Pathological Femoral Shaft Fracture With McCune-Albright Syndrome With Hyperthyroidism Managed With Oral Alendronate: A Case Report

Kuldeep Bansal, Avijeet Prasad, Sumedha Singh, Ankita Chauhan, Shubham Singla

1. Orthopaedics, University College of Medical Sciences, New Delhi, IND
2. Radiodiagnosis, Institute of Medical Sciences and SUM Hospital, Bhubaneswar, IND
3. Dermatology, University College of Medical Sciences, New Delhi, IND

Corresponding author: Shubham Singla, drshubhamsingla@gmail.com

Abstract
McCune-Albright syndrome (MAS) is a complex endocrinopathy with polyostotic fibrous dysplasia and café-au-lait spots. Pathological femoral shaft fracture along with MAS is not a common occurrence. Bisphosphonates can be used for the management of pathological fractures.

A seven-year-old male child presented with pathological femoral shaft fracture with MAS (café-au-lait macules and hyperthyroidism). The patient was managed conservatively with closed reduction along with hip spica and oral alendronate. Fracture consolidation occurred within six to eight weeks with improvement in bone pains. Oral alendronate can be used as an adjuvant to conservative therapy for pathological fracture in MAS in children.

Introduction
McCune-Albright syndrome (MAS) is a rare mosaic disorder that often manifests as a trio consisting of polyostotic fibrous dysplasia (FD), café-au-lait macules, and varied endocrine abnormalities (especially precocious puberty) [1]. FD can be monostotic or polyostotic; it can occur alone or as a component of MAS [1]. The most prevalent site of FD is the proximal femur, and the resultant fracture rates are the highest in childhood and decrease gradually with age [1]. We present the case of a seven-year-old boy with a pathological femoral shaft fracture, peculiar café-au-lait macules crossing the midline, and hyperthyroidism who was treated conservatively with oral alendronate.

Case Presentation
A seven-year-old boy reported to the emergency department of orthopedics with a sudden onset of pain in the left thigh and an inability to bear weight on the left lower extremity following a minor fall while playing one week prior. Considering the damage to be a mild soft tissue injury, no medical consultation was sought for the first week after the incident. As the child’s symptoms did not improve, he was brought in for evaluation. On general physical examination, the patient was lean and thin built, with three to four hyperpigmented macules on the back, abdomen (Figure 1), and right lower limb (Figure 2). The two lesions on the back were light to dark brown, ranging in size from 15 × 8 cm to 16 × 15 cm, with irregular margins (“Coast of Maine”) and crossing the midline, whereas the lesions on the abdomen and right lower limb were segmental in distribution, did not extend to the left side, and resembled café-au-lait spots.
FIGURE 1: Presence of large hyperpigmented macules with irregular margins over the back and crossing the mid-line.
A plain radiograph of the patient’s left hip-thigh-knee revealed a small oblique fracture line at the junction of the upper and middle thirds of the femur, along with deformed proximal femoral anatomy with a ground-glass appearance. The case was reviewed by a multidisciplinary panel that included an orthopedist, dermatologist, radiologist, and endocrinologist. In accordance with MAS, the patient was assessed with a skeletal survey and an endocrine profile which included serum parathyroid hormone (PTH), thyroid-stimulating hormone (TSH), cortisol, and growth hormone (GH). Serum values of calcium, phosphorus, 25-OH vitamin D, and PTH were normal. Alkaline phosphatase (ALP) and 24-hour urinary hydroxyproline levels were higher, indicating increased bone turnover. The blood count, electrocardiogram, and serum liver
enzymes were all within normal limits. There were no signs of precocious puberty; however, the TSH level in the serum was high. Based on a retrospective evaluation, a history of a wide-based gait with minimum to no activity restriction was obtained. The skeletal survey revealed bilateral short deformed tubular metacarpal bones (Figure 3), as well as a bilateral proximal femoral deformity with a ground-glass appearance (Figure 4), with no scoliosis or craniofacial asymmetry.

![Figure 3: Short deformed tubular metacarpal bones.](image-url)
The objective was to fix the femoral shaft fracture following a discussion with the parents regarding the nature of the disease, but the parents refused. Consequently, the patient was treated conservatively with closed reduction and application of a hip spica, and tablet alendronate (ALN) (70 mg tablet/2 mg per kg with 200 mL of water) was administered once a week; initially, under medical supervision. After taking the drug, he was instructed to maintain fasting and an upright position for at least 30 minutes. After two weeks, the patient continued to take his prescription at home without any problems. The ALN treatment was continued for 12 weeks. At six weeks, a second X-ray revealed an early callus bridging the fracture (Figure 5).
FIGURE 5: Early bridging callus across fracture at six weeks.

At 12 weeks, the fracture was completely consolidated, with full knee and hip range of motion. We detected radiological thickening of the femoral shaft cortex, progressive ossification of the fibrodysplastic lesion, and periosteal bone. The parents were advised on the importance of close monitoring and follow-up for skeletal deformities.

Discussion

MAS is a rare condition characterized by the classical triad of polyostotic FD, skin hyperpigmentation (café-au-lait patches), and endocrine abnormalities, especially precocious puberty [1]. FD is caused by activating mutations in GNAS on chromosome 3, which leads to dysregulated GαS-protein signaling in affected tissues, resulting in the replacement of healthy bone with unhealthy fibro-osseous tissue. MAS is a mosaic disorder with asymptomatic to complete disability [2-4].
For simple fractures of the shaft of the femur in children, various treatment options are available; however, with MAS, the preferred method for pathological fractures of the shaft of the femur is fixation as fracture union may be delayed in FD, and conservative treatment results in proximal femoral deformities such as coxa vara/valgus [5,6].

In this case report, because the patient’s parents refused surgery, we managed him conservatively with hip spica and oral ALN. The patient was referred to the endocrine division for additional examination and care of hyperthyroidism. Several research organizations have published case reports and patient series about the treatment of FD with bisphosphonates (pamidronate, ALN, and zoledronate). There are numerous accounts on the use of pamidronate and zolendronate [7-10]. However, to the best of our knowledge, this is the first report of FD in MAS treated with ALN. Chapurlat et al. [11] also observed that pamidronate therapy (180 mg) alleviated bone pain and reduced bone resorption with improvements in radiological results in less than half of 62 individuals treated. Liens et al. described a group of 30 patients treated with pamidronate (180 mg) at six-month intervals for three cycles, followed by annual administration [12].

Bisphosphonates administered intravenously have a satisfactory safety record overall. Hypocalcemia, as well as flu-like symptoms and brief fever, are among the adverse effects of the drug [10]. In our case, oral bisphosphonates had no side effects.

Conclusions

Oral ALN therapy can be, in the future, considered an alternative treatment for pathological fractures in MAS because it does not alter or delay fracture healing time and reduces bone pains associated with polyostotic FD.

This case represents a child with a rare condition, MAS, with an atypical presentation of skin lesions crossing the midline on the back and being confined to the right side of the midline in the front. To our knowledge, the existence of pathological fracture in the proximal femur (FD) on the opposite side of skin involvement has not been previously recorded. For pathological fractures in MAS, an alternate therapy in the form of oral ALN and hip spica could be considered. More studies with a larger sample size need to be done to establish it as an alternative modality.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Dumitrescu CE, Collins MT: McCune-Albright syndrome. Orphanet J Rare Dis. 2008, 3:12. 10.1186/1750-1172-3-12
2. Weinstein LS, Shenker A, Geiman PV, Merino MJ, Friedman E, Spiegel AM: Activating mutations of the stimulatory G protein in the McCune-Albright syndrome. N Engl J Med. 1991, 325:1688-95. 10.1056/NEJM199112223252403
3. Bianco P, Riminucci M, Majoulaghe A, et al.: Mutations of the GNAS1 gene, stromal cell dysfunction, and osteomalacic changes in non-McCune-Albright fibrous dysplasia of bone. J Bone Miner Res. 2000, 15:120-8. 10.1359/jbmr.2000.15.1.120
4. Hartley I, Zhadina M, Collins MT, Boyce AM: Fibrous dysplasia of bone and McCune-Albright syndrome: a bench to bedside review. Calcif Tissue Int. 2019, 104:517-29. 10.1007/s00223-019-00550-z
5. Javaid MK, Boyce A, Appelman-Dijkstra N, et al.: Best practice management guidelines for fibrous dysplasia/McCune-Albright syndrome: a consensus statement from the FD/MAS international consortium. Orphanet J Rare Dis. 2019, 14:139. 10.1186/s13023-019-1102-9
6. Hefti F, Donnan L, Krieg AH: Treatment of shepherd’s crook deformity in patients with polyostotic fibrous dysplasia using a new type of custom made retrograde intramedullary nail: a technical note. J Child Orthop. 2017, 11:64-70. 10.1007/s00223-016-1207-2
7. Wang Y, Wang O, Jiang Y, Li M, Xia W, Meng X, Xing X: Efficacy and safety of bisphosphonate therapy in McCune-Albright syndrome-related polyostotic fibrous dysplasia: a single-center experience. Endocr Pract. 2019, 25:23-30. 10.4158/EP-2018-0328
8. Aragão AL, Silva IN: Oral alendronate treatment for severe polyostotic fibrous dysplasia due to McCune-Albright syndrome in a child: a case report. Int J Pediatr Endocrinol. 2010, 2010:432060. 10.1155/2010/432060
9. Lala R, Matarazzo P, Andreo M, Marzari D, Bellone J, Corrias A, de Sanctis C: Bisphosphonate treatment of bone fibrous dysplasia in McCune-Albright syndrome. J Pediatr Endocrinol Metab. 2006, 19 Suppl 2:583-93. 10.1515/jpem.2006.19.2.583
10. Majoor BC, Appelman-Dijkstra NM, Fiocco M, van de Sande MA, Dijkstra PS, Hamdy NA: Outcome of long-term bisphosphonate therapy in McCune-Albright syndrome and polyostotic fibrous dysplasia. J Bone Miner Res. 2017, 32:264-76. 10.1002/jbmr.2999

11. Chapurlat RD: Medical therapy in adults with fibrous dysplasia of bone. J Bone Miner Res. 2006, 21 Suppl 2:114-9. 10.1359/jbmr.06s222

12. Liens D, Delmas PD, Meunier PJ: Long-term effects of intravenous pamidronate in fibrous dysplasia of bone. Lancet. 1994, 343:953-4. 10.1016/s0140-6736(94)90369-8

8 of 8