Partial excision of the nidus of an atypical cancellous osteoid osteoma by use of a bone marrow biopsy needle under fluoroscopic guidance

Atipik kansellöz osteoid osteomada floroskopi kılavuzluğunda kemik iliği biyopsi iğnesi ile nidusun kısmi eksizyonu

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Received: 28 November 2018 Accepted: 18 May 2019

ABSTRACT

Aim: The widely accepted method in treatment of osteoid osteoma is either the complete excision or destruction of the nidus. The aim of this study was to evaluate the efficacy of partial nidus excision using a bone marrow biopsy needle as a minimally invasive technique for management of atypical cancellous osteoid osteomas.

Material and Method: Partial excision of the nidus was performed in four cases using an 11-G bone marrow biopsy needle under fluoroscopic guidance. The lesions were located in the capital femoral epiphysis, the posterosuperior side of the femoral neck, the distal tibial epiphysis and the olecranon process of the ulna.

Results: The patient’s pain resolved the night following excision of the nidus. No recurrence was observed at the 72, 36, 32 and 24 month follow-ups.

Conclusion: Partial excision of the nidus may be considered a remarkable technique especially in the treatment of atypical cancellous osteoid osteoma, where surgical intervention is challenging.

Keywords: osteoid osteoma, partial excision, fluoroscopy, biopsy needle
ÖZ

Amaç: Osteoid osteoma tedavisinde yaygın olarak kabul edilen yöntem nidusun ya tam eksizyonu ya da tahrib edilmesidir. Bu çalışmanın amacı, atipik kansellöz osteoid osteoma tedavisinde minimal invaziv teknik olarak kemik iliği biyopsi iğnesi ile kısmi nidus eksizyonu etkinliğini değerlendirmekti.

Gereç ve Yöntem: Dört olguda 11-G kemik iliği biyopsi iğnesiyle floroskopi kılavuzluğunda nidusun kısmi eksizyonu uygulandı. Lesyonlar femur başı epifizi, femur boynu posterosüperior tarafı, tibia distal epifizi ve olekranon çıkıntida yerlesmişlerdi.

Bulgular: Hastaların ağrısi nidus eksizyonunu takip eden gece geçti. 72, 36, 32 ve 24. aylardaki takipte nüks görülmedi.

Sonuç: Nidusun kısmi eksizyonu özellikle cerrahi girişimin zor olduğu atipik kansellöz osteoid osteoma tedavisinde dikkate değer bir teknik olarak düşünülebilir.

Anahtar kelimeler: osteoid osteoma, kısmi eksizyon, floroskopi, biyopsi iğnesi

INTRODUCTION

The Osteoid Osteoma (OO) is the third most common benign bone tumor, and is commonly seen in the second and third decades of life. The OO lesion was first proposed as a distinct entity by Jaffe in 1935 [1]. More than half of these lesions occur in the long bones of the lower extremities. The proximal femur, mainly in the neck portion, is the most frequent location for the development of an OO [2-4]. OO lesions can occur anywhere within the bone, including the cortex, medulla, subperiosteal and intracapsular areas. OO lesions are rarely seen in the epiphysis of the femur, the carpals, the phalanges, the bones of the pelvis and the vertebrae [5-10].

It has been accepted that a complete recovery from an OO is possible through the removal or destruction of the nidus. A residual nidus has been reported to be a source of recurrence [11]. In our study, we evaluated the efficacy of interventional partial nidus excision technique using a bone marrow biopsy needle under fluoroscopic guidance in patients with atypical cancellous bone OOs.

MATERIAL AND METHOD

Four patients with atypical cases of OO underwent partial nidus excision under fluoroscopic guidance by an 11G bone marrow biopsy needle (11G Jamshidi bone marrow biopsy needle, CareFusion, San Diego, CA) at the Orthopedics Department of our training and research hospital between 2010 and 2016. Approval of the local ethics committee was obtained. Informed consent forms were signed by the patients or their parents. All patients presented to our clinic complaining of pain on their involved extremity. The patients’ pain increased at night, and was relieved with salicylates or non-steroidal anti-inflammatory drugs. The average onset of pain was 5.8 months (range 2 to 14) before the intervention. Radiographic examinations were performed by obtaining AP and lateral X-ray views of the painful bones, which along with computed tomography (CT). In all cases we performed a partial excision of the nidus of the same lesion twice sequentially in the same session.

Case 1

Our first atypical OO was in the left capital femoral epiphysis near the physeal line of a 10-year-old boy. The patient had been treated for Legg-Calve Perthes disease for a year at another hospital. The lesion had a sclerotic zone of approximately 2 cm with a partially discernible lucent focus on plain radiographs (Figure 1a). CT scans showed a well-delineated sclerotic area 17 mm in size with a central lytic focus of 5 mm (Figure 1b). We performed an interventional procedure to excise a 2 mm portion of the nidus using an 11G Jamshidi biopsy needle under fluoroscopic guidance (Figure 1c). Histopathologic diagnosis confirmed that the lesion was an OO. CT scans at the one-year follow-up revealed that the nidus was calcified, and surrounded by a dense peripheral sclerosis (Figure 1d). Plain radiographs 72 months after intervention showed a complete resolution of the sclerosis, and the central lucency was not visible (Figure 1e). A repeat CT also demonstrated a calcified nidus with a diminished sclerotic area and excessive ossification of the femoral head (Figure 1f).
Figure 1. a) The OO of the capital femoral epiphysis with a sclerotic area together with a fairly delineated, centrally lucent focus b) CT scan of the OO of the capital femoral epiphysis, revealing a well-delineated sclerotic area with a central lytic focus c) Partial excision of the nidus using an 11G Jamshidi biopsy needle under fluoroscopy d) A CT image of the OO of the capital femoral epiphysis at the 1-year follow-up. The nidus began to calcify and dense sclerosis is evident e) An anteroposterior plain radiograph of the capital femoral epiphysis 72 months later. No significant findings were identified. The lucent focus has completely disappeared f) A CT image of the capital femoral epiphysis 72 months later. The nidus was completely calcified and the sclerotic rim had regressed in size and was mostly ossified
Case 2

Our second atypical OO case was a 12-year-old boy with a 4 mm radiolucent lesion of the left distal epiphysis of the tibia, with cortical thickening on the medial side (Figure 2a). The lesion was located on the margin of the distal tibia and eccentric sclerosis was detected on CT imaging (Figure 2b). An excision was performed by placing an 11G Jamshidi biopsy needle into the centrally located nidus under fluoroscopy. Histopathologic analysis confirmed the diagnosis. Follow-up plain radiographs showed a complete resolution of the OO 32 months later (Figure 2c), and no lesion was detected on CT.

Figure 2. a) The atypical OO of the left distal tibial epiphysis with a marginal lytic lesion and cortical thickening b) A coronal CT scan of the left distal tibia, revealing a marginal cortical bone defect with an eccentric dense bone reaction c) An anteroposterior plain radiograph of the left distal tibia showing the complete resolution of the OO 32 months later

Case 3

The third patient with an atypical OO was a 18-year-old boy with a lesion located on the right proximal femur in the posterosuperior portion of the femoral neck. Plain pelvic radiographs revealed a mildly radiolucent lesion on the posterosuperior aspect of the femoral neck. A lytic lesion with peripheral sclerosis was seen on CT scan (Figure 3a). The size of the nidus was measured as 4 mm. The same partial excision procedure as in the first two cases was performed. Histopathologic diagnosis confirmed an OO. Plain radiographs taken at the 3-year follow-up were normal. The lucent area appeared calcified on CT (Figure 3b).
Case 4

The fourth case was a 21-year-old male patient with an atypical OO on the left olecranon process of the ulna. Plain radiographs showed a lucent area with a sclerotic rim inside the olecranon process, with an irregularity of the posterior cortex (Figure 4a). A CT scan of the lesion confirmed a well-delineated lytic area that was 3 mm in size, located lateral to the olecranon process. The lesion had a calcified focus and a mildly sclerotic rim (Figure 4b). The patient had followed-up at another center for five months due to complaints of pain at the elbow, but had not been diagnosed with an OO during that period. The same treatment protocol was used on this patient. Histopathologic diagnosis confirmed the lesion as OO. Follow-up plain radiographs of the elbow showed a complete disappearance of the nidus 2 years later (Figure 4c).

RESULTS

In our patients, the average size of the OO nidus was 4 mm (range 3 to 5 mm). The patient’s pain resolved the night following excision of the nidus. No recurrence was observed at the 72, 36, 32 and 24 month follow-ups.

DISCUSSION

Classical invasive surgical methods to treat an OO include en bloc resection and burr-down excision techniques. However, percutaneous radiofrequency ablation and percutaneous laser coagulation have become popular as minimally invasive techniques [2,9,10,12]. Other minimally invasive techniques include CT-guided excision [13,14] and excision with arthroscopy [15]. Another method that was recently reported is the MRI-guided focused ultrasound ablation technique [16]. However, this ablation method is not yet widely used. Resolution of pain is a good indicator of the success of surgery. The long-term resolution of pain following the partial excision of the OO was previously reported [17]. Some authors reported on clinically and radiographically diagnosed OO cases that spontaneously regressed within a matter of years [11,18,19]. It has been also been reported that the nidus first calcifies, then ossifies with a concurrent resolution of symptoms over time [20,21]. If the OO symptoms are mild and relieved by non-steroidal anti-inflammatory drugs, clinical observation with follow-up is therefore the preferred approach.

An epiphyseal OO is extremely rare and has been reported in only a few cases in the literature [5,8,9,22,23]. The diagnosis of an epiphyseal OO is challenging and may present with similar radiographic findings to those of a chondroblastoma and a Brodie abscess [1]. A CT scan with contrast is preferable to differentiate these lesions from an OO. Legg Calve Perthes disease also has to be considered in cases without typical imaging findings [12]. Epiphyseal OOs may occur in or near intra-articular sites and may occasionally present with clinical findings ranging from joint effusion to decrease in range of movement. No similar findings were observed during the follow-up of our patients with an OO in the capital femoral epiphysis and the posterosuperior aspect of the femoral neck. An OO located in the tibial distal epiphysis [24] and the olecranon process of the ulna [25] is a rare pathology, whose diagnosis is often delayed due to insignificant symptoms and minimal radiographic findings during the onset of the disease.
The limited number of patients and enrolling only patients with atypically located cancellous osteoid osteoma constitute limitations of our study. We do not assert that the technique we propose, performed on a limited number of patients, will be an immediate alternative to the current successful methods [2,9] exhibited in the literature. However, taking the positive outcomes achieved in all patients following partial excision of the nidus into account, we concluded that the technique may be applied on atypical cancellous osteoid osteoma cases, surgically challenging and open to complications as in the osteoid osteoma of the femoral epiphysis in pediatric patients. Our results on the posterosuperior femoral neck, distal epiphysis of the tibia and olecranon process of the ulna, where a surgical intervention is relatively easier, is encouraging for performing partial excision of the nidus. The technique we utilized in the treatment of OOs is different from its predecessors, which advocate complete excision or destruction of the nidus. The technique also differs from the burr-down excision technique. Complete excision of a nidus with a diameter of 3 to 5 mm is geometrically impossible by two sequential punches of the 11G bone marrow biopsy needle. We therefore preferred a partial excision of the nidus. By performing the partial excision technique twice in the nidus of the same lesion, we aimed to increase our chances of excising a significant amount of material from the nidus.

In conclusion, partial excision of the nidus using a bone marrow biopsy needle is an efficient and easily applicable technique. With the minimally invasive technique we propose, partial excision of the nidus may be considered a remarkable alternative in the treatment of atypical cancellous osteoid osteoma where surgical intervention is challenging and complications may be encountered. Further studies enrolling large population of patients are warranted for validation of our technique.

DECLARATION OF CONFLICT OF INTEREST

The authors received no financial support for the research and/or authorship of this article. There is no conflict of interest.

Figure 4. a) The lucent area with a mildly sclerotic rim of the olecranon process and an irregularity of the posterior cortex b) A CT image showing a well-delineated lytic area with a calcified focus on the radial side of the olecranon process c) A lateral plain radiograph of the elbow 2 years later. The nidus has completely disappeared. Soft tissue calcification on the posterior aspect of the elbow.
REFERENCES


