A Case of Giant Molluscum Contagiosum in an Immunocompetent Child

Keywords: Giant molluscum contagiosum; Immunocompetent; Child

A 4-year-old boy presented with an asymptomatic, erythematous mass on the abdomen. His mother said that the lesion developed a few months ago and has been gradually increasing in size. His weight was 19 kg, height was 95 cm and showed normal development. The patient did not show any signs of immunodeficiencies. The past history and family history was unremarkable. On close examination, a solitary, 0.5 cm-sized, pedunculated, erythematous mass lesion was present on the abdomen (Figure 1). Multiple, tiny, umbilicated, skin-colored papules were also noticed on the trunk and extremities. Punch biopsy was performed at the erythematous lesion. Histopathologic examination showed lobular epithelial hyperplasia with central keratinization. Diffuse lymphocytic infiltration and intracytoplasmic inclusion bodies were observed, which was consistent with molluscum contagiosum (Figure 2).

Molluscum contagiosum is a self-limiting skin infection caused by a member of the molluscum contagiosum virus (MVCV1-4), the poxvirus family. Incidence peaks in pre-school children and reports of infection have also been made in immunocompromised adults. The virus directly enters into the epidermis from skin defect or can be disseminated by simple contact. The disease is self-limiting in immunocompetent individuals but severe and prolonged cases have been reported in Human Immunodeficiency Virus (HIV) infected patients [1]. Clinically, it is characterized by 1 to 3 mm-sized, skin-colored, umbilicated papules that usually appear on the face, trunk, and limbs [2]. There is typical central umbilication from which cheesy keratinous material can be expressed. The incubation period is around 2 weeks. Widespread, multiple MC lesions are more common in children. Atypical presentations, such as facial, large, hyperkeratotic lesions can be seen and are more frequent in immunocompromised adults.

Molluscum contagiosum with a diameter greater than 0.5-1 cm is classified as a ‘Giant MC’. The giant atypical MC rarely occurs in healthy individuals. The giant variant of MC can be a single lesion and as they mostly occur at the eyelids, scalp, and soles, they can be easily confused with other conditions like basal cell carcinoma, abscess, furuncle, and various adenomas. They usually occur in conditions with altered immunity as atopic dermatitis, patients on immune-suppressive therapy, leukemias, Wiskott-Aldrich syndrome and AIDS [3].

The diagnosis of MC is usually made based on the typical clinical findings. However, atypical forms may require biopsy to confirm the diagnosis. The treatment does not differ the classic forms of MC. Surgical methods including curettage, electrodessication, cryotherapy, and laser surgery are often adopted. Topical cytodestructive agents including cantharadin, iodine, lactic acid, tretinoin, trichloroacetic acids and systemic chemotherapeutic agents including cidofovir, interferon, imiquimod can be considered as alternative treatment options [2]. In our case, the giant MC lesion was completely removed...
with a 6 mm punch biopsy and there has been no sign of recurrence at 6 months follow-up.

Solitary giant MC, although a rare condition in an immunocompetent patient, should be suspected if typical clinical presentations such as a central umbilication is found. Complete removal with a punch and concurrent biopsy is considered as one of the best diagnostic and therapeutic option for giant MC, especially when the diagnosis is uncertain.

References


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