Cancer of the oral cavity accounts for approximately 3% of all malignancies diagnosed annually in 270,000 patients worldwide. Oral cancer is the 12th most common cancer in women and the 6th in men. Many oral squamous cell carcinomas develop from potentially malignant disorders (PMDs). Lack of awareness about the signs and symptoms of oral PMDs in the general population and even healthcare providers is believed to be responsible for the diagnostic delay of these entities. The aim of this article is to update and improve the knowledge of healthcare providers about oral PMDs.

Key words: Precancerous conditions, oral cavity neoplasms, malignancy, leukoplakia, oral cancer.
signs and symptoms of oral PMDs among general population and even physicians are believed to be responsible for the diagnostic delay of these entities.\textsuperscript{1,3} The aim of this article is to update and improve the knowledge of healthcare providers about oral PMDs.

What are PMDs?

The clinical concept of malignant transformation in oral mucosa has been proposed for more than 100 years. Sir James Paget\textsuperscript{15} first described malignant transformation of an oral lesion into tongue carcinoma in 1870. Schwimmer\textsuperscript{16} also reported the same finding in 1877. Several years later, the term "potentially malignant disorders" was defined by World Health Organization (WHO) as the risk of malignancy being present in a lesion or condition either during the time of initial diagnosis or at a future date. WHO also classified PMDs into two subgroups as follows: a) precancerous lesion, a benign lesion with morphologically altered tissue, which has a greater than normal risk of transforming into malignancy; b) precancerous condition, a disease or patients’ habit that does not necessarily alter the clinical appearance of local tissues but is associated with a greater than normal risk of precancerous lesion or cancer development in that tissue (Table 1).\textsuperscript{3,4} Although little information is available regarding the real prevalence of PMDs in the general population, a commonly accepted prevalence of 1–5% has been reported.\textsuperscript{5} Average age of patients with PMDs is 50–69 years, which is 5 years before occurrence of oral cancer. Unfortunately, in recent years 5% of PMDs has been observed in persons under 30.\textsuperscript{1} Premalignant disorders are usually found on the buccal mucosa, followed by gingivae, tongue and floor of the mouth.\textsuperscript{1,16}

Characteristics of precancerous disorders are summarized in Table 2.

\textit{Leukoplakia (Leukokeratosis)}

Schwimmer\textsuperscript{16} first used this term in 1877 to describe a white plaque on the tongue. Since 1980, World Health Organization has changed the definition of leukoplakia as follows:\textsuperscript{3}

1) A white patch or plaque that cannot be characterized clinically or histologically as any other disease.
2) A predominantly white lesion of the oral mucosa that cannot be characterized as any other definable lesion.
3) A white plaque of questionable risk having excluded (other) known disease or disorders that carry no increased risk for cancer.

This lesion is seen most often in middle-aged and older men. Less than 1% of males under 30 years of age have leukoplakia, but the prevalence increases to an alarming 8% in men and 2% in women over the age of 70.\textsuperscript{10,17} Although leukoplakia is more common in males than in females, the latter has a higher risk of developing oral cancer.\textsuperscript{4} Approximately 70% of oral leukoplakias are seen on the buccal mucosa, lip vermilion and gingivae. However, lesions on the floor of the mouth (42.9%), tongue (24.2%) and lip vermilion (24%) account for more than 90% of those with dysplastic or malignant changes. The rate of dysplastic or malignant alterations in oral leukoplakia has been reported to be between 15.6% and 39.2%. Leukoplakia usually occurs about 5 years earlier than OSCC.\textsuperscript{4,10} Waldron and Shafer\textsuperscript{18} demonstrated that 19.9% of leukoplakia had some degrees of dysplasia, 3.1% were frank carcinoma, 4.6% showed severe dysplasia or carcinoma in situ, and 12.2% showed mild to moderate dysplasia.

There are 4 subdivisions of leukoplakia: early or thin, homogenous or thick, granular or verruciform and speckled or erythroleukoplakia. Each subdivision has a different malignant transformation potential. For example, thin leukoplakia often becomes malignant without clinical changes. Thick leukoplakia undergoes malignant transformation in 1–7% of cases. The frequency of malignant changes in verruciform and speckled leukoplakia ranges from 4% to 15% and 18% to 47%, respectively.\textsuperscript{4}

In a recent study, Liu et al\textsuperscript{19} found that 20.8% of patients with verrucous leukoplakia developed cancer. They pointed out that age and lesion site were risk factors for malignancy, whereas sex, smoking, and alcohol intake were not. Leukoplakia exhibiting moderate to severe dysplasia warrants complete removal by electrocautery, cryosurgery or laser ablation. Leukoplakia not exhibiting dysplasia usually is

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Table 1. Potentially Malignant Disorders (PMDs)

| Premalignant lesions     | Premalignant conditions                       |
|--------------------------|-----------------------------------------------|
| Leukoplakia              | Lichen planus                                 |
| Erythroplakia            | Discoid lupus erythematosus                   |
| Proliferative verrucous leukoplakia (PVL) | Epidermolysis bullosa                        |
| Viadent leukoplakia      | Verruciform xanthoma                          |
| Candida leukoplakia      | Grafi-versus-host-disease                     |
| Reverse smokeings’ palate | Cheilitis glandularis                        |
| Verrucous hyperplasia    | Xerodermia pigmentosum                       |
| Oral verrucous carcinoma | Syphils (third stage)                        |
| Dyskeratosis congenita   | Plummer-Vinson syndrome                      |
| Actinic cheilosis        | Malnutrition                                |
| Keratoacanthoma          | Vitamin A, B, C deficiency                   |
| Oral submucous fibrosis  | Immunosuppressive diseases [AIDS]             |
not excised, but clinical assessment every 6 months is recommended.

**Erythroplakia (Erythroplasia, Erythroplasia of Queyrat)**

Fournier and Darier first described erythroplasia as a malignant dyskeratosis with unknown etiology in 1893 and designated it as épithéliome papillaire. WHO defined oral erythroplakia or erythroplasia during the years as follows:

1. Bright red, velvety plaques which cannot be characterized clinically or pathologically as being due to any other condition.
2. Red areas that cannot be diagnosed as any other definable lesion.
3. A predominantly red lesion of the oral mucosa that cannot be characterized as any other definable lesion.

Clinically, erythroplakia is classified into three types: homogeneous, granular, and speckled. Erythroplakia most frequently occurs in males aged 50-70. The point prevalence rate of erythroplakia in the oral cavity has been estimated to be 1 per 2500 adults. The lowest prevalence of erythroplakia was

**Table 2. Characteristics of potentially malignant disorders**

| Disorders                          | Clinical features                          | Locations                                | Risk of malignancy |
|------------------------------------|-------------------------------------------|------------------------------------------|--------------------|
| Leukoplakia                        | White plaque                              | Cheeks, lips, gingivae                   | 15.6–39.2%         |
| Early (thin)                       |                                           |                                          | NA                 |
| Homogenous                         |                                           |                                          | 1–7%               |
| Verruciform                        |                                           |                                          | 4–15%              |
| Speckled                           |                                           |                                          | 18–47%             |
| Erythroplakia (Erythroplasia, Erythroplasia of Queyrat) | A predominantly red lesion | Mouth floor, tongue, retromolar pad, soft palate | 51% |
| Proliferative verrucous leukoplakia (PVL) | Multifocal white patch or plaque + rough surface projections | Gingivae | 63.3–100% |
| Viadent leukoplakia                  | White patch or plaque                     | Gingivae, buccal and labial vestibule   | NA                |
| Candida leukoplakia                 | Firm, white leathery plaques              | Cheeks, lips, palate                     | 4–5 times more common than leukoplakia |
| Smokeless tobacco keratosis         | White plaque                              | Buccal or labial vestibule              | NA                |
| Palatal keratosis associated with reverse smoking | White patches and plaques                        | Palate, tongue | 83.3% dysplasia  |
| Verrucous hyperplasia               | Extensive thick white plaque               | Buccal tongue                            | 68% dysplasia      |
| Oral verrucous carcinoma            | Extensive thick white plaque               | Buccal mucosa                            | 20%                |
| Dyskeratosis congenita              | Oral leukoplakia                           | Buccal mucosa, tongue, oropharynx       | NA                |
| Actinic cheilosis                   | Diffuse, poorly defined atrophic, erosive, ulcerative or keratotic plaques | Lower lip | 6–10% |
| Keratoacanthoma                     | Firm, sessile non tender nodule + a central plug of keratin | Lips, tongue, sublingual region | 24% |
| Oral submucous fibrosis             | Mucosal rigidity                          | Buccal mucosa, retromolar area, tongue, soft palate | 7–26% |
| Lichen planus                      | Reticular, erosive, atrophic, bullous, ulcerative, popular, plaque like | Posterior buccal mucosa, tongue, gingivae, palate, vermilion border | 0.4–3.7% |
| Discoid lupus erythematosus         | White plaques with elevated borders, radiating white striae and telangiectasia | Cheeks, lips, palate | NA |
| Epidermolysis bullosa               | Bullae and vesicle formation following mild trauma | Cheeks, tongue, palate | 25% |
| Verruciform xanthoma                | A well demarcated mass with a yellow-white or red color and a papillary or verruciform surface | Gingivae, tongue, buccal mucosa, vestibular mucosa, floor of the mouth | NA |
| Graft-versus host disease           | Atrophy, erythema, erosions, ulcers, lichenoid lesions | Cheeks, tongue, lips, buccal & labial vestibule | NA |

NA: not assigned
reported to be 0.01% by Lumerman et al\textsuperscript{21} and the highest was 0.2% according to Hashibe et al.\textsuperscript{22} Common sites of involvement are floor of the mouth, tongue, retromolar pad, and soft palate. This lesion is usually asymptomatic, but some patients may complain of a burning sensation.\textsuperscript{4,10} According to Shafer and Waldron,\textsuperscript{23} 51% of erythroplakias transformed into SCC, 40% were carcinoma in situ and 9% showed mild to moderate dysplasia. Because of 90% malignant transformation rate, early detection and immediate surgical excision are recommended. Early cancers of the oral cavity and lip (stage I, stage II) have a better prognosis; therefore, surgery or radiation are the treatments of choice. Chemotherapy is added to surgery and/or radiation in stage III and higher.\textsuperscript{4}

**Proliferative Verrucous Leukoplakia**

Proliferative verrucous leukoplakia (PVL) is a distinct and aggressive form of oral leukoplakia, which was first described by Hansen et al\textsuperscript{24} in 1985. Most patients with PVL are women (female-to-male ratio of 4:1) without any history of tobacco use. Clinically, PVL presents on the oral mucosa as a multifocal white patch or plaque with rough surface projections. These lesions tend to spread slowly and involve other oral mucosal regions. PVL occurs more frequently on the gingiva; however, other sites may be affected as well.\textsuperscript{4} Bagan et al\textsuperscript{25} and Batsakis et al\textsuperscript{26} found SCC development in 63.3% and 100% of PVL, respectively. Transformation into SCC usually occurs within 8 years of initial PVL diagnosis.\textsuperscript{4} PVL is also characterized by resistance to generally approved treatments such as surgery, CO\textsubscript{2} laser evaporation, laser surgery, chemotherapy and radiotherapy.\textsuperscript{27}

**Viadent Leukoplakia (Sanguinaria-associated Keratosis)**

Viadent leukoplakia is a white patch or plaque, which is associated to sanguinaria mouth rinse. Sanguinaria extract is a mixture of benzophenanthridine alkaloids derived from bloodroot plant.\textsuperscript{28} According to Damm et al\textsuperscript{29} an increased prevalence of leukoplakia has been observed in the maxillary alveolar mucosa in patients using sanguinaria mouthwash. It seems that sanguinaria extract has a carcinogenic effect. Therefore, viadent leukoplakia should be considered a premalignant lesion and mouth rinses containing sanguinaria should be avoided until the risk for transformation into malignancy is determined.\textsuperscript{30}

**Candida Leukoplakia (Candida Hyperplasia, Chronic Hyperplastic Candidiasis)**

Candida leukoplakia (CL) is a chronic form of candidiasis which is characterized by firm, white, leathery plaques on the cheeks, lips, palate and tongue. Generally, epithelial dysplasia occurs four to five times more frequently in CL than in leukoplakia.\textsuperscript{28}

**Smokeless Tobacco Keratosis (Snuff Dipper’s Lesion, Spit Tobacco Keratosis, Tobacco Pouch Keratosis)**

Smokeless tobacco keratosis is characterized by a white plaque in the buccal or labial vestibule where the tobacco is held. The degree of mucosal alteration depends on the quality and type of tobacco, the frequency of tobacco use and host susceptibility. This entity has been observed in 60% of snuff dippers and 15% of chewing tobacco users. Chewing tobacco or snuff is usually started at 8-14 years of age. Smokeless tobacco keratosis takes about 5 years to develop. Epithelial dysplasia has been found in this lesion and the risk of developing OSCC is four times greater in smokeless tobacco users compared to non-users. In 98% of patients, breaking the habit leads to a normal mucosa within 2 to 6 weeks. A lesion that remains more than 6 weeks should be considered a true leukoplakia or carcinoma and should be biopsied.\textsuperscript{4,31-33}

**Palatal Keratosis associated with Reverse Smoking**

Reverse smoking is seen in patients of low socioeconomic class and is more common in females (91.3%), especially after the third decade of life. The frequency of oral involvement is high (97.8%), and the most commonly affected areas are the palate and the tongue.\textsuperscript{34} According to Barie et al\textsuperscript{35} the risk of malignant transformation in oral lesions of reverse smokers was 19 times more than that of nonsmokers. In addition, Alvarez-Gómez et al\textsuperscript{34} found epithelial dysplasia and OSCC in 83.3% and 12.5% of reveres smokers, respectively.

**Verrucous Hyperplasia**

Verrucous hyperplasia (VH) was first described as an extensive, thick, white plaque by Shear and Pindborg\textsuperscript{36} in 1980. There are histopathological differences between VH and verrucous carcinoma. VH is characterized by the thickened epithelium with superficial to adjacent normal epithelium, whereas verrucous carcinoma is characterized by an invaded pushing border of the hyperplastic epithelium to connective tissue, but the basement membrane is intact. In addition, Zhu et al\textsuperscript{37} demonstrated that VH located on the lower lip had a higher risk of malignant transformation compared with other sites. Despite the rarity of VH in Western countries, it is a
quite common oral premalignant lesion in Taiwan. The approved treatment is surgical excision. However, photodynamic therapy has also been recommended.48

**Oral Verrucous Carcinoma (Snuff Dipper’s Cancer, Ackerman’s Tumor)**

Oral verrucous carcinoma (OVC) is a low-grade variant of OSSC, which was first described in 1984. It has been demonstrated that 1–10% of all OSCCs are OVC. The incidence rate of OVC has been estimated as one oral lesion per 1,000,000 of the population each year. OVC predominantly occurs in patients with the habit of areca chewing, cigarette smoking and alcohol drinking. Malignant transformation has been found in 20% of OVC cases. Surgical excision, radiotherapy and chemotherapy have been recommended for treatment of OVC.4,38

**Dyskeratosis Congenita (Cole-Engman Syndrome or Zinsser-Cole-Engman Syndrome)**

Dyskeratosis congenita (DC) is a rare inherited disease, which is characterized by the classic triad of nail dystrophy, reticular skin pigmentation, and oral leukoplakia.39 DC may affect one in 1,000,000 of the population.40 It mainly occurs in men (male-to-female ratio of 13:1) and manifests between 5 and 13 years of age.39 Oral lesions, as white keratotic patches, occur in about 80% of cases. Typically, the buccal mucosa, tongue and oropharynx are affected. Ectodermal anomalies and neutropenia may lead to severe periodontal destruction. DC has an increased potential of malignant transformation.39-41 In a recent case report, Ray et al41 found malignant transformation on the patients’ tongue with DC. There is no effective treatment for DC. However, surgery and treatment with bleomycin, vitamin A, steroids and testosterone have been recommended for the management of this condition.39

**Actinic Cheilosis (Actinic Cheilitis)**

Actinic cheilosis (AC) is a premalignant condition frequently involving the lower lip. The main clinical presentations of AC are diffuse poorly-defined atrophic, erosive, ulcerative or keratotic plaques. Other clinical signs are dryness, scaly patches, swelling, transverse fissures, crusting and blotching.42,43 There is a strong likelihood of progressing to a malignancy. AC is one of the main causes of lip cancer, which is regarded as the 10 most common cancers in men. Six to ten percent of AC cases evolve into SCC over time.4,43,44

In a recent study, Kwon et al42 demonstrated that lip SCC originating from AC is more prone to metastasis than SCC arising from its cutaneous counterpart. Furthermore, they pointed out that the presence of bleeding, indurations, disease recurrence and persistent pain should be considered as markers of AC transformation into SCC.

There are various treatment methods for AC, including surgery, cryotherapy, electrosurgery, topical retinoids, 5-flourouracil cream, photodynamic therapy, CO2 laser ablation and vermilionectionomy.4,43,44

**Keratoacanthoma (Self-healing Carcinoma, Pseudocarcinoma, Keratoacarcinoma)**

Keratoacanthoma (KA) was first described in 1888. This entity appears as a firm, sessile, non-tender nodule with a central plug of keratin. It occurs more frequently on hair-bearing skin.45

The presence of this lesion was also reported on the lip and tongue by Ramos et al46 and Chen et al,47 respectively. According to Sanchez Yus et al48 at least 24% of KA showed malignant transformation into SCC. KA must be totally excised and studied for any evidence of malignancy.

**Oral Submucous Fibrosis**

Oral submucous fibrosis (OSF) is a chronic, progressive, scarring, precancerous condition which is characterized by mucosal rigidity. It has been associated with long-lasting dipping of betel-nut or paan in the mouth. OSF predominantly occurs in Indians and Southeast Asians. This condition is more common in young adults, aged 20-40. The most frequently affected sites in the oral cavity are buccal mucosa, retromolar area, tongue and soft palate. Clinical signs and symptoms of OSF are vesicles, ulcerations, petechiae, melanoses, xerostomia, burning sensation, and limitation in jaw movement.1,49 Overall, patients with OSF are at least 19 times more likely to develop OSCC than healthy people.4

According to a recent study by Guo et al50 OSSC originating from OSF occurs at a younger age (45.8 vs. 55.9 years), is more common in men (male-to-female ratio 32.1:1 vs. 2.3:1) and is more invasive and potentially metastatic than OSSC not originating from OSF. Surgical excision is an accepted treatment method for OSF. Furthermore, intra-lesional injections of interferon-γ have been recommended.5

**Oral Lichen Planus**

Lichen planus (LP) is a common chronic, immunologically mediated muco-cutaneous disease, which was first described in 1869. Most patients with lichen planus are middle-aged (over 40), and it is rare
in children. Females account for at least 65% of patients. Cutaneous lichen planus is seen in 1% and oral lichen planus (OLP) affects 0.1 - 2.2% of the population. Clinical variations of OLP are reticular, erosive, atrophic, bullous, ulcerative, papular, and plaque-like. The most commonly affected site in the oral cavity is posterior buccal mucosa, followed by tongue, (lateral and dorsal), gingivae, palate and vermilion border. The risk of malignant change in OLP has been controversial for a long time and reported to be between 0.4% and 3.7%. Patients often experience this complication after 10 years. Hence, OLP should be followed for a long period. Topical or systemic corticosteroids are usually recommended for the management of OLP.

Discoid Lupus Erythematosus

Discoid lupus erythematosus (DLE) is a chronic, scarring, immunologic, mucocutaneous disease, which is characterized by white keratinized plaques with elevated borders, radiating white striae and telangiectasia. The prevalence of DLE is less than 5 in 10,000 individuals and is more common in women than men with female to male ratio of 1.8:1. The frequency of oral involvement in DLE is lower than skin lesions, and is found in about 20% of cases. Malignant transformation of DLE into SCC is usually observed on the lower lip, and in Caucasians. However, there are some case reports in African Americans. Ma et al demonstrated that 5.5% of lower lip SCCs arise from lesions of DLE. This rate was reported to be 6.8% by Liu et al. According to Liu et al, epithelial dysplasia and age over 60 are risk factors for malignant transformation in DLE, whereas sex, location of the lesion, smoking and alcohol intake were not considered to be risk factors. Topical or systemic corticosteroids are usually recommended for management of DLE.

Epidermolysis Bullosa

Epidermolysis bullosa (EB) is a blistering disease of the skin and mucosa. Approximately 500,000 individuals are affected worldwide. There are three types of EB: simplex, junctional, and dystrophic. Each type consists of several forms of disorders. Oral involvement has been reported in the junctional and dystrophic forms of the disease, which is characterized by bulla and vesicle formation following mild physical trauma. Malignant transformation of EB has been reported most commonly in junctional and dystrophic types. In the oral cavity, lingual mucosa is more susceptible to malignant transformation. Malignancy usually develops in areas of chronic ulceration during the second to third decades of life. According to Yuen and Jonkman, developing a SCC among adults with junctional EB is noticed in 25% of cases. Risk of SCC in EB has also been documented by Oliveira et al. 60 and Yuen and Jonkman. Moreover, Fine and Mellerio showed that EB is even at risk for developing basal cell carcinoma (BCC) and malignant melanoma. Topical or systemic corticosteroids are usually recommended for management of EB.

Verruciform Xanthoma

Verruciform xanthoma (VX) is a rare benign lesion first described by Shafer in 1971. The lesions appear as a well-defined, asymptomatic, slightly elevated mass with a yellow-white or red color and a papillary or verruciform surface. It mainly occurs in the oral mucosa. The gingiva is the most commonly affected site, followed by tongue, buccal mucosa, vestibular mucosa, floor of the mouth, soft palate and lower lip. There is limited information about malignant transformation of VX. Mannes et al described association of VX with SCC and carcinoma in situ, respectively.

Verruciform xanthoma is treated with conservative surgical excision. Recurrence after removal of the lesion is rare.

Graft-versus-host Disease

Graft-versus-host disease (GvHD) is a common complication in patients treated with allogenic peripheral stem cell transplantation. Oral involvement has been observed in 80% of patients, characterized by atrophy, erythema, erosion, ulcers, lichenoid lesions, xerostomia, and oral pain. Malignant transformations in oral lesions have been documented by Salum et al and Montebugnoli et al.

Topical corticosteroids may facilitate the healing of focal oral ulcerations associated with GvHD. In some cases narcotic analgesics may be required.

Cheilitis Glandularis

Cheilitis glandularis (CG) is characterized by hyperplasia of minor salivary glands, first described by von Volkman in 1870. Frequently the lower lip is involved but lesions on the upper lip have also been reported. The main causes of CG are syphilis, bacterial infection, solar radiation, tobacco, poor oral hygiene and genetic factors. Transformation of CG into SCC has been found by Nico et al. Generally, a significant percentage of patients...
(18–36%) have been associated with the development of SCC of overlying epithelium of the lip.75

Xeroderma Pigmentosum

Xeroderma pigmentosum (XP) is a rare autosomal recessive disease, which is associated with cutaneous malignant transformation. Oral involvement is usually observed in patients under 20.4 Squamous cell carcinoma of the lower lip and tongue has been reported in patients with XP by Chidzonga76 and Patelttla.77 Consequently, XP should be considered an oral premalignant disorder. Patients with XP are advised to avoid sunlight and unfiltered fluorescent light, and to wear appropriate protective clothing and sunscreens if they cannot avoid sun exposure.4

Syphilis (Third Stage), Iron Deficiency Anemia (Plummer-Vinson or Patterson-Kelly Syndrome), Malnutrition, Vitamin A, B, and C Deficiency and Immunosuppressive Diseases (HIV)

Although the above conditions have been mentioned with malignant potential, data regarding their malignant transformation is controversial. However, they should be considered as predisposing factors of oral cancers.78-80

Conclusion

Most cancers and their related complications are preventable through early detection. Unfortunately, oral premalignant disorders are usually misdiagnosed due to lack of adequate knowledge among the general population and even medical professionals. Therefore, improvement of physicians’ or dentists’ level of knowledge about oral PMDs may play a key role in saving patients’ lives.

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