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Intraoperative Identification and Mosaicplasty in a Case of Femur Subchondral Osteoid Osteoma

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Abstract

Osteoid osteomas (OOs) are non-malignant primary bone abnormalities marked by a central nidus surrounded by reactive sclerosis. They typically manifest as aggravated nocturnal pain that responds to non-steroidal anti-inflammatory drugs (NSAIDs). These growths are most frequently found within the intracortical bone and the diaphysis of elongated bones. Within the realm of uncommon conditions, intra-articular OOs (IAOOs) exhibit distinctive presentations, often leading to postponed or inaccurate diagnoses. We present a patient with OO at the distal femur, accessible through the knee joint, which was intraoperatively identified and localized using a *needle pricking technique* and treated by arthrotomy and mosaicplasty.

Categories: Orthopedics, Trauma

Keywords: needle pricking method, subchondral, arthrotomy, mosaicplasty, osteoid osteoma

Introduction

Osteoid osteoma (OO) is a non-malignant, osteoblastic, and painful tumor-like lesion. It ranks as the third most prevalent benign bone lesion, with intra-articular cases comprising around 5% to 13% of occurrences [1-3]. Within the category of intra-articular OOs (IAOOs), sometimes referred to as "juxta-articular OOs," these are characterized by their development within synovial cavities, specifically in subsynovial or subchondral locations. Subchondral IAOOs stand out due to their proximity to and engagement with articular cartilage [2]. Unlike their extra-articular counterparts located in the diaphysis, IAOOs often exhibit atypical radiographic and advanced imaging manifestations, contributing to delays in diagnosis, taking over two years on average, in contrast to 8.5 months for extra-articular cases [1,2,4,5].

Upon recognition, intervention is generally required for IAOOs. Minimally invasive approaches aided by imaging, such as CT-guided radiofrequency ablation (CT-RFA), present potential strategies. However, these methods carry theoretical risks of affecting cartilage and lack comprehensive exploration for subchondral IAOO treatment [6,7]. Surgical extraction can be done through open or arthroscopic procedures, yet preserving articular cartilage proves challenging, particularly in weight-bearing regions [3,8,9].

Presented here is a case of an IAOO situated in the distal femur at a subchondral location. This case was identified and localized intraoperatively using a needle pricking technique and treated with arthrotomy and mosaicplasty.

Case Presentation

Clinical history

A 36-year-old male reported to the OPD and complained of a 4-year history of atraumatic pain in the left knee which was undiagnosed on three previous MRIs.

The pain emerged gradually, with a subtle onset and mild intensity at first, but over the course of several months, it steadily worsened. This persistent discomfort was present throughout both daytime and nighttime, and while its severity was somewhat relieved by non-steroidal anti-inflammatory drugs (NSAIDs), it persisted despite treatment.

Upon physical examination, there was noticeable but mild swelling and tenderness at the medial femoral condyle. No joint effusion or quadriceps atrophy was observed. The range of motion of the knee extended from 0 to 110 degrees, showing symmetrical symmetry with the opposite knee. Evaluation of the patellofemoral joint revealed no abnormalities.

A knee X-ray was conducted, yielding unremarkable results (Figure 1). Subsequently, a CT scan (Figure 2) and an MRI were performed (Figure 3). These imaging techniques revealed a distinct, well-defined circular lesion measuring 8x6 mm located in the front part of the medial femoral condyle. Furthermore, the MRI indicated areas of uneven bone marrow swelling within the medial femoral condyle, along with a noticeable

 $accumulation \ of \ fluid \ in \ the \ knee \ joint. \ The \ differential \ diagnosis \ encompassed \ subchondral \ cysts \ and \ OOs.$

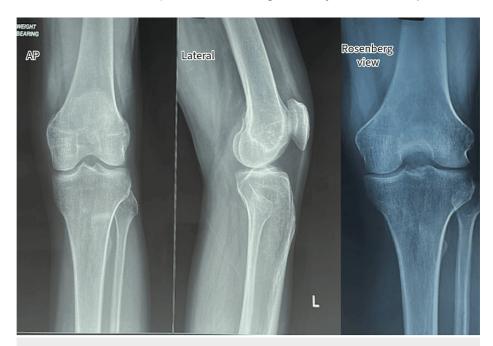


FIGURE 1: Preoperative plain radiograph



FIGURE 2: CT scan showing left knee joint

Arrows showing the lesion

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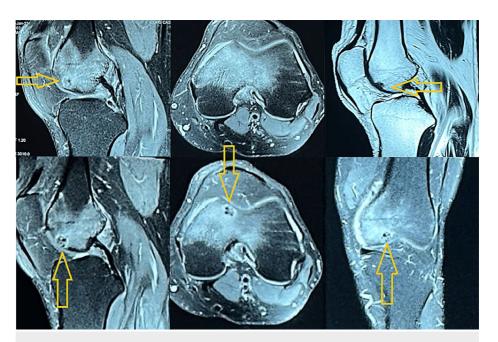


FIGURE 3: MRI showing the left knee joint

Arrows showing the lesion

We had planned an open arthrotomy of the knee joint and excision of the tumor with reconstruction by mosaicplasty. The individual was positioned in a supine stance with spinal anesthesia administered. A medial parapatellar incision was made, and the patella was flipped laterally, revealing the lesion located over the distal medial femoral condyle. The cartilage over the trochlea and medial femoral condyle was normal. On C-Arm, the lesion was not identified. The preoperative CT scans had pinpointed a lesion measuring between 6 and 8 mm in diameter and around 1 mm below the subchondral bone. As a very thin bone wall was present between the normal cartilage and the tumor, we decided to identify the tumor by our needle pricking technique with an 18G needle for precise intraoperative identification, localization, and depth assessment (as shown in Figure 4A). We measured the preoperative distances of the tumor from three bony edges, and intraoperatively, we marked the same measurements with a marker pen (as shown in Figure 4A). After identifying the small area between the marked measurements, an 18G needle was used to prick the normal cartilage. At the site of the tumor, the needle was pierced deep around 1 cm, and repeating the same, we localized the whole tumor. The cartilage over the tumor was removed, and curettage of the tumor was done. After curettage, we had a defect of 14x8 mm. Utilizing a mosaicplasty kit (8 mm Arthrex), we prepared two recipient wells at the site of the excised tumor. Two osteochondral grafts of size 8 mm were harvested from the non-weight-bearing part of the lateral femoral condyle and put in the prepared recipient site. With mosaicplasty, we could reconstruct the cartilage defect (as depicted in Figure 4B-4C).

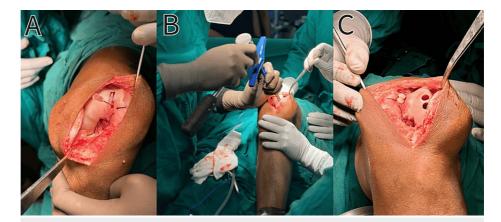


FIGURE 4: Intraoperative pictures

(A) Intraoperative identification of the lesion using a needle pricking technique. (B) Lesion taken out using a harvester. (C) Graft harvested from the lateral femoral condyle and inserted into the medial condyle

Intraoperative samples were sent for histopathological examination which confirmed the diagnosis to be OO (Figure 5), and a postoperative X-ray the of left knee joint was done (Figure 6).

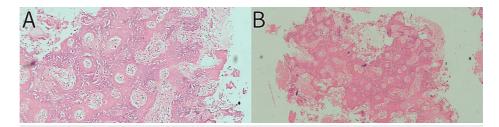


FIGURE 5: Histopathological slides of the intraoperative sample confirmed the diagnosis to be OO

(A) Magnification: 400X. (B) Magnification: 100X. Stain - hematoxylin and eosin

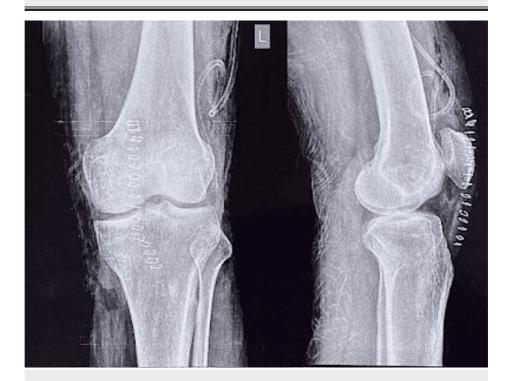


FIGURE 6: Postoperative plain radiograph of the left knee joint

Postoperative rehabilitation

The patient was kept partial weight bearing with a long knee brace for six weeks. The knee range of motion was restricted up to 60 degrees for three weeks and later improved to complete flexion by the end of eight weeks, and full weight bearing was allowed at the end of eight weeks. Further quadriceps strengthening and sports activities were started at around six months. The patient was comfortable with a full range of motion and joined his job in the security forces at the end of eight months.

Discussion

IAOO is commonly observed in the hip joint, with a primary prevalence in the upper part of the femur. Nonetheless, occurrences of IAOO have been noted in different joints such as the knee, ankle, elbow, hand, subtalar, and metatarsophalangeal joints [1,2,10]. Regular X-rays often fail to promptly detect IAOO since the distinctive hardened border is usually absent in most cases [2,5].

The plan can be to make a patient-specific 3D printed jig to localize the tumor unavailable at our institute. However, MRI plays a vital role in distinguishing osteochondral defects, subchondral cysts, infections, or stress fractures, but it does not display IAOO early as profound marrow edema, effusion, and no sclerosis around the lesion in the early stage, making it difficult to differentiate with infection [6,11-13]. CT scans are the favored diagnostic imaging technique, especially when using 3 mm thick sections or smaller to precisely

identify small niduses [6,12].

While NSAIDs are the mainstay for OO treatment, addressing IAOOs is more complex due to its connection with synovitis-associated joint complications, resulting in heightened morbidity [4,14]. CT-RFA is the recognized interventional treatment for classic, extra-articular OOs, boasting success rates exceeding 90% [15]. However, applying this technique to IAOOs raises concerns about potential chondrocyte demise attributed to thermal injury. Previous studies involving cadavers and animals have indicated a 1 cm area of damage resulting from CT-RFA [16-18]. Utilizing radiofrequency ablation poses a potential risk of cartilage damage above the lesion and possibly insufficiently addressing pathologically compromised cartilage. Conversely, while straightforward excision may offer pain relief, it fails to adequately address the subsequent loss of articular cartilage. Nevertheless, various studies indicate the use of CT-RFA as a treatment for IAOO, demonstrating results similar to those seen in cases of diaphyseal OO [7,15,19,20]. It's worth noting that the ablation procedure typically avoids a transarticular approach, especially in hip IAOO cases, with only a limited number of instances involving direct subchondral sites. To address this uncommon scenario, we propose an alternative treatment approach, involving the needle pricking technique described in the procedure with arthrotomy followed by mosaicplasty. However, it can be done only if the tumor wall is 1-2 mm thin and cannot be performed in deeper IAOOs. This innovative method has demonstrated positive outcomes, including effective pain alleviation and the restoration of the articular surface.

The primary surgical intervention for IAOOs has commonly involved straightforward excision, employing both open and arthroscopic methods [3,8,9,11]. Various accounts of distal femoral lesions underscore favorable pain alleviation and functional outcomes achieved through excision. However, it's important to note that these accounts exclusively pertain to subsynovial lesions that do not induce damage to the weightbearing zone of the cartilage [3,8,9]. Hence, in specific patient cases, osteochondral grafting presents various benefits. This approach addresses a range of potential diagnoses, including osteochondral defects, allowing appropriate treatment for either condition. After complete excision, a definitive diagnosis is more feasible due to the sufficient biopsy sample, as confirmed pathology results were only achieved in 73% of a significant series of OOs [15]. Osteochondral grafting represents a well-established approach for addressing cartilage defects in the distal femur, thus presenting a rational therapeutic choice for this specific site. While the patient in our case had reached skeletal maturity, it is conceivable that this technique could be adapted for pediatric patients with adequate epiphyseal size, allowing for resection without influencing the growth plate.

Conclusions

We introduce a case involving a subchondral IAOO located within the distal femur. In such cases, intraoperative identification of the tumor is difficult because of the normal overlying cartilage. For the localization of the tumor, we used our needle pricking technique with preoperative CT planning, which is easy and cheaper than making a 3D-printed patient-specific jig. We advocate using this technique in tumors with a very thin wall of subchondral bone between the tumor and cartilage. This ingenious method exemplified positive outcomes, effective pain alleviation, and the restoration of the articular surface.

Additional Information

Author Contributions

All authors have reviewed the final version to be published and agreed to be accountable for all aspects of the work.

Concept and design: Sushant Kumar, Rishabh Aggarwal, Vinod Nair, Vishal S. Patil, Archit Gupta

Acquisition, analysis, or interpretation of data: Sushant Kumar, Archit Gupta

 $\textbf{Drafting of the manuscript:} \ \textbf{Sushant Kumar, Rishabh Aggarwal, Archit Gupta}$

Critical review of the manuscript for important intellectual content: Sushant Kumar, Vinod Nair, Vishal S. Patil, Archit Gupta

Supervision: Vinod Nair, Vishal S. Patil

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.

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References

- Rolvien T, Zustin J, Mussawy H, Schmidt T, Pogoda P, Ueblacker P: Intra-articular osteoid osteoma as a differential diagnosis of diffuse mono-articular joint pain. BMC Musculoskelet Disord. 2016, 17:455. 10.1186/s12891-016-1313-3
- Szendroi M, Köllo K, Antal I, Lakatos J, Szoke G: Intraarticular osteoid osteoma: clinical features, imaging results, and comparison with extraarticular localization. J Rheumatol. 2004, 31:957-64.
- Abnousi F, Saliman JD, Fanton GS: Arthroscopic visualization and assisted excision of osteoid osteoma at the knee: a case report and review. Am J Sports Med. 2008, 36:375-8. 10.1177/0363546507307865
- Cassar-Pullicino VN, McCall IW, Wan S: Intra-articular osteoid osteoma. Clin Radiol. 1992, 45:153-60. 10.1016/s0009-9260(05)80631-2
- Bauer TW, Zehr RJ, Belhobek GH, Marks KE: Juxta-articular osteoid osteoma. Am J Surg Pathol. 1991, 15:381-7
- Pikoulas C, Mantzikopoulos G, Thanos L, Passomenos D, Dalamarinis C, Glampedaki-Dagianta K: Unusually located osteoid osteomas. Eur J Radiol. 1995, 20:120-5. 10.1016/0720-048X(95)00636-5
- Mylona S, Patsoura S, Galani P, Karapostolakis G, Pomoni A, Thanos L: Osteoid osteomas in common and in technically challenging locations treated with computed tomography-guided percutaneous radiofrequency ablation. Skeletal Radiol. 2010, 39:443-9. 10.1007/s00256-009-0859-7
- 8. Gunes T, Erdem M, Bostan B, Sen C, Sahin SA: Arthroscopic excision of the osteoid osteoma at the distal femur. Knee Surg Sports Traumatol Arthrosc. 2008, 16:90-3. 10.1007/s00167-007-0380-2
- Kang S, Kim YS, Lee HM, Lee MY, Kim JP: Arthroscopic Excision of an intraarticular osteoid osteoma in the Distal femur. Clin Orthop Surg. 2016, 8:475-80. 10.4055/cios.2016.8.4.475
- Corbett JM, Wilde AH, McCormack LJ, Evarts CM: Intra-articular osteoid osteoma: a diagnostic problem. Clin Orthop Relat Res. 1974, 98:225-30.
- Harun M, Hayrettin Y, Serhat M, Engin C, Kamil C, Armagan A, Sancar PA: Atypical location of an osteoid osteoma with atypical anterior knee pain. Int J Surg Case Rep. 2014, 5:873-6. 10.1016/j.ijscr.2014.10.015
- Iyer RS, Chapman T, Chew FS: Pediatric bone imaging: diagnostic imaging of osteoid osteoma . AJR Am J Roentgenol. 2012, 198:1039-52. 10.2214/AJR.10.7313
- Leeman JJ, Motamedi D, Wildman-Tobriner B, O'Donnell RJ, Link TM: Intra-articular osteoid osteoma at the femoral trochlea treated with osteochondral autograft transplantation. J Radiol Case Rep. 2016, 10:22-9. 10.3941/jrcr.v10i6.2817
- 14. Kneisl JS, Simon MA: Medical management compared with operative treatment for osteoid-osteoma . J Bone Joint Surg Am. 1992, 74:179-85.
- Rosenthal DI, Hornicek FJ, Torriani M, Gebhardt MC, Mankin HJ: Osteoid osteoma: percutaneous treatment with radiofrequency energy. Radiology. 2003, 229:171-5. 10.1148/radiol.2291021053
- Caffey S, McPherson E, Moore B, Hedman T, Vangsness CT Jr: Effects of radiofrequency energy on human articular cartilage: an analysis of 5 systems. Am J Sports Med. 2005, 33:1035-9. 10.1177/0363546504271965
- Edwards RB 3rd, Lu Y, Uthamanthil RK, Bogdanske JJ, Muir P, Athanasiou KA, Markel MD: Comparison of mechanical debridement and radiofrequency energy for chondroplasty in an in vivo equine model of partial thickness cartilage injury. Osteoarthritis Cartilage. 2007, 15:169-78. 10.1016/j.joca.2006.06.021
- Rosenthal D, Callstrom MR: Critical review and state of the art in interventional oncology: benign and metastatic disease involving bone. Radiology. 2012, 262:765-80. 10.1148/radiol.11101384
- Filippiadis DK, Velonakis G, Kostantos C, Kouloulias V, Brountzos E, Kelekis N, Kelekis A: Computed tomography-guided radiofrequency ablation of intra-articular osteoid osteoma: a single centre's experience. Int J Hyperthermia. 2017, 33:670-4. 10.1080/02656736.2017.1294711
- Papagelopoulos PJ, Mavrogenis AF, Kyriakopoulos CK, Benetos IS, Kelekis NL, Andreou J, Soucacos PN: Radiofrequency ablation of intra-articular osteoid osteoma of the hip . J Int Med Res. 2006, 34:537-44. 10.1177/147323000603400512