakia, lupus erythematoses, linear IgA disease, mucous membrane pemphigoid and some other white or red lesions of the oral mucosa.17 In some of these cases the medical history or other clinical features are helpful in establishing the correct diagnosis. Unfortunately, a biopsy is not always helpful in such cases. Some clinicians will always do a biopsy in case of oral lichen planus, while others take a biopsy only in case of doubt about the diagnosis.

Histopathology

In case of a distinct clinical diagnosis of lichen planus the histopathologic features are usually quite characteristic (Figure 3.). In scientific studies usually reference is made
Is oral lichen planus potentially malignant and can such event be prevented?

Oral lichen planus is by some regarded as a premalignant condition, the annual malignant transformation rate being less than 1%. If this event occurs, it usually concerns the tongue. The suggestion has been made that a certain cell surface marker on hematopoietic stem cells and early progenitor cells in the bone marrow, referred to as CD133, correlates with the progression to oral squamous cell carcinoma. Others have not found any evidence for a potentially malignant behavior. Anyhow, such malignant transformation can not be prevented, although it has been speculated that the use of topical steroids may prevent such event.

Treatment

Oral lichen planus may last for many years, if not lifelong. There is no cure for oral lichen planus. Treatment can only be symptomatic and most commonly consists of topical or, occasionally systemic, corticosteroids. The use of corticosteroids may enhance the risk of the development of candidiasis, sometimes necessitating antifungal therapy. Also the use of many other drugs, e.g. tacrolimus, thalidomide, topical aloe vera, oral curcuminoids and lysopine has been reported. The efficacy of all these drugs is rather questionable. There are opposing views about the effect of removing all amalgam restorations, not just the ones directly related to a lesion. It has been shown that plaque control improves the symptoms of OLP with gingival involvement. A variety of other treatment modalities has been reported, such as surgical excision, laser evaporation, laser excision and photodynamic treatment. Such treatment modalities may be useful in persistent localized areas of lichenoid involvement.

Follow-up

Since the issue of premalignancy is still under debate, it seems safe practice to check the patient annually for any possible unfavorable change. However, it is questionable whether such follow-up is really effective in either preventing malignant transformation or in resulting in better survival in case of cancer development.

REFERENCES


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