Fordyce's Disease Treated with Pimacrolimus: A Rare Case Report

*T Hoque1, AZMM Islam2

ABSTRACT
Fordyce's disease, a rarely found disease of lips has been reported recently in department of Skin and VD, Gonosashthaya Somaj Vittic Medical College Hospital, Savar, Dhaka. Occasionally it may not be possible to identify the cause. The patient presented with identical features of Fordyce's disease and lip biopsy for histopathology showed the features of Fordyce's disease. Then patient was treated with Pimecrolimus cream and improved. Fordyce's disease is an extremely rare disorder. So its cutaneous findings, histopathology and treatments are highlighted here.

Key Words: Fordyce's disease, histopathology, Pimacrolimus.

Introduction
Fordyce's disease is ectopically located sebaceous glands, over mucous membrane of the mouth and lips1,14, characterized by the presence of whitish or yellowish, scanty or abundant, discrete, aggravated and often coalescent milium like bodies.2 It occurs commonly inside of the mouth laterally along the line of the teeth as far back as the last molar and possibly somewhat less frequently on the vermilion or mucous and inner surface of the lips, cheeks, less often glans penis, labia mejora and minora.1,15 The lesions are from 1mm to 3 mm in size and usually of a pale or oatmeal color.3 They are almost invariably imperceptible to touch, being situated on a level with the buccal mucosa, but at times they may send out hairy like projections which penetrates the mucous membrane. Patient usually unaware about the condition, as subjective symptoms are lacking.

Although the exact cause for Fordyce's condition is not known, yet the study suggest its generic connection.4 According to researchers, the heredity disorder can be the main factor behind the condition is nearly 40% of cases. Viral infection and overgrowing sebaceous glands can be the most common cause behind the condition. The development of yellowish papules and their location may indicate actopic sebaceous glands due to abnormal disposition during embryonic development.5 Sometimes warts are mistakenly diagnosed as Fordyce's condition, because of similarity in involvement of vermilion border in both cases.6 In cases the Fordyce's condition affected genital area that need for biopsy or blood examination considering the similarity in appearance with some of Sexually transmitted diseases. Details of Fordyce's condition perhaps have not yet been reported in our country. Here, a rare disease of sebaceous gland of lips, Fordyce's disease diagnosed clinically and histopathologically is reported.

Case Report
A 20 years young male came to the skin VD department of Gonosashthaya Somaj Vittic Medical College Hospital, Savar, Dhaka, with the complains of flat topped pinhead sized skin colored popular lesions over both upper and lower lips for 2 years (Fig 1), which
was painless and non-itching in types. He has no history of smoking, betel leaf or any other habit of smoking.

His family have no history of this complaint. He took medication of antifungal and antibacterial with local steroid ointment for 2 years but there was no improvement.

Physical and systemic examinations were essentially unremarkable. Local examination shows both lips were affected, no pain, no itching, only pinching sensation of lip mucosa. Hematological investigation with liver and kidney function tests were normal.

Skin biopsy from the lip showed the non-keratinized stratified squamous epithelium lining with minimal hyperplasia. The subepithelial area revealed lobulated sebaceous glands deep in the lamina propria (Fig 2). Each lobules consisted of polygonal cells with small nuclei and abundant clear cytoplasm. No hair follicle was observed. This findings similar with the diagnosis of Fordyce's disease.

The patient was treated with Pimecrolimus cream twice daily for 3 months and then once daily for further 3 months. After a month he had marked improvement of symptoms. By 3 months, the papules had flattened. The lesion found absent on follow up nine months after completing therapy.

**Fig-1:** flat topped pinhead sized skin colored popular lesions over both upper and lower lips.

**Fig-2:** non-keratinized stratified squamous epithelium lining with minimal hyperplasia. The subepithelial area revealed lobulated sebaceous glands deep in the lamina propria.

**Discussion**

Fordyce spots are named after the American dermatologist John Addison Fordyce (1858-1925) who first described these clinically in a medical journal. He also coined the terms Fox-Fordyce disease, Fordyce's disease, Fordyce's lesions and Brooke-Fordyce trichoepithelioma. Fordyce condition is noncontagious. Some serious cases of Fordyce's condition especially those of cosmetic concern may require treatment with vaporizing laser. The oral lesion that become frequent with the growing age can be seen in many elderly person. Even a healthy person can be affected by this condition. So much so that Fordyce's condition can involve several other medical condition. Fordyce's condition is commonly seen in men & women of any age group. According to a study, nearly 90% man are affected by Fordyce's condition. Some studies reported a male predilection or no significant difference in prevalence between male & female. According to Oliver, Fordyce's spots in selected South African population reach a peak between 20 & 29 years of age.

Fordyce's spot are a normal feature of sebaceous gland consisting of a single sebaceous lobule or gland located in the dermis or submucosa. The well formed lobule consist of small clusters of mature sebocytes with a sebaceous duct. This lesion is characterized by the presence of an opening directly onto the epithelial surface. However these Fordyce granules are actopic glands and have no connection
to hair follicle & open out directly. The vermilion border of upper lip is the most common site of lesion found on the lip.

Fordyce found parakeratosis & acanthosis, with cytoplasmic changes in the superficial strata of epidermis. The basal layer was normal. The protoplasm of many of prickle cells appeared granular and glistering. Perinuclear haloes were not infrequent.8 Lustgarten (quoted by Fordyce) considered that these changes corresponded to the normal granular metamorphosis of epidermis. D.W. Montgomery and Hay examined 2 cases of disease microscopically and found large number of apparently normal sebaceous glands underlying the affected area. These concluded that the yellowish coloration of lesion was due to subepidermal collection of fatty matter contained in these glands.14 CO2 Laser and oral isotretinoin can be considered as treatment options.9 However CO2 ablation can leave scars afterwards and isotretinoin cannot be taken for long periods of time. There is a report on 5-aminolevulenic acid photodynamic therapy for Fordyce’s spots.10 However side effects, such as a burning sensation, vesiculation and post inflammatory hyperpigmentation have been reported. Recently successful therapy that combines CO2 Laser ablation & topical trichloroacetic acid or bichloroacetic acid has been reported. Cauterization and cryosurgery causes extreme cold temperature to destroy the spots. A balanced diet comprising folic acid and vitamin C is necessary; an antioxidant helps build up the immune system.

Fordyce’s disease runs a prolonged course. As well as the discomfort associated with pruritis and the lesions are cosmetically dis-figuring. Case reports describe treatment with limited efficacy or associated with side effects the benefits of treatment. Treatments include topical clindamycin, corticosteroid, tretinoin, benzyl peroxide, oral isotretinoin, UV light, electrocoagulation, copper vapours, Laser & liposuction associated curettage. To date, there has been one previous case report demonstrating a successful response to eight weeks of Pimacrolimus in 3 female patients after unsuccessful treatment with topical corticosteroid and tretinoin.11 Tretinoin use regulates in order to ensure prevention of new lesion.12 The current case also has successful response and cure after 3 months use of the Pimacrolimus.

Pimacrolimus is currently use for mild to moderate atopic eczema. It is an ascomycin derivatives. It binds to cytosolic ligand receptor FK 506 binding protein. This complex inhibits the enzyme calcineurine phosphorylation of the cytoplasmic component of the nuclear factor of activated T-cell. The transcription of a number of inflammatory cytokines is therefore inhibited. This cytokines include IL-2, IFNγ, IL-4 and IL-10. Other cytokines such as IL-5 and TNF α are decreased in a dose dependent manner. Pimacrolimus also inhibits the transcription and synthesis of cytokines from mast cells and the release of performed mediators serotonin and β-l exosaminidase. Pharmacokinetic studies of Pimacrolimus have demonstrated negligible systemic absorption following topical application.13

**Conclusion**

Pimacrolimus is a relatively safe, easy to use option. So I suggest that it may be considered as first line therapy for Fordyce’s disease. Pimacrolimus was developed specifically for the treatment of inflammatory skin disease. Inflammation secondary to rupture of apocrine duct in Fordyce’s disease may be reduced by Pimacrolimus. There is no currently known mechanism to account for Pimacrolimus preventing the keratin plugging and initial obstruction of apocrine duct. However, Pimacrolimus may play a role in the view of a recent case of Pityriasis Rubra Pilaris, another keratinization disorder, completely clearing after use of Pimacrolimus.

**Conflict of interest:** We have no conflict of interest.

**References**


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