A Rare Case of Gorham's Disease of the Third Finger Managed by Surgical Reconstruction

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Abstract
A 35 year old male presented with pain in right middle finger since 5 y associated with shortening of right middle finger since 3 y. He had dull aching pain and blackish discoloration of the finger since 5 y. Examination revealed painful, restricted movements at inter-phalangeal joint and shortening of the finger of about 4 cm. After a thorough work up of the patient, he was diagnosed to have Gorham’s disease of the third metacarpal which was confirmed by histopathology reports. The patient was managed by surgical reconstruction to provide a functional and cosmetically acceptable hand.

Keywords: Gorham disease; Hand; Metacarpal; Proximal phalanx; Surgical reconstruction

Introduction
Gorham-Stout disease (vanishing or disappearing bone disease, phantom bone) is a rare disease of unknown aetiology that is characterized by massive osteolysis and excessive intra-osseous proliferation of small blood or lymphatic vessels, resulting in progressive resorption of bone [1]. The disease is often seen in children and young adults of either gender, and mainly affects bones that develop by intra-membranous ossification (shoulder girdle, pelvis, jaw, ribs, and spine). Approximately 200 cases of vanishing bone disease have been reported in the literature (Table 1). Numerous names have been used in the literature to describe this condition such as phantom bone disease, massive osteolysis, disappearing or vanishing bone disease and acute spontaneous absorption of bone.

Gorham and Stout presented the first overview of vanishing bone disease in 1955 and reported 24 cases [2]. They concluded progressive osteolysis in those cases was associated with angiomatosis of blood or lymphatic vessels. This is now known as Gorham disease. The aetiology remains speculative, the prognosis is unpredictable.

Case Report
A 35 year old male presented with pain in right middle finger since 5 y and shortening of middle finger since 3 yrs. He had dull aching pain and blackish discoloration of the finger since 5 y. Examination revealed painful, restricted movements at inter-phalangeal joint and shortening of the finger of about 4 cm (Figure 1).

Investigations
X-ray right hand
Showed absence of middle phalanx of middle finger and hazy borders of head of proximal phalanx.

Table 1: Sites of involvement of vanishing bone disease reported in the literature.

| Location                | No of cases |
|-------------------------|-------------|
| Skull                   | 8           |
| Maxillofacial           | 42          |
| Spine                   | 18          |
| Pelvis                  | 14          |
| Trunk (including clavicle and ribs) | 35 |
| Upper extremity (including scapula) | 41 |
| Lower extremity         | 22          |
| Multicentric            | 11          |
| Total                   | 191         |

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MRI right hand
Suggestive of vascular malformation of the middle finger of the right hand with non-visualization of the middle phalanx, anterior subluxation of the distal phalanx with flexion deformity of the digit, acro-osteolysis and soft tissue thickening (Figure 2).

Histopathology
Showed multiple sections of spicules of lamellar bone displaying many blood vessels in the bone marrow, stroma is loose with few occasional plasma cells and lymphocytes, which are suggestive of angiomatous hyperplasia (Gorham’s disease).

Diagnosis Gorham’s Disease of Right Middle Phalanx

Surgical reconstruction
Ray amputation of right middle finger, 3rd metacarpal with stump closure and reconstruction of the hand.

Incision:
1. A ‘Y’ shape incision was made over dorsum aspect of the right hand from 2nd and 3rd web space extending proximally,
2. ‘V’ shape incision given over the palmar aspect at the base of the 3rd metacarpal.

Soft tissue dissection:
Soft tissue released, phalanx explored and periosteum elevated till base of the 3rd metacarpal and excised extensor tendon at the same level. ‘V’ shape osteotomy done at base of the 3rd metacarpal and released distal part. Half of the interossei are excised on either side of the 3rd metacarpal and flexor digitorum profundus tendon was excised.

Fixation:
The retained halves of the interossei were approximated and sutured. A 2 mm K-wire passed from head of 5th metacarpal to the 2nd head of metacarpal in a transverse manner (Figure 3).

Immobilization:
Immobilised with below elbow slab.

Post-operative rehabilitation
3 weeks: K-wire removal
6 weeks: achieved full range of movements
3 months: 80% hand grip
6 months: 100% hand grip achieved (Figure 4).

Discussion
Vanishing bone disease is a rare idiopathic disease leading to extensive loss of bony matrix, which is replaced by proliferating thin-walled vascular channels and fibrous tissue [3]. Although the disease can be monostotic or polyostotic, multicentric involvement is unusual [4]. The patients whom we present had monostotic pattern of the disease.

Gorham’s disease may be associated with history of trivial trauma, although as many as half of the patients have no history of trauma. As with many other diseases, the role of trauma in vanishing bone disease may be to signal the presence of a pre-existing abnormality. Our patient also did not present with any history of trauma.
Most cases occur in children or in adults less than 40 years. The bones of the upper extremity and the maxillofacial region are the predominant osseous sites of the disease. Leriche’s hypothesis that post-traumatic arterial hyperaemia was responsible for bone resorption was rejected first by Mouchet, and later by Gorham et al. [5]. The same authors postulated that an angioma might act as a shunt increasing local oxygen tension [5]. The process may affect the appendicular or the axial skeleton. The shoulder and the pelvis are the most common sites of involvement. Gorhams disease affecting the hand is rare and we are presenting this case for its rarity.

Our search in English and international literature revealed only five cases of Gorham’s disease affecting the hand and wrist [2,6-9]. In all the described cases, the disease was unifocal and the phalanges were minimally affected (Table 2).

The diagnosis of vanishing bone disease is based on clinical examination, radiologic imaging studies and histo-pathological study of the affected bone. Vanishing bone disease is not accompanied by general symptoms. Dull aching, weakness in the affected extremity, swelling and skeletal deformities are the usual presenting symptoms [10]. The chief complaints presented by our patient were discolouration, shortening and dull aching pain of the third finger of the right hand since five years.

Radiographs are the best tools for detecting vanishing bone disease [11]. The radiographic appearance becomes diagnostic of vanishing bone disease when unilateral partial or total disappearance of contiguous bones, tapering of bony remnants, and absence of a sclerosing or osteoblastic reaction are observed. In our case, radiography revealed partial disappearance of the third metacarpal with absence of corresponding middle phalanx.

Histologically, the appearance depends on the phase in which the disease is diagnosed. In 1983, Heffez et al. [12] described two phases. The first phase represents increased vascular concentration in the bone-displacing fibrous tissue part; in the second phase, only fibrous tissue is found. The presence and number of osteoclasts vary significantly in vanishing bone disease. In most cases, osteoclastic activity is minimal or nonexistent, whereas in other cases, osteoclasts are easily identifiable [13]. If present, osteoclastic activity is concentrated in the interface between the vascular channels and the cortex [14]. Our patient’s biopsy report was in concurrence with the literature by the fact that there were multiple bone spicules with increased blood vessels in bone marrow. This suggests that there was increased vascularity (angiomatic hyperplasia) and was probably in the first phase of the disease.

The differential diagnosis should exclude other causes of osteolysis such as skeletal angiomas, essential osteolysis, hereditary osteolysis, infection, trauma (Sudeck atrophy), endocrinal abnormalities, rheumatoid arthritis and tumours [15].

Various modalities of treatment have been described for Gorhams disease. We chose to surgically reconstruct his hand because he wanted to perform his activities of daily living. His main requirements were good hand grip and pincer grip.

**Conclusion**

Gorham disease is a combined clinical, radiographic and histological entity. It is characterized by a nonfamilial, histologically benign vascular proliferation originating in bone and producing complete lysis of all or a portion of the bone [16]. Usually, the prognosis depends on complications, such as neurological deficits or pleural effusion. It has been reported that more than 15% of patients die as a result of this disease. Life expectancy is not affected if the extremities are involved. The treatment of vanishing bone disease is controversial. A review of the literature led to the conclusion that there is no consensus about the most efficacious treatment. Synergistic action of zoledronic acid and Interferon is a powerful antiangiogenic therapy, which is currently giving the best results [17-19].

Radiotherapy acts by accelerating sclerosis of the proliferating blood vessels and prevents regrowth of these vessels. Although the use of total doses from 30 Gy to 45 Gy has been reported to be effective, some authors reported excellent results while using a total dose of 15 Gy in a case that involved the upper extremity [20-23].

Surgical intervention has been suggested as the method of choice when feasible and involves local resection of the affected bone, with or without replacement prostheses or bone grafts [24].

**References**

1. Gorham LW, Stout AP (1955) Massive osteolysis (acute spontaneous absorption of bone, phantom bone, disappearing bone); its relation to hemangiomatosis. J Bone Joint Surg Am 37-37A: 985-1004.

2. Devin RD, Bone HG 3rd, Roodman GD (1996) Interleukin-6: a potential mediator of the massive osteolysis in patients with Gorham-Stout disease. J Clin Endocrinol Metab 81: 1893-1897.

3. Vélez A, Herrera M, Del Rio E, Ruiz-Maldonado R (1993) Gorham’s syndrome. Int J Dermatol 32: 884-887.

4. Loppnow H, Libby P (1990) Proliferating or interleukin 1-activated human blood vessels and prevents regrowth of these vessels. Although the use of total doses from 30 Gy to 45 Gy has been reported to be effective, some authors reported excellent results while using a total dose of 15 Gy in a case that involved the upper extremity [20-23].

5. Dickson GR, Hamilton A, Hayes D, Carr KE, Davis R, et al. (1990) An investigation of vanishing bone disease. Bone 11: 205-210.

6. Luhnow H, Libby P (1990) Proliferating or interleukin 1-activated human vascular smooth muscle cells secrete copious interleukin 6. J Clin Invest 85: 731-738.

7. Dickson GR, Hamilton A, Hayes D, Carr KE, Davis R, et al. (1990) An investigation of vanishing bone disease. Bone 11: 205-210.

8. Lehnhardt M, Steinau HU, Homann HH, Steinstraesser L, Druceck D (2004) [Gorham-Stout disease: report of a case affecting the right hand with a follow-up of 24 years]. Handchir Mikrochir Plast Chir 36: 249-254.
9. Tunon JB, Gonzalez FP (1977) Angiomatosis of the metacarpal skeleton. Hand. 9: 86-91.
10. Möller G, Priemel M, Ameling M, Werner M, Kuhlmey AS, et al. (1999) The Gorham-Stout syndrome (Gorham’s massive osteolysis). A report of six cases with histopathological findings. J Bone Joint Surg Br 81: 501-506.
11. Naden BA (1995) When bone disappears. RN 58: 26-28.
12. Heffez L, Doku HC, Carter BL, Feeney JE (1983) Perspectives on massive osteolysis. Report of a case and review of the literature. Oral Surg Oral Med Oral Pathol 55: 331-343.
13. Dickson GR, Mollan RA, Carr KE (1987) Cytochemical localization of alkaline and acid phosphatase in human vanishing bone disease. Histochemistry 87: 569-572.
14. Hvos AG (1980) Bone tumours, diagnosis, treatment and prognosis. J Clin Pathol 33: 415.
15. Holroyd I, Dillon M, Roberts GJ (2000) Gorham’s disease: a case (including dental presentation) of vanishing bone disease. Oral Surg Oral Med Oral Pathol Oral Radiol Endod 89: 125-129.
16. Høgborg H, Lamberg K, Aström G (1997) Alpha-2b interferon and oral doxorubicin for Gorham’s disease. Lancet 350: 1822-1823.
17. Hammer F, Kerr W, Wesselmann U, Hobfauer LC, Delling G, et al. (2005) Gorham-Stout disease—stabilization during bisphosphonate treatment. J Bone Miner Res 20: 350-353.
18. Mawrk JR, Obukhov SK, Nichols WD, Wynne TD, Odell JM, et al. (1997) Successful conservative management of Gorham disease of the skull base and cervical spine. Childs Nerv Syst 13: 622-625.
19. Fontanesi J (2003) Radiation therapy in the treatment of Gorham disease. J Pediatr Hematol Oncol 25: 816-817.
20. Dunbar SF, Rosenberg A, Mankin H, Rosenthal D, Sult HD (1993) Gorham’s massive osteolysis: the role of radiation therapy and a review of the literature. Int J Radiat Oncol Biol Phys 26: 491-497.
21. Hanly JG, Walsh NM, Bresnihan B (1985) Massive osteolysis in the hand and response to radiotherapy. J Rheumatol 12: 580-582.
22. Listewnik MJ, Marzeczi Z, Kordowski J (1986) Disappearing bone disease: massive osteolysis of the ribs treated with radiotherapy. Report of a case. Strahlenther Onkol 162: 338-341.
23. Grätz KW, Prein J, Remagen W (1987) [A reconstruction attempt in progressive osteolysis (Gorham disease) of the mandible. A case report]. Schweiz Monatschr Zahnmed 97: 980-984.
24. Picault C, Comtet JJ, Imbert JC, Boyer JM (1984) Surgical repair of extensive idiopathic osteolysis of the pelvic girdle (Jackson-Gorham disease). J Bone Joint Surg Br 66: 148-149.