

presented with skin lesions on his head was admitted to our hospital on 22 September 2009. Thirteen years ago, an indurated papule appeared on his scalp, without pain and itch. From then on, the same papules spread peripherally, and became confluent. Similar lesions involved his nose, chest, and both lower extremities gradually. The lesions were skin colored and hairless. However, his intelligence was normal. Six years ago, he underwent skull drilling to clear subdural hematoma following brain trauma. Otherwise, he denied any other medical conditions such as chronic eczema, acne, pyoderma, psoriasis, endocrine diseases, or other tumors. The patient was the product of a non-consanguineous marriage. His mother had no history of exposure to drugs, chemical materials, or radiation during her pregnancy and lactation. There was no family history of similar illness.

The patient was well developed. The size of his head was in normal range. Scattered papules could be seen on his scalp, face [Figure 1], chest, and ankle, which were 0.1-0.5 cm in diameter, skin colored, and hard on palpation. On his head, there was a well circumscribed hairless giant tumor (15×9×0.5 cm). He was otherwise healthy. The plain CT scan of skull showed multiple calcifications in the scalp, especially the right frontal part and parietal part; the lesion hardly adhered to the skull. The other laboratory tests (including liver and renal function tests, calcium, phosphate, and parathormone level) were almost in the normal range. Under local anesthesia, a biopsy was performed from the edge of the lesion. The histopathological examination revealed that the epidermis was normal, and bone tissue including bone cells, bony trabeculae, and fatty tissue appeared in the entire dermis. Calcification also could be seen to some extent under microscopy. Osteoma cutis was diagnosed by the histopathology, so the patient was confirmed as a case of the primary osteoma cutis.

Under local anesthesia, a 300 ml skin tissue expander was embedded beneath the normal scalp adjacent to the giant tumor. The sterile saline water of 510 ml was infused during the 62 days after initial expander insertion [Figure 2]. The giant lesion of the head was resected after the tissue expander was removed on 29 December 2009 [Figure 3].

Osteoma cutis is a rare, benign disease that was first described by Virchow in 1864.<sup>[1]</sup> It is characterized by the formation of morphologically normal bone within the dermis or the subcutaneous tissue. There are two major classes of osteoma cutis: primary and

## A case of giant primary osteoma cutis successfully treated with tissue expansion and surgical excision

Sir,  
A 40-year-old Canadian Chinese male patient who



**Figure 1:** On his head, there was a well circumscribed hairless giant tumor (15×9×0.5 cm). Scattered papules could be seen on his scalp and face



**Figure 2:** A 300 ml skin tissue expander was embedded beneath the normal scalp adjacent to the giant tumor. The sterile saline water of 510 ml was infused during the 62 days after initial expander insertion



**Figure 3:** The giant lesion of the head was resected after the tissue expander was removed

secondary. The latter is more common than the former. It can be a sequel of multiple disorders including nevi, scleroderma, pilomatricoma, dermatomyositis, basal cells carcinoma, scars, inflammation, trauma, calcification, fibrous proliferations, and venous stasis.<sup>[2]</sup> The primary osteoma cutis is characterized by *de novo* bone formation in the skin without a known associated or pre-existing cutaneous disorder.<sup>[1]</sup> It has been described as four different clinical variants: solitary, widespread, plaque-like, and multiple military osteomas of the face.<sup>[3]</sup> The osteoma cutis may be single or multiple and have been reported on the scalp, face, trunk, breast, extremities, and buttocks.<sup>[4]</sup>

At present, the pathogenesis of primary osteoma cutis remains unclear. Diagnosis of osteoma cutis may be suspected clinically or radiographically, but can

be confirmed only by histopathology. Microscopic findings consist of well-formed and calcified bone arranged in spicules of spongy bone or sheets of compact bone. There are always identifiable osteocytes, and occasionally Haversian system. Howship's lacunes and osteoclasts are seen only rarely.<sup>[5]</sup>

The osteoma cutis is a cosmetically distressing problem with different treatment alternatives. These methods include non-invasive treatment such as tretinoin application or invasive treatments such as combinations of dermabrasion and punch biopsy, CO<sub>2</sub> laser, and erbium: YAG laser, scalpel incisions and curettage, and needle microincision extirpation.<sup>[1,4]</sup>

As in our case, the lesions on his scalp were up to 15×9 cm in size, so the treatment by surgical resection after skin tissue expansion was performed. The objective was to replace excised lesions with expanded adjacent scalp skin that had normal sensation and hair follicles. Fortunately, the scalp was ideal for skin tissue expansion in this case. The operation was pretty successful in not only removing the lesion, but also achieving the cosmetic results. It is beyond doubt that our case is the first described, indicating that it is a safe and effective approach to treat giant osteoma cutis with tissue expansion and surgical excision.

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Access this article online	
<b>Quick Response Code:</b>	<b>Website:</b> www.ijdvl.com
	<b>DOI:</b> 10.4103/0378-6323.75005

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