Oral candidiasis
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Oral candidiasis is a common opportunistic infection of the oral cavity caused by an overgrowth of *Candida* species, the commonest being *Candida albicans*. The incidence varies depending on age and certain predisposing factors. There are three broad groupings consisting of acute candidiasis, chronic candidiasis, and angular cheilitis. Risk factors include impaired salivary gland function, drugs, dentures, high carbohydrate diet, and extremes of life, smoking, diabetes mellitus, Cushing’s syndrome, malignancies, and immunosuppressive conditions. Management involves taking a history, an examination, and appropriate antifungal treatment with a few requiring samples to be taken for laboratory analysis. In certain high risk groups antifungal prophylaxis reduces the incidence and severity of infections. The prognosis is good in the great majority of cases.

Oral candidiasis is an opportunistic infection of the oral cavity. It is common and underdiagnosed among the elderly, particularly in those who wear dentures and in many cases is avoidable with a good mouth care regimen. It can also be a mark of systemic disease, such as diabetes mellitus and is a common problem among the immunocompromised. Oral candidiasis is caused by an overgrowth or infection of the oral cavity by a yeast-like fungus, *candida*. The important ones are *C. albicans* (the commonest; see fig 1), *C. tropicalis*, *C. glabrata*, *C. parapsilosis*, and *C. stellatoidea*. *C. albicans*, *C. glabrata*, and *C. tropicalis* represent more than 80% of isolates from clinical infection. Oral candidiasis is the most common human fungal infection especially in early and later life. In the general population, carriage rates have been reported to range from 20% to 75% without any symptoms. The incidence of *C. albicans* isolated from the oral cavity has been reported to be 45% in neonates, 45%–65% of healthy children, 30%–45% of healthy adults, 50%–65% of people who wear removable dentures, 65%–88% in those residing in acute and long term care facilities, 90% of patients with acute leukaemia undergoing chemotherapy, and 95% of patients with HIV. *C. albicans* is a normal commensal of the mouth and generally causes no problems in healthy people. Overgrowth of candida, however, can lead to local discomfort, an altered taste sensation, dysphagia from oesophageal overgrowth resulting in poor nutrition, slow recovery, and prolonged hospital stay. In immunocompromised patients, infection can spread through the bloodstream or upper gastrointestinal tract leading to severe infection with significant morbidity and mortality. Systemic candidiasis carries a mortality rate of 71% to 79%.

It is important for all physicians looking after older patients to be aware of the risk factors, diagnosis, and treatment of oral candidiasis. In a recent study 30% of doctors said they would prescribe nystatin for oral candidiasis on the request of nursing staff without examination of the oral cavity. This is unfortunate as other pathology may be missed, the diagnosis may be incorrect, and failure to address risk factors may lead to recurrence of the candidiasis.

**CLASSIFICATION**

There are a number of different types of oropharyngeal candidiasis including acute pseudomembranous, acute atrophic, chronic hyperplastic, chronic atrophic, median rhomboid glossitis, and angular cheilitis. The most discrete lesion represents conversion from benign colonization to pathological overgrowth.

**Pseudomembranous candidiasis (thrush)** is characterized by extensive white pseudomembranes consisting of desquamated epithelial cells, fibrin, and fungal hyphae (see fig 2). These white patches occur on the surface of the labial and buccal mucosa, hard and soft palate, tongue, periodontal tissues, and oropharynx. The membrane can usually be scraped off with a swab to expose an underlying erythematous mucosa. Diagnosis is usually straightforward as it is easily seen and is one of the commonest forms of oropharyngeal candidiasis accounting for almost a third. Diagnosis can be confirmed microbiologically either by staining a smear from the affected area or by culturing a swab from an oral rinse. Predisposing factors include extremes of age, diabetes mellitus, patients who have HIV/AIDS or leukaemia, those using steroid aerosol inhalers, broad spectrum antibiotics, and psychotropic drugs, and patients who are terminally ill. Other conditions that can give rise to white patches in the mouth are lichen planus, squamous cell carcinoma, lichenoid reaction, and leukoplakia.

**Acute atrophic candidiasis** is usually associated with a burning sensation in the mouth or on the...
tongue. The tongue may be bright red similar to that seen with a low serum B12, low folate, and low ferritin. Diagnosis may be difficult but should be considered in the differential diagnosis of a sore tongue especially in a frail older patient with dentures who has received antibiotic therapy or who is on inhaled steroids. A swab from the tongue/buccal mucosa may help diagnosis.

**Chronic hyperplastic candidiasis** characteristically occurs on the buccal mucosa or lateral border of the tongue as speckled or homogenous white lesions (see fig 3). The lesions usually occur on the buccal mucosa or lateral borders of the tongue. There is an association with smoking and complete resolution appears to be dependent on cessation of smoking. This condition can progress to severe dysplasia or malignancy and is sometimes referred to as candidal leukoplakia. *Candida* spp are not always isolated from lesions of oral leukoplakia and it has been suggested that the finding of *Candida* spp in these premalignant lesions is a complicating factor rather than a causative one. This condition may be confused with lichen planus, pemphigoid/pemphigus, and squamous cell carcinoma.

**Chronic atrophic candidiasis** also known as “denture stomatitis” is characterised by localised chronic erythema of tissues covered by dentures. Lesions usually occur on the palate and upper jaw but may also affect mandibular tissue. Diagnosis requires removal of dentures and careful inspection; swabs may be taken for confirmation. It is quite common with incidence rates of up to 65% reported.

**Median rhomboid glossitis** is a chronic symmetrical area on the tongue anterior to the circumvallate papillae. It is made up of atrophic filiform papillae. Biopsy of this area usually yields *candida* in over 85% of cases. It tends to be associated with smoking and the use of inhaled steroids.

**Angular cheilitis** is an erythematous fissuring at one or both corners of the mouth (see fig 4), and is usually associated with an intraoral candidal infection. Other organisms implicated are staphylococci and streptococci. In the case of staphylococci the reservoir is usually the anterior region of the nostrils and spread to the angles of the mouth has been confirmed by phage typing. Facial wrinkling at the corners of the mouth and along the nasolabial fold especially in older people leads to a chronically moist environment that predisposes to this lesion. This wrinkling is worse in long term denture wearers because there is resorption of bone on which the dentures rest leading to a reduction in height of the lower face when the mouth is closed. Other factors implicated in the aetiology of this condition are iron deficiency anaemia and vitamin B12 deficiency.

**RISK FACTORS**

1. **Pathogen**

*Candida* is a fungus and was first isolated in 1844 from the sputum of a tuberculous patient. Like other fungi, they are non-photosynthetic, eukaryotic organisms with a cell wall that lies external to the plasma membrane. There is a nuclear pore complex within the nuclear membrane. The plasma membrane contains large quantities of sterols, usually ergosterol. Apart from a few exceptions, the macroscopic and microscopic cultural characteristics of the different candida species are similar. They can metabolise glucose under both aerobic and anaerobic conditions. Temperature influences
Oral candidiasis can be classified as follows:

1. Acute candidiasis
   - Acute pseudomembranous candidiasis (thrush).
   - Acute atrophic (erythematous) candidiasis.

2. Chronic candidiasis
   - Chronic hyperplastic candidiasis (candidal leukoplakia).
   - Denture induced candidiasis (chronic atrophic (erythematous) candidiasis).
   - Median rhomboid glossitis.

3. Angular cheilitis (stomatitis)

Box 2: Classification

Oral candidiasis can be classified as follows:

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   - Denture induced candidiasis (chronic atrophic (erythematous) candidiasis).
   - Median rhomboid glossitis.

3. Angular cheilitis (stomatitis)

Box 3: Risk factors for oropharyngeal candidiasis

<table>
<thead>
<tr>
<th>Factors</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pathogen</td>
<td>Pathogen has peculiar properties that increase its infectivity rate in the right environment.</td>
</tr>
<tr>
<td>Host factors</td>
<td>Host factors could be local and/or systemic.</td>
</tr>
<tr>
<td>Local factors</td>
<td>Local factors include wearing dentures, impaired salivary gland function, inhaled steroids, and oral cancer.</td>
</tr>
<tr>
<td>Systemic factors</td>
<td>Systemic factors include extremes age, smoking, diabetes mellitus, Cushing’s syndrome, immunosuppression, malignancies, nutritional deficiencies, and antibiotics.</td>
</tr>
</tbody>
</table>

Phenotypic switching of Candida albicans is associated with certain pathogenic variables. Adhesion of candida to epithelial cell walls, an important step in initiation of infection, is promoted by certain fungal cell wall components such as mannose, C3d receptors, mannoprotein, and saccharins. The degree of hydrophobicity has also been reported to be important in the initial stages of infection. Other factors implicated in germ tube formation, presence of mycelia, persistence within epithelial cells, endotoxins, induction of tumour necrosis factor, and proteinases. Phenotypic switching which is the ability of certain strains of C albicans to change between different morphologic phenotypes has also been implicated.

(2) Host

Local factors

Impaired salivary gland function can predispose to oral candidiasis. Secretion of saliva causes a dilutional effect and removes organisms from the mucosa. Antimicrobial proteins in the saliva such as lactoferrin, sialoperoxidase, lysozyme, histidine-rich polypeptides, and specific antischistosoma antibodies, interact with the oral mucosa and prevent overgrowth of candida. Therefore conditions such as Sjögren's syndrome, radiotherapy of the head and neck, or drugs that reduce salivary secretions can lead to an increased risk of oral candidiasis.

Drugs such as inhaled steroids have been shown to increase the risk of oral candidiasis by possibly suppressing cellular immunity and phagocytosis. The local mucosal immunity reverts to normal on discontinuation of the inhaled steroids. Dentures predispose to infection with candida in as many as 65% of elderly people wearing full upper dentures. Wearing of dentures produces a microenvironment conducive to the growth of candida with low oxygen, low pH, and an anaerobic environment. This may be due to enhanced adherence of Candida spp to acrylic, reduced saliva flow, and the nature of the denture fittings, improperly fitted dentures, or poor oral hygiene.

Other factors are oral cancer/leukoplakia and a high carbohydrate diet. Growth of candida in saliva is enhanced by the presence of glucose and its adherence to oral epithelial cells is enhanced by a high carbohydrate diet.

Systemic factors

Extremes of life predispose to infection because of reduced immunity. Drugs such as broad spectrum antibiotics alter the local oral flora creating a suitable environment for candida to proliferate. The normal oral flora is restored once the antibiotics are discontinued. Immunosuppressive drugs such as the antineoplastic agents have been shown to predispose to oral candidiasis by altering the oral flora, disrupting the mucosal surface and altering the character of the saliva. Ninané found that 15–60% of people with malignancies will develop oral candidiasis while they are immunosuppressed.

In those with HIV infection rates of between 7% to 48% have been quoted and more than 90% has been reported in those with advanced disease. Relapse rates are between 30% and 50% on completion of antifungal treatment in severe immunosuppression.

MANAGEMENT

Taking a history followed by a thorough examination of the mouth, looking at the soft and hard palate, and examining the buccal mucosa in those wearing dentures after they have been removed are usually good starting points. Predisposing factors are identified as mentioned above and resolved if possible, and the type, severity, and chronicity of the infection are assessed.

The right diagnosis is usually made on finding the characteristic lesion, ruling out other possibilities, and the response to antifungal treatment. Acute pseudomembranous and chronic atrophic candidiasis can be treated based on clinical features but culture and sensitivity testing should be undertaken if initial therapy is unsuccessful. Imprint cultures, where sterile foam pads dipped in Sabouraud's broth are placed for 30 seconds on the lesion and then placed on Sabouraud's agar containing chloramphenicol for an hour after which they are incubated, have also been used for identification of Candida spp. Acute atrophic and chronic hyperplastic forms may mimic other lesions and a biopsy is recommended in addition to empirical therapy to rule out more serious lesions such as squamous cell carcinoma.

Oral hygiene and topical antifungals are usually adequate for uncomplicated oral candidiasis.

Oral hygiene involves cleaning the teeth, buccal cavity, tongue, and dentures, if present, daily. Dentures should be cleaned and disinfected daily and left out overnight or for at least six hours daily. The dentures should be soaked in a denture cleaning solution such as chlorhexidine which is more effective in eliminating candida than brushing. This is because dentures have irregular and porous surfaces to which candida easily adheres and brushing alone cannot remove them. When rinsing the mouth with the topical antifungal, dentures should be removed to allow contact between the mucosa and the antifungal. The patient should ensure that the
whole mucosa is coated with the antifungal and held in the mouth for a few minutes. The incorporation of an antifungal with a denture liner has been recommended for patients with dentures who find it difficult to hold the antifungal in their mouth for a few minutes. Also the mucosal surface should be brushed regularly with a soft brush. After disinfection, dentures should be allowed to air dry as this also kills adherent candida on dentures. Chlorhexidine can discolour both dentures and natural dentition if not removed adequately after disinfection. A referral to a dentist might be necessary for those with poorly fitting dentures as these predispose to infection by breaking down the epithelial barrier. Other conditions where topical antifungal agents are used are those with poorly fitting dentures as these predispose to infection by breaking down the epithelial barrier. Other denture cleaning methods not routinely used but shown to be effective are ultrasonic cleaning tanks with a suitable solution.

Regular oral and dental hygiene with periodic oral examination will prevent most cases of oral candidiasis in those with dentures. Combining nystatin with chlorhexidine digluconate, an antiseptic used to disinfect dentures, inactivates both drugs. Therefore this combination should not be used. The dentures should be removed each time the mouth is rinsed with the oral antifungal preparation in established cases of recurrent infection. The systemic adverse effects and drug interactions that occur with the systemic agents do not occur with topical agents.

Treatment in the early part of the 20th century was with gentian violet, an aniline dye, but because of resistance developing and side effects, such as staining of the oral mucosa, it was replaced by a polyene antibiotic, nystatin, discovered in 1951 and amphotericin B, discovered in 1956. They act by binding to sterols in the cell membrane of fungi and, altering cell membrane permeability.

Nystatin and amphotericin B are not absorbed from the gastrointestinal tract and are used by local application in the mouth. Miconazole, an imidazole, can be used as a topical cream and many preparations are available. The incorporation of an antifungal agent reduces the dose and duration of systemic treatment required. The systemic adverse effects and drug interactions that occur with the systemic agents do not occur with topical agents. Treatment in the early part of the 20th century was with gentian violet, an aniline dye, but because of resistance developing and side effects, such as staining of the oral mucosa, it was replaced by a polyene antibiotic, nystatin, discovered in 1951 and amphotericin B, discovered in 1956. They act by binding to sterols in the cell membrane of fungi and, altering cell membrane permeability.

Systemic antifungal therapy in oral candidiasis is appropriate in patients intolerant of or refractory to topical treatment and those at high risk of developing systemic infections.

Both nystatin oral rinses and clotrimazole troches have a high success rate and if tooth decay is a concern patients with oral candidiasis complicated by diabetes, steroid use or an immunocompromised state, triazoles which include fluconazole or itraconazole once per day has been found to be effective in these cases. Ketoconazole is also as effective as fluconazole and itraconazole but its use in elderly patients is not recommended due to drug interactions and side effects, which include hepatotoxicity.

Fluconazole is a potent and selective inhibitor of fungal enzymes involved in the synthesis of ergosterol, an important constituent of the plasma cell membrane. It therefore disrupts cell wall formation leading to leakage of cellular contents and cell death. It is well absorbed by the gastrointestinal tract and the plasma levels are over 90% of the levels achieved with intravenous administration and the levels in saliva and sputum are also similar to that in the plasma. It is preferred, as it does not have the same hepatotoxicity as the imidazoles. It is now listed in the dental practitioners’ formulary as well as the British National Formulary and is therefore widely used both in dental as well as medical practice but there are problems with resistance.

Itraconazole has a wider spectrum of activity than fluconazole and is therefore valuable in salvage treatment of the immunocompromised patients with fluconazole resistant candidiasis. Increasing resistance to antifungals has become increasingly common since the introduction of fluconazole especially in patients with advanced HIV disease, and recurrent and long term treatment.

Angular cheilitis is treated with antifungal steroid creams and ointments and any concurrent intraoral lesion is also treated at the same time and dietary deficiencies should be excluded and treated if found.

Failure to respond to therapy especially in chronic atrophic candidiasis is usually due to non-compliance with treatment. Prophylaxis with antifungal agents reduces the incidence of oral candidiasis in patients with cancer undergoing treatment and fluconazole has been found to be more effective than topical polyenes.

Prophylaxis on either a daily or weekly basis with antifungals reduces the incidence of oral candidiasis in patients with HIV with the reductions being most marked in those with low CD4 counts and recurrent oral candidiasis. The use of a chlorhexidine rinse only in bone marrow transplant patients as prophylaxis was found to be very effective.

PROGNOSIS

The prognosis is good for oral candidiasis with appropriate and effective treatment. Relapse when it occurs is more often than not due to poor compliance with therapy, failure to remove and clean dentures appropriately, or inability to resolve the underlying/predisposing factors to the infection.

Box 4: Management

- Diagnosis is usually made on clinical grounds with laboratory testing to exclude potentially other serious oral lesions especially squamous cell carcinoma.
- Predisposing factors should be treated or eliminated where feasible.
- Good oral hygiene is important.
- Topical antifungals given for two weeks are usually effective.
- Systemic antifungals should be given in certain circumstances.