

Bipolar fresh osteochondral allograft of the shoulder

SANDRO GIANNINI, ELISA SEBASTIANI, ALBA SHEHU, MATTEO BALDASSARRI
SUSANNA MARALDI, FRANCESCA VANNINI

1st Clinic of Orthopaedics and Traumatology, Rizzoli Orthopaedic Institute, Bologna, Italy

Abstract

Purpose: to describe an original technique and preliminary results of bipolar fresh osteochondral allograft implantation for the treatment of end-stage glenohumeral osteoarthritis.

Methods: three patients underwent bipolar fresh osteochondral allograft implantation to the shoulder. Clinical and radiographical evaluations were carried out periodically through to final follow-up.

Results: constant Score increased from 38.3 ± 2.9 preoperatively to 78.7 ± 16.2 at 12 months, 72.3 ± 15.3 at 24 months, and 59.3 ± 22.0 at 34 months. Arthritis and partial reabsorption of the implanted surfaces were evident radiographically.

Conclusions: the clinical results obtained in these patients seem to support the applicability of bipolar fresh osteochondral allograft implantation in the shoulder in subjects with severe post-traumatic arthritis and intact rotator cuff. The development of arthritis of the implanted surfaces, while not impacting the clinical result, is a cause of concern.

Level of evidence: level IV, therapeutic case series.

Key Words: allograft, arthritis, bipolar fresh osteochondral allograft, shoulder.

Introduction

Shoulder pain due to osteoarthritis of the glenohumeral joint is a therapeutic challenge for the surgeon (1, 2). In patients with an intact rotator cuff, the most

widely used surgical options are total shoulder arthroplasty and hemiarthroplasty with or without biological glenoid resurfacing, even in patients younger than 55 years (3, 4).

Nevertheless, shoulder arthroplasty in young patients carries a high percentage of unsatisfactory results, even though recent reports have shown favorable outcomes in the short to medium term (5-7).

Bipolar fresh osteochondral allograft (BFOA) placement, first proposed for the ankle and knee joint, is a fascinating option for biological joint reconstruction (8-14). The rationale of BFOA is to provide viable cartilage that can survive transplantation, being supported by an intact subchondral bony structure, which is progressively integrated and replaced by host bone over time (12, 15, 16).

To our knowledge, the literature contains no reports of bipolar total allograft procedures applied to the shoulder (12, 17).

The purpose of the present study was to describe the surgical technique for BFOA of the shoulder and to report the outcome of BFOA of the shoulder in three cases at more than 30 months' follow-up.

Methods

Three patients (two men and one woman), mean age 44.0 ± 3.6 years, were referred for evaluation of long-standing shoulder pain and mechanical symptoms at the authors' institute. The diagnosis of post-traumatic arthritis of the glenohumeral surfaces with intact rotator cuff was confirmed by MRI. Substitution of the bare articular surfaces with a fresh bipolar allograft was indicated as a surgical option. The proposed procedure was approved by the institute's ethics committee and informed consent to the procedure was obtained from the patients.

Corresponding Author:

Francesca Vannini MD, PhD
1st Clinic of Orthopaedics and Traumatology,
Rizzoli Orthopaedic Institute, Bologna, Italy
Phone: +39 051/6366669
E-mail: France_vannini@yahoo.it

The affected shoulder of each patient was examined in order to assess the level of pain and the functional impairment. Clinical evaluation before allograft reconstruction was performed and rated according to the Constant Score. Radiographic evaluation was performed with standard radiographs and a CT scan of the affected shoulder was also performed in order to digitally measure the size of the shoulder. The supero-inferior and antero-posterior diameters of the glenoid were taken into account, together with the diameter of the humeral head considered as a sphere, and used to match the donor size with the recipient site.

The candidates were put on a waiting list pending the availability of an appropriate-sized donor joint. The donors were identified through our institute's Bone Bank program for musculoskeletal tissue transplantation. Using a standard surgical procedure, in a sterile regimen, the entire shoulder joint was harvested "en bloc" from each donor. Once harvested, a mini-incision was performed in the capsule in order to visually check the status of the cartilage and to allow diffusion of the storage medium. Following a CT scan, the allograft was placed in a sterile container with L-glutamine, NaHCO₃ and antibiotic solution and stored at 4°C. A MRI was finally performed on the harvested shoulder in order to verify articular cartilage integrity, while the adequacy of the shoulder size for the patient was verified through a CT scan. Allograft implantation was performed 14.0 ± 1.0 days after harvesting. The surgical session consisted of two steps: graft preparation and graft implantation.

Step 1. On a separate surgical table, all soft tissues were carefully removed from the harvested shoulder, taking care not to damage the cartilage. Then the articular surfaces of the glenoid and the humeral head were carefully cut with a standard pneumatic saw, keeping the whole articular surface and about 10-12 mm of subchondral bone intact. The prepared articular surfaces were then temporarily placed in a container of saline solution.

Step 2. The patient was placed in the beach-chair position under general anesthesia. A standard delto-pectoral approach was performed. With a blunt dissection the cephalic vein was identified and dislocated laterally. The subscapularis muscle was then identified and detached from its insertion. The capsule was opened and the humeral head was dislocated

anteriorly. Under fluoroscopic control two Kirschner wires were placed in the humeral neck (perpendicular to the axis of the neck) and another two Kirschner wires were placed in the base of the glenoid, parallel to the articular surface, in order to define the plane for the bone cut (Fig. 1). The cut was then made using a standard pneumatic saw. The articular surfaces of the humeral head and the glenoid were removed taking care to remove the whole articular surfaces and 1 cm of subchondral bone (Fig. 2). Finally, the articular surfaces were replaced with the previously prepared allograft components.

The newly implanted glenoid was fixed with two small fragment screws, whereas the humeral head was fixed with two twist-off screws (Fig. 3). The joint was reduced and fluoroscopic control was performed in order to verify the correct placement of the graft; the shoulder range of motion (ROM) was also checked. Finally, the subscapularis muscle was reinserted, a drainage tube was placed and a routine suture was performed. Postoperative radiographs were taken.

Each patient underwent a light postoperative immunosuppressive therapy regimen: cyclosporin 3 mg/kg per day for six months and prednisone 10 mg per day for the first month then 5 mg per day for a further two months. Nine to 12 months after surgery the patients underwent a second operation for removal of the hardware and biopsy evaluation of the implanted allograft. Biopsy samples of cartilage, bone and synovial tissue were taken and immunological and immunohistochemical analysis of the specimens was carried out. The Constant Score at 12, 24 and 34 months after surgery was used to assess the clinical outcome.

Results

No intraoperative or postoperative complications occurred.

At final follow-up all the patients had a full and painless ROM with no additional complaints of resting pain or pain related to activities; the Constant Score increased from 38.3 ± 2.9 preoperatively to 78.7 ± 16.2 at 12 months, 72.3 ± 15.3 at 24 months and 59.3 ± 22.0 at 34 months. Two patients declared themselves to be satisfied with the treatment and the result. One patient was not satisfied with the treatment



Fig. 1. Illustration showing the line of the cut on the recipient surfaces.

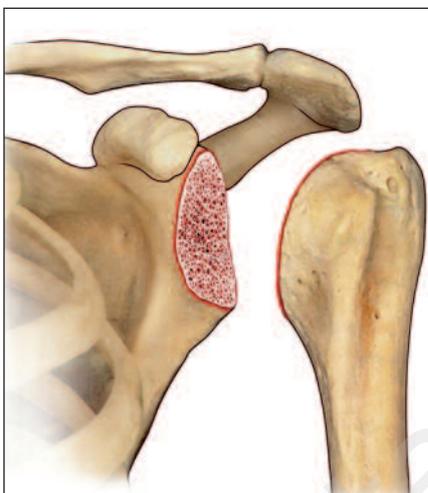


Fig. 2. The articular surfaces of the humeral head and the glenoid were removed taking care to remove the whole articular surfaces and 1 cm of subchondral bone.

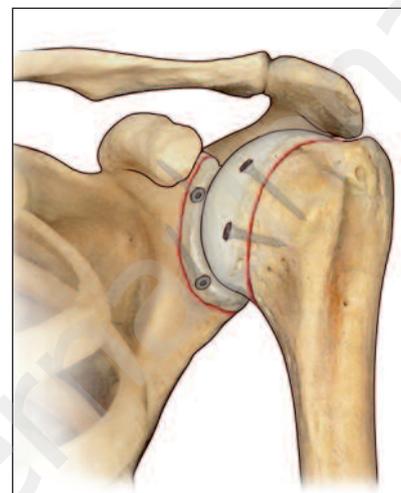


Fig. 3. Illustration showing the correct position of the allograft. The newly implanted glenoid was fixed with two small fragment screws, whereas the humeral head was fixed with two twist-off screws.

because of instability and reduced strength, but to date has refused revision surgery.

All the allografts showed signs of integration on CT scan at four months of follow-up and were considered healed at six months on the basis of X-ray results. Follow-up MRI at six months showed a well-maintained cartilage layer in all cases. There was no intra-articular effusion, although subchondral bone edema was evident. No partial reabsorption of the implanted surfaces was evident at the final follow-up.

The degree of osteoarthritis was not found to impact on the clinical score at any follow-up, nor was it influenced by the immunosuppressive therapy.

Immunohistochemical analysis of the biopsy specimens revealed vital cartilage with a high proteoglycan content, normal cell distribution and type 2 collagen expression. Moreover, on the bony interface, the biopsy showed complete osseointegration between the graft and the recipient bone (Fig. 4). Finally, synovial biopsy analysis revealed a limited number of macrophages, without prominent perivascular inflammatory cell infiltrates or lymphoid aggregates.

Discussion

Bipolar fresh osteochondral allograft was found to be a viable option for the treatment of severe post-trau-

matic arthritis of the shoulder in the presence of an intact rotator cuff. The technique allowed biological resurfacing and may represent a fascinating alternative to traditional prosthetic substitution.

It has been established that BFOA is applicable for treatment of the ankle joint, however, only partial or monopolar allografts have, to date, been reported in other joints (11, 14, 17-20). Although the concept of allograft implantation is well established in the treatment of shoulder instability (21) and in revision shoulder arthroplasty (15), there are no reports of bipolar total fresh allografting.

The arthritic shoulder may represent an ideal indication for BFOA implantation since it is a non-weight-bearing joint, but is subject to compressive and shear forces that can easily lead to mobilization of prosthetic components; the technique was previously presented in a single case report (16). The recipient showed complete integration of the allograft, which allowed him to achieve almost normal shoulder biomechanics and therefore an almost normal shoulder ROM (16).

These results were confirmed in the present case series, even though the number of patients considered is very small: shoulder ROM was restored and the patients were pain free. Unfortunately, one case developed laxity of the joint, which was the cause of the significant decrease in this patient's final score.

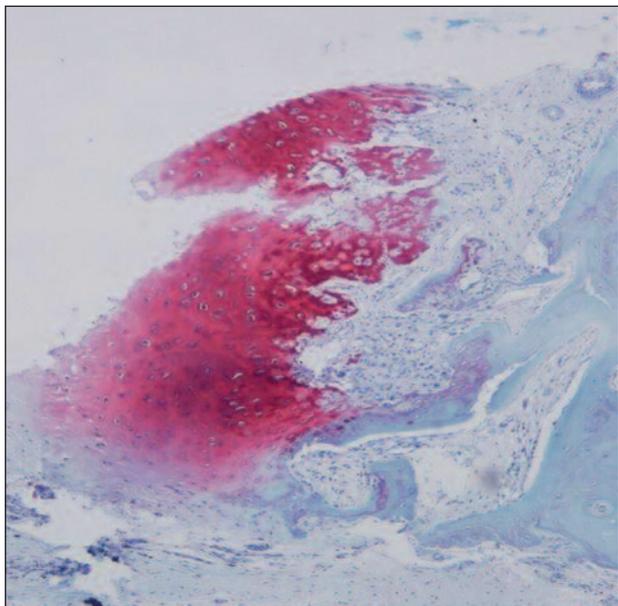


Fig. 4. Histological findings of the allograft. Biopsy specimen showed vital cartilage with a high proteoglycan content, normal cell distribution and complete osseointegration between the graft and the recipient bone. Safranin-O staining. Magnification 20x.

The immunosuppressive therapy was performed on the basis of the idea that the immune system plays a role in rejection of transplanted cartilage; the aim was to prevent the resulting damage from causing arthritis (22-24). The protocol we used was a light immunosuppressive therapy, already used in rheumatology, which was administered for six months. This protocol was easily tolerated by the patients and no adverse events were noted, however it was found to show poor protective efficacy against arthritis. Further studies are required in order to understand how to prevent arthritic changes in transplanted cartilage.

BFOA is a technically demanding surgical procedure, requiring a multidisciplinary approach and a bone bank facility. The significance of the present work lies in the fact that its results support the applicability of BFOA as an interesting surgical alternative to traditional arthroplasty in severe post-traumatic shoulder arthritis in young, active patients. The limited number of patients and the short follow-up are limitations of this analysis. Nevertheless, the fact that arthrodesis is poorly applicable for the shoulder, and shoulder replacement has limited indications in young patients, makes this possible alternative solution extremely appealing. However, the

indications for its application are extremely limited because of the need for an intact rotator cuff.

In conclusion, the clinical results obtained in the cases described were extremely satisfactory in terms of pain resolution, ROM, overall function of the transplanted shoulder, and patient satisfaction and quality of life. In one patient, however, the development of shoulder laxity led to clinical impairment; to date, no revision surgery has been performed in this patient. The histological results also confirmed good survivorship of the chondrocytes after implantation and good integration of the graft. In all the patients, despite correct graft positioning and size, the radiographic results at final follow up were disappointing, even though they did not impact on the clinical result.

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