

## CASE REPORT

# A Case of Hand-Foot-Mouth Disease observed during Routine Dental Checkup in an Immunocompetent Healthy Adult Patient

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## ABSTRACT

Signs of oral lesions are one of the initial indications for many major diseases. As we come across different lesions during routine dental checkups, we should be aware about the diseases associated with those lesions. A 47-year-old immunocompetent adult patient reported with small erythematous lesions in the hard palate and small ulcers in the tonsillar pillars and right buccal mucosa with mild pyrexia of 37.8°C. Maculopapular lesions were found on the palms and soles. He was diagnosed with hand-foot-mouth disease, which is caused by the following viruses: Enterovirus-71, Coxsackie Virus (CV)-16, CV-A6, and CV-A10. The severity of this condition leads to meningitis, paralytic polio, and onychomadesis (falling of nails).

**Keywords:** Erythematous lesions, Hand-foot-mouth disease, Immunocompetent adult, Maculopapular lesions, Meningoencephalitis.

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## INTRODUCTION

Hand-foot-mouth disease (HFMD) was first described by Robinson et al<sup>1</sup> following an outbreak in Toronto in 1957. It is characterized by the appearance of vesicles on the mouth, hands, and feet. Coxsackie Virus (CV)-A16 was isolated. The disease rarely occurs as an epidemic, but in countries like China around 500,000 to 1 million cases are noted per year.<sup>2</sup> It usually occurs in early summer season,<sup>3</sup> but studies show a higher incidence rate in the temperature range of 21.1°C to 26.6°C,<sup>4</sup> with

greater predilection in children between the ages of 0 and 5 years<sup>4,5</sup> and in immunocompromised adults. As far as the sex predilection is concerned, there is greater incidence in males than females.<sup>4,6,7</sup> However, it can occur in immunocompetent adults<sup>5,8</sup> in rare instances, the key causative agents being enterovirus (EV)-71, CV-A16, and also CV-A6 and CV-A10.<sup>9-11</sup> Main routes of transmission are from person to person via oropharyngeal secretion or by direct vesicle contact. Feco-oral route and contaminated objects are also possible modes of transmission.<sup>12</sup> Incubation period ranges from 3 to 7 days starting with initial viral implantation in the oral cavity and ileum spreading to the regional lymph nodes within 24 hours. Viremia followed by pyrexia occurs within 72 hours followed by secondary infection and viral seeding in areas, such as oral mucosa, hand, and feet. Usually after a week, there is an increase in antibody levels and the disease begins to disappear.<sup>13</sup> Oral lesions are the first and usually the only clinical signs of the disease, so the dentists must be aware of this to quickly diagnose the condition. Only later do the lesions begin to appear in the extremities.<sup>12</sup>

Diagnosis is primarily made by observing the clinical signs of the disease, such as fever and characteristic lesions presenting in the oral cavity, hands, and feet. Confirmation of the diagnosis is carried out by isolating the virus responsible or by identifying virus-neutralizing antibodies in the patient's serum.<sup>5</sup>

The disease is mostly self-limiting and due to lack of a virus-specific therapy, the treatment is usually symptomatic. To relieve oral discomfort, an oral rinse with an anesthetic is prescribed along with an antipyretic for controlling fever. In addition, patients are advised plenty of fluids and to avoid spicy and acidic food. In most cases, prognosis is good with scabs or scars; however, there have been reports of onychomadesis (falling of nails)<sup>14-16</sup> associated with the disease. In some severe cases, there have been reports of neurological disorders, such as meningitis (EV-4), Guillain-Barré syndrome, meningoencephalitis as well as paralytic polio, myoclonus mainly in children under the age of 5 years (who are the main risk group of the population). It has also been demonstrated that these patients have low levels of vitamin A and reduced immunity.<sup>17</sup>

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**CASE REPORT**

A 47-year-old immunocompetent male reported to the clinic with a complaint of discomfort in the oral cavity and difficulty in swallowing food along with burning and itching sensation in the throat. The patient also had mild pyrexia of 37.8°C and general malaise at that time, but medical history seemed to be normal.

Intraoral examination revealed small erythematous lesions located on his hard palate and small ulcers on the tonsillar pillars and right buccal mucosa (Figs 1A and B). The examination also found maculopapular lesions on the palms of his hands and on the soles of his feet (Figs 2A and B), some of which were in the form of blisters

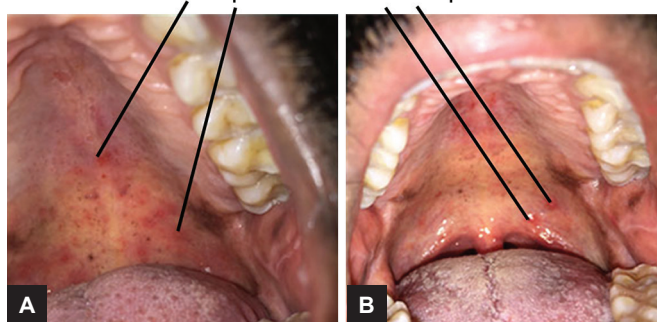
and upon further questioning, the patient stated that few days ago (about 8–10 days) his son aged 9 years suffered a similar condition and was examined and diagnosed with HFMD by the pediatrician. Based on this history, we came to a provisional diagnosis of the same condition and we took a detailed record with relevant photographs of the oral cavity, lesions of hand and feet, and also referred him to a medical specialist with this provisional diagnosis mentioned in the case sheet.

The medical specialist also confirmed the diagnosis of HFMD and treated him with an antipyretic (paracetamol 650 mg) and an antihistamine for 5 days and topical calamine lotion for his hands and feet. We called the patient after a week and observed that the oral lesions had completely subsided, but the hand and foot lesions were healing and the skin showed peeling with red discoloration in his palms and soles.

**CONCLUSION**

Hand-foot-mouth disease is a highly contagious disease mainly caused by EVs. The initial presentation includes erythematous papules on the palms, soles, and in the oral cavity accompanied by prodromal symptoms, such as myalgia, mild pyrexia, and abdominal pain. The lesion usually evolves into vesicles which are painful, covered by a yellowish pseudomembrane and surrounded by an erythematous halo measuring up to about 5 mm in diameter. Constant movement in the

Intraoral picture showing lesions on hard palate and tonsillar pillars



**Figs 1A and B:** (A) Intraoral view of lesions on hard palate; and (B) tonsillar pillars



**Figs 2A and B:** (A) Maculopapular lesions on hands; and (B) lesions on the soles of feet

oral cavity results in disruption of the membrane covering the lesion. Vesicles present in the skin can vary in number and usually appear on the sides and back of the fingers and soles of the feet. The condition is usually asymptomatic. Children and immunocompromised adults are more susceptible, but HFMD is rare in immunocompetent adults. In view of the differential diagnosis for this condition, the possible lesions to be considered are primary herpetic gingivostomatitis, herpangina, erythema multiforme, aphthous stomatitis, and varicella. Inappropriate drugs must not be administered without a correct diagnosis.<sup>18,19</sup> Diagnosis is mainly through clinical features along with histopathological examination of the vesicles. Recent diagnostic method studied by Yu et al<sup>20</sup> is by use of immunoglobulin M enzyme-linked immunosorbent assay in EV-71 and CV-A16 infection to correctly identify the virus causing the disease in patients. Onychomadesis need not be treated except by keeping the nail area clean, and in most cases, it is self-healing over a course of several weeks.<sup>16</sup> All studies correspond to EV-71 being the causative agent for manifestation of HFMD, and children in age group of 2 to 4 years are the prime risk group.

Vaccination has been a key in the prevention of this infection. The most recent vaccine for the year 2015 is formalin-inactivated EV-71 (FI-EV-71) developed and verified through human trials and found to provide adequate protection against EV-71, but nevertheless, there is no approved vaccine available for treating and preventing EV-71 infection. An important aspect of preventing the disease is that HFMD patients should be kept isolated until the fever and skin and mucosal lesions have resolved as this stage of the disease is potentially contagious. Disease spread also occurs through sharing utensils or through feces and oronasal secretion, so frequent handwashing should be given importance.

Emphasis is to be laid on the fact that dentists play an important role in identifying this condition earlier than other specialists and awareness must be given to patients to visit the dentist for periodic checkups, which aid in early identification of the condition, thereby potentially preventing its spread to the community as the condition is contagious and the patient must be educated to prevent spread of the disease to other members of his/her family.

## REFERENCES

1. Robinson CR, Doane FW, Rhodes AJ. Report of an outbreak of febrile illness with pharyngeal lesions and exanthema: Toronto, summer 1957; isolation of group A Coxsackie virus. *Can Med Assoc J* 1958 Oct 15;79(8):615-621.
2. Wang JF, Guo YS, Christakos G, Yang WZ, Liao YL, Li ZJ, Li XZ, Lai SJ, Chen HY. Hand foot and mouth disease: spatiotemporal transmission and climate. *Int J Health Geogr* 2011 Apr 5;10:25.
3. Blomqvist S, Klemola P, Kaijalainen S, Paananen A, Simonen ML, Vuorinen T, Roivainen M. Co-circulation of coxsackieviruses A6 and A10 in hand, foot and mouth disease outbreak in Finland. *J Clin Virol* 2010 May;48(1):49-54.
4. Wang Y, Feng Z, Yang Y, Self S, Gao Y, Longini IM, Wakefield J, Zhang J, Wang L, Chen X, et al. Hand, foot, and mouth disease in China: patterns of spread and transmissibility. *Epidemiology* 2011 Nov;22(6):781-792.
5. Zhang X, Yan HP, Huang C, Tan YF, Ma DM, Zhang HP, Liu Y, Wang SZ. The etiology and clinical manifestations of 70 patients with hand-foot-mouth disease. *Zhonghua Yu Fang Yi Xue Za Zhi* 2009 Oct;43(10):872-874.
6. Zeng M, Li YF, Wang XH, Lu GP, Shen HG, Yu H, Zhu QR. Epidemiology of hand, foot, and mouth disease in children in Shanghai 2007–2010. *Epidemiol Infect* 2012 Jun;140(6):1122-1130.
7. Yan XF, Gao S, Xia JF, Ye R, Yu H, Long JE. Epidemic characteristics of hand, foot, and mouth disease in Shanghai from 2009 to 2010: enterovirus 71 subgenotype C4 as the primary causative agent and a high incidence of mixed infections with Coxsackie virus A16. *Scand J Infect Dis* 2012 Apr;44(4):297-305.
8. Shin JU, Oh SH, Lee JH. A case of hand-foot-mouth disease in an immunocompetent adult. *Ann Dermatol* 2010 May;22(2):216-218.
9. Kaminska K, Martinetti G, Lucchini R, Kaya G, Mainetti C. Coxsackie virus A6 and hand, foot and mouth disease: three case reports of familial child-to-immunocompetent adult transmission and a literature review. *Case Rep Dermatol* 2013 Aug;5(2):203-209.
10. Stewart CL, Chu EY, Introcaso CE, Schaffer A, James WD. Coxsackie virus A6-induced hand-foot-mouth disease. *JAMA Dermatol* 2013 Dec;149(12):1419-1421.
11. Harris PNA, Wang AD, Yin M, Lee CK, Archuleta S. Atypical hand, foot, and mouth disease: eczema coxsackium can also occur in adults. *Lancet Infect Dis* 2014 Nov;14(11):1043.
12. Park SK, Park B, Ki M, Kim H, Lee K, Jung C, Sohn YM, Choi SM, Kim DK, Lee DS, et al. Transmission of seasonal outbreak of childhood enteroviral aseptic meningitis and hand-foot-mouth disease. *J Korean Med Sci* 2010 May;25(5):677-683.
13. Delgado-Azañero W, Concha-Cusihuallpa H, Guevara-Canales JO. Infección de la mucosa oral por coxsackie virus: enfermedad de boca mano pie. *Rev Estomatol Herediana* 2007;17(1):35-39.
14. Davia JL, Bel PH, Ninet VZ, Bracho MA, González-Candelas F, Salazar A, Gobernado M, Bosch IF. Onychomadesis outbreak in Valencia, Spain associated with hand, foot, and mouth disease caused by enteroviruses. *Pediatr Dermatol* 2011 Jan-Feb;28(1):1-5.
15. Guimbao J, Rodrigo P, Alberto MJ, Omeñaca M. Onychomadesis outbreak linked to hand, foot, and mouth disease, Spain, July 2008. *Euro Surveill* 2010 Sep;15(37):19663.
16. Wei SH, Huang YP, Liu MC, Tsou TP, Lin HC, Lin TL, Tsai CY, Chao YN, Chang LY, Hsu CM. An outbreak of coxsackievirus A6 hand, foot, and mouth disease associated with

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- onychomadesis in Taiwan, 2010. *BMC Infect Dis* 2011 Dec 14; 11:346.
17. Chen S, Yang Y, Yan X, Chen J, Yu H, Wang W. Influence of vitamin A status on the antiviral immunity of children with hand, foot and mouth disease. *Clin Nutr* 2012 Aug;31(4):543-548.
  18. Muppa R, Bhupatiraju P, Duddu M, Dandempally A. Hand, foot and mouth disease. *J Indian Soc Pedod Prev Dent* 2011 Apr-Jun;29(2):165-167.
  19. Na SY, Son YM, Lee HY, Baek JO, Roh JY, Lee JR. A case of varicella combined with hand-foot-mouth disease in a healthy child. *Ann Dermatol* 2009 Feb;21(1):98-101.
  20. Yu N, Guo M, He SJ, Pan YX, Chen XX, Ding XX, Hao W, Wang YD, Ge SX, Xia NS, et al. Evaluation of human enterovirus 71 and coxsackievirus A16 specific immunoglobulin M antibodies for diagnosis of hand-foot-and-mouth disease. *Virol J* 2012 Jan 11;9:12.