Alma Mater Studiorum Università di Bologna Archivio istituzionale della ricerca

Surgical treatment of early knee osteoarthritis with a cell-free osteochondral scaffold: Results at 24 months of follow-up

This is the final peer-reviewed author's accepted manuscript (postprint) of the following publication:

Published Version:

Di Martino, A., Kon, E., Perdisa, F., Sessa, A., Filardo, G., Neri, M.P., et al. (2015). Surgical treatment of early knee osteoarthritis with a cell-free osteochondral scaffold: Results at 24 months of follow-up. INJURY, 46(S8), S33-S38 [10.1016/S0020-1383(15)30052-8].

Availability:

This version is available at: https://hdl.handle.net/11585/540937 since: 2016-06-07

Published:

DOI: http://doi.org/10.1016/S0020-1383(15)30052-8

Terms of use:

Some rights reserved. The terms and conditions for the reuse of this version of the manuscript are specified in the publishing policy. For all terms of use and more information see the publisher's website.

This item was downloaded from IRIS Università di Bologna (https://cris.unibo.it/). When citing, please refer to the published version.

(Article begins on next page)

This is the final peer-reviewed accepted manuscript of:

Di Martino, A.; Kon, E.; Perdisa, F.; Sessa, A.; Filardo, G.; Neri, M. P.; Bragonzoni, L.; Marcacci, M. Surgical Treatment of Early Knee Osteoarthritis with a Cell-Free Osteochondral Scaffold: Results at 24 Months of Follow-Up. *Injury* **2015**, *46 Suppl 8*, S33-38.

The final published version is available online at:

https://doi.org/10.1016/S0020-1383(15)30052-8

Rights / License:

The terms and conditions for the reuse of this version of the manuscript are specified in the publishing policy. For all terms of use and more information see the publisher's website.

This item was downloaded from IRIS Università di Bologna (https://cris.unibo.it/)

When citing, please refer to the published version.

Surgical treatment of early knee osteoarthritis with a cell-freeosteochondral scaffold: results at 24 months of follow-up

Alessandro Di Martino^a, Elizaveta Kon^a,*, Francesco Perdisa^a, Andrea Sessa^a, Giuseppe Filardo^a, Maria Pia Neri^a, Laura Bragonzoni^b, Maurilio Marcacci^a

a II Clinic, Biomechanics Laboratory, Rizzoli Orthopaedic Institute, Bologna, Italy

Purpose: "Early Osteoarthritis (EOA)" has been defined combining clinical, imaging and surgical param- eters, with the aim to identify patients in early degenerative phases, who might benefit from the use of available regenerative procedures. Aim of this first clinical trial is to prospectively evaluate the results obtained in a group of patients meeting the inclusion criteria of "EOA" as proposed by the ESSKA Cartilage Committee, and surgically treated with the implantation of a multi-phasic osteochondral scaffold.

Methods: 23 patients were prospectively evaluated at 12 and 24 months of follow-up. Etiology of the chondral or osteochondral defect was rated as microtraumatic or degenerative in 18 cases, and trau-matic in 5 cases. Patients included were complaining of clinical symptoms like knee pain and affected by chondral and osteochondral lesions located at the femoral condyles or trochlea and MRI findings demonstrating articular cartilage degeneration and/or meniscal degeneration and/or subchondral bone marrow lesions.

Results: All patients increased significantly in any clinical score adopted. The IKDC subjective score increased from 42.8 ± 13.8 at basal evaluation to 74.3 ± 17.4 at 12 months' (p < 0.0005), being stable (74.9 ± 20.4) up to the final follow-up of 24 months. Tegner score showed a statistically significant improvement in sports activity from 3.3 ± 2.7 pre-operative to 4.6 ± 2.2 at 12 months (p < 0.005), with a slight improvement to the final evaluation (4.7 ± 2.1 ; n.s.). However, the activity level was significantly lower than the pre-injury one (6.1 ± 2.6 ; p = 0.004). A significant difference was shown between patients younger versus older than 40 years, with younger patients had better clinical improvement (76.0 ± 18.6 vs 45.1 ± 38.8 respectively, p = 0.037).

Conclusions: The implantation of a multi-phasic osteochondral scaffold represents a good option after failure of conservative management for Early OA patients, where younger age represent an important factor for a better outcome.

Longer follow-up is needed to evaluate the benefit over time.

Introduction

Osteoarthritis (OA) is one of the most widespread orthopaedic diseases, characterized by joint pain and decreased function [1]. It is most common in patients over 50 years old, but with the increase of physical activity in the aging population it is not rare to find degenerated knees even in younger patients. In recent years, there has been an increasing interest in identi-fying early phases of OA, so that the

degenerative processes might be arrested or delayed. In this context, the definition of early OA has been recently proposed by combining both clinical, imaging, and surgical parameters, to identify patients in these early degen- erative phases at risk for progression, who might still benefit from the use of regenerative procedures. In fact, young patients with OA symptoms frequently present with an altered joint biomechanical environment responsible for an imbalance between the mechanical demand and the ability of the joint to maintain and repair itself, thus resulting in premature degenerative changes [2,3]. Thus, once identified, predisposing factors can be addressed to prevent fur-ther degeneration, favour

b Biomechanics Laboratory, Rizzoli Orthopaedic Institute, Bologna, Italy

a better chance of success with biolog-ical treatments and delay the need for more invasive procedures.

In fact, whereas metal resurfacing can provide a high success rate and satisfaction for older patients, high functional demand and longer life expectancy of young patients are an issue for joint arthroplasty. However, recommendations have tradition- ally excluded regenerative procedures for the treatment of carti-lage lesions in OA patients, because of the poor expected results. However, leaving degenerative defects untreated accelerates the rate of cartilage loss [4,5], and early OA patients, too young for joint replacement, might benefit from a biological reconstruction to restore the articular surface.

One of the main challenges in the treatment of degenerative lesions is that cartilage is not the only tissue involved. For exam- ple recently there has been increasing interest and awareness in the importance of the subchondral bone, that may be affected by the degenerative pathological processes as well [6]. Thus, to obtain a repair tissue that closely resembles the native articu-lar surface and might last over time, restoring the physiological properties of the entire osteochondral unit may be key in these degenerative patients. An osteochondral treatment is challeng-ing, due to the different healing capabilities of the two tissues, but new regenerative procedures are emerging as promising approaches to manage these lesions [7,8]. Among these, a bio- mimetic nanostructured osteochondral scaffold [9,10] was pro- posed to restore the articular surface, by addressing both the chondral layer and the underlying subchondral bone.

The aim of this clinical trial is to evaluate prospectively the results obtained in a group of patients meeting the criteria of "Early OA" and surgically treated with the implantation of a multi-phasic cell-free osteochondral scaffold.

Materials and methods

The study was approved by the Hospital Ethics Committee and Internal Review Board, and informed consent was obtained from each patient.

The patients met the previously proposed criteria for early OA

[11] "clinical symptoms, such as knee pain (at least two episodes of pain for more than 10 days in the last year) and MRI findings demonstrating articular cartilage degeneration and/or meniscal degeneration and/or subchondral BMLs", and affected by chon- dral and osteochondral lesions located at the femoral condyles or trochlea. Exclusion criteria were: lesions at the patella or tib-ial plateau, osteochondritis dissecans (OCD), and patients with non-corrected misalignment or instability of the knee. Patients presenting infectious, neoplastic, metabolic and inflammatory pathologies, as well as those not able to comply with the required post-operative rehabilitation regimen, were also excluded from this study. Conversely, patients with an axial deviation or an anterior cruciate ligament (ACL) lesion who underwent realign- ment or ligament reconstruction in the same surgical session as the scaffold implantation were included.

Twenty-four patients were consecutively enrolled and treated and 23 of them (19 men and 4 women) were prospectively evalu- ated pre-operatively, at 12 and 24 months of follow-up, whereas one patient was lost to follow-up. Mean age was 38.0 ± 8.2 years and average BMI was 25 ± 2.9 . Five patients had multiple lesions, mak- ing a total of 29 defects treated, located as follows: 12 medial fem- oral condyle (MFC), 9 lateral femoral condyle (LFC), 6 trochlea, plus 1 tibial plateau and 1 patella lesion that were included and treated as secondary lesions. Average defect size was 3.2 ± 1.9 cm². Etiology was rated as microtraumatic or degenerative in 18 cases, and trau-matic (not acute) in 5 cases. Seven patients were surgically treated for the first time, whereas 17 patients had undergone previous sur-geries (6 of them had previous cartilage surgery): 11 meniscectomy, 6 ACL reconstruction, 4 microfracturing, 4 debridement, 1 loose body removal, 1 tibial plateau fracture treatment, and 1 autologous chon- drocyte transplantation. In 17 patients other procedures were per-formed at the same time as surgery: 4 ACL reconstruction, 3 meniscal scaffold implantation, 2 meniscectomy, 1 loose body removal, 2 high tibial osteotomy, 2 distal femoral osteotomy, 1 meniscal allograft implantation, 1 microfracturing, and 1 postero-lateral corner repair.

Surgical procedure

The surgical procedure was performed with the patient under general or spinal anesthesia and in the

supine position with a pneumatic tourniquet around the proximal thigh. The defects were exposed through medial or lateral mini-arthrotomic para-patellar approach and prepared with an osteotome by removing the sclerotic subchondral bone. An 8-mm-deep lodging with perpendicular sides was created to allow press-fit fixation of the implant [12]. Stability was then visually tested by cyclic bending of the knee, both before and after tourniquet removal.

Patients evaluation

The patients were prospectively evaluated preoperatively and postoperatively at 12 and 24 months of follow-up. The clin- ical outcome was assessed for each patient using the Cartilage Standard Evaluation Form as proposed by the ICRS (International Cartilage Repair Society) [13]. The sport activity level was anal- ysed with the Tegner score and compared with pre-operative and pre-injury values.

Also, magnetic resonance imaging (MRI) evaluation of the implant was performed for 26 defects in 21 patients at 12 months and 21 lesions in 16 patients at 24 months of fol-low-up. Examinations were performed using a 1.5-T supercon-ducting magnet (General Electric Co, Fairfield, Connecticut) with a dedicated quadrature detection knee coil (Quadknee; diameter, 18 cm), using the same sequences previously described for this imaging analysis [14]. The MOCART scoring system was applied for the evaluation of the implant at follow-up times [15]. The eval-uation was performed in consensus by an orthopaedic surgeon and a musculoskeletal radiologist both experienced in cartilage procedures, who blindly assessed and reviewed the images.

Statistical methods

All continuous data were expressed in terms of the mean and the standard deviation of the mean, the categorical data were expressed as frequency and percentages. The Kolmogorov Smirnov test was performed to test normality of continuous var- iables. The Repeated Measures General Linear Model (GLM) with Sidak test for multiple comparisons was performed to assess the differences at different follow-up times. The Friedman non parametric test, followed by the Wilcoxon post hoc pairwise comparison corrected by Bonferroni method for multiple compari- sons, was used to the differences at different follow-up times of not normally distributed scores. The ANOVA test was performed to assess the between groups differences of continuous, normally distributed and homoscedastic data, the Mann Whitney test was used otherwise. The ANOVA test followed by the Scheffè post hoc pairwise comparison was used also to assess the among groups differences of continuous, normally distributed and homoscedas- tic data, the Kruskal Wallis test followed by the Mann Whitney test with the Bonferroni correction for multiple comparison was used otherwise. The Spearman rank Correlation was used to assess correlation between continuous data. The Kendall tau cor- relation was used to assess correlation between ordinal data. The Pearson Chi square test evaluated by Exact Methods for small samples was performed to investigate the relationships between grouping variables. The analysis on the MRI findings were eval-uated by the Monte Carlo method for small samples. For all tests p < 0.05 was considered significant.

All statistical analysis was performed using SPSS v.19.0 (IBM Corp., Armonk, NY, USA).

Results

No major adverse events were reported in the present series, besides two patients who required knee mobilization under nar- cosis, respectively 2 and 4 months after surgery, due to articular stiffness. Two patients (8.3%) failed, according to a more com- prehensive definition [16] including both surgical and clinical criteria, being re-treated due to persistent symptoms within the follow-up period.

The patients significantly improved in all the clinical scores adopted. In detail, the IKDC subjective score increased from

 42.8 ± 13.8 at basal evaluation to 74.3 ± 17.4 at 12 months (p < 0.0005), being stable (74.9 ± 20.4 , n.s.) up to the final follow-up of 24 months (Fig. 1). The Tegner score showed a statistically significant

improvement in sports activity from

 3.3 ± 2.7 pre-operatively to 4.6 ± 2.2 at 12 months (p < 0.005), with stable results at the final evaluation (4.7 ± 2.1; n.s.). However, the activity level at follow-up was significantly lower than that before injury (6.1 ± 2.6; p = 0.004) (Fig. 2).

A positive trend was also recorded by analysing the IKDC objective score. At the basal evaluation 12 knees (64.5%) were considered "normal" or "nearly normal" (6 grade A, 6 grade B). At the 12 months' evaluation a significant improvement was documented: 21 knees (14 grade A; 7 grade B) were considered "normal" or "nearly normal" (93.3%; p < 0.0005). This result was later confirmed at the final follow-up evaluation (19 "normal" or "nearly normal" knees of which 13 were grade A and 6 grade B).

A further analysis was performed to identify patient charac-teristics that might influence the clinical outcome. The follow- ing parameters did not correlate with the clinical results in this series: sex, BMI, lesion size, lesion site, and aetiology, concurrent or previous surgeries, while a significant difference in terms of IKDC subjective score improvement was shown for patients under 40 years old, who had better results (86.7 \pm 9 vs 67.3 \pm 22, respectively p = 0.037) (Fig. 3).

Twenty-six lesions in 21 patients and 21 lesions in 16 patients were evaluated by MRI at 12 and 24 months, respectively. At 12 months the MOCART parameters showed a complete filling of the cartilage layer in 65.3% of the lesions, complete integration of the graft in 76.9% of cases, intact surface of the repair tissue in 38.5% of the cases, homogeneous structure of the repair tis- sue in half of the cases, and iso-intense graft signal intensity with the adjacent native cartilage in 69.2% of the cases, inde-pendently from the sequence used. Subchondral bone alterations were present in 42.3%, and the lamina was intact in only 2 sites (7.7%). One patient had adhesions (3.9%), whereas effusion was observed in 65.4 % of the cases. At 24 months a complete fill- ing of the cartilage was shown in 61.9% of the lesions, complete integration of the graft was detected in 66.7% of cases, the repair tissue surface was intact in 57.1%, while its structure was nonhomogeneous in 61.9% of the cases. The graft signal intensity was iso-intense with the adjacent native cartilage in 57.1% and 66.6% of the cases in dual T2-FSE and 3D-GE-FS sequences, respec- tively. Subchondral bone alterations were observed in 57.1% of the cases, and the subchondral lamina appeared intact in 2 cases (9.5%). No adhesions were shown and 57.1% had some effusion. Finally, the total MOCART score was stable between 12 and 24 months of follow-up (72.9 \pm 13.6 and 70.8 ± 13.2, respectively), and no correlation was found for either the MOCART variables and total score with the clinical parameters.

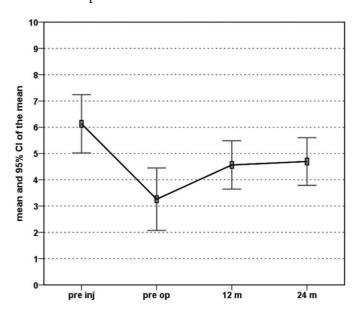


Fig. 1. IKDC subjective score evaluation up to 24 months of follow-up.

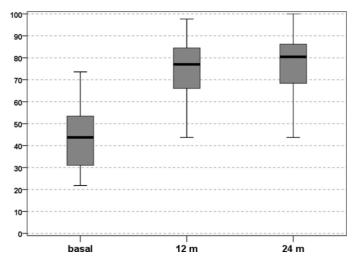


Fig. 2. The Tegner score significantly improved from pre-operative level, even if remained significantly lower than the pre-symptoms value.

Discussion

The main finding of the present study is that the implanta- tion of this osteochondral scaffold offered satisfactory results at short-term follow-up and may therefore be a possible treatment solution for patients affected by Early OA of the knee.

Since their introduction into the clinical practice two decades ago, regenerative approaches have shown they can promote the restoration of a hyaline-like articular surface [17] with satisfac-tory clinical outcomes [18,19]. In particular, the literature reports that these techniques may be successfully applied in young active patients with large chondral lesions, whereas limits emerged when applied to degenerative or complex joint disease [20-22].

Tissue engineering applied to the treatment of articu- lar degenerative lesions presents some additional problems: healthy tissues are key to provide stable sides for the implant, whereas in a degenerative process the surrounding areas may be involved, thus limiting the stability of the graft, and the altered

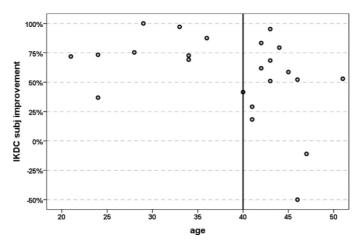


Fig. 3. Age differences: patients under 40 years old had better IKDC subjective score improvement.

joint environment may hinder the offered outcomes [23,24]. Preclinical trials have shown that a disturbed joint environment produces unfavorable conditions for tissue regeneration, with a negative

influence on cartilage formation [25-27]. Rodrigo et al.

[26] found that synovial fluid contains factors able to stimulate chondral healing in the acute period following traumatic injury, whereas it has inhibitory effects when the lesions turn chronic.

However, some authors have shown that regenerative proce-dures may still produce satisfactory results also in joints affected by degenerative changes. Hollander et al. [28] observed tissue regeneration even for implants inside OA joints, and laboratory studies confirmed the potential usefulness of regenerative pro-cedures in joints with degenerative lesions, even when the osteo- arthritic process has already started [29,30], thus suggesting that OA does not inhibit the regeneration process and justifying a pos- sible clinical use [29].

Interesting findings have been reported using regenerative procedures in humans. Minas et al. [20] treated 153 patients with autologous chondrocyte transplantation (ACT) for early-stage OA. Good function was observed at 5 years in 92% of them, thus delaying the need for joint replacement. Ossendorf et al. [31] reported pain, symptoms, and quality of life improvement after the implantation of a polymer-based autologous cartilage graft for mild degenerative cartilage lesions or focal knee OA defects. Subsequently, Kreuz et al. [32] confirmed stable results for 4 years, with a significant improvement in the patients' condition and a good defect filling on MRI. However, Filardo et al. [22] analysed the clinical outcome of patients treated with 2nd generation ACT for small or medium isolated degenerative knee cartilage lesions, with no signs of OA, and showed poorer results and higher failure rates than those reported in not degenerative patient populations [32-34], and even worse results have been reported by the same group treating focal lesions in OA joints [21]. Finally, Nehrer et al. used MACT as a salvage procedure for patients with kissing lesions or early OA changes, previously treated for the same carti-lage defect, and found 87.5% of failures in this category [35].

A possible reason for the poor reported outcome with carti-lage regenerative procedures in degenerative lesions is the long time lapse between cartilage damage and onset of the clinical symptoms, which is likely in case of OA changes and may result in treatment delay, that has been shown to be a factor limiting the success of regenerative procedures [34]. In fact, in degen-erative lesions an imbalance of joint homeostasis has already occurred with subchondral bone involvement in the pathologic changes [27,36].

Recently, the development of biomaterial technology has led to new constructs, thus offering the possibility to treat subchon-dral bone disease by implanting multi-layered scaffolds designed to provide the regeneration of cartilage and bone tissues, both altered in a degenerative joint environment [9]. A previous com-parative study showed that the same ostechondral scaffold used in this series produced better results than a chondral one for the treatment of complex knee lesions [37], and promising findings had been reported even in unicompartmental OA patients [38].

The results of this study confirm the potential of this cell- free multi-layer scaffold in providing a successful outcome in early OA joints. Nonetheless, patients presented mainly a partial improvement and the overall results are poorer than those pre-viously reported using the same procedure in different groups of patients [12,39,40]. Thus, although an osteochondral scaffold may be successful in restoring the damaged articular surface, this approach may not take into account other important aspects key for a complete success in this patient population. Results are probably limited by the degenerative and inflammatory changes that affect the entire environment of a joint that has already undertaken the path of OA [2], thus making a treatment of the damaged articular surface, even with the correction of concur- rent biomechanical alterations, inadequate to address the multi- ple targets related to the disease.

Moreover, early OA typically affects patients in an age range where the biological healing potential may be already reduced. The importance of age has been confirmed in this series, where older patients presented a reduced clinical improvement. This is not surprising, since the potential of cartilage procedures, in particular in case of a cell-free approach, relies on the body's self-renewing potential, which is an age-related feature [41]. Age leads to degenerative changes in all cartilage elements [42], thus impairing its properties and healing potential [43]. Accordingly, older patients have traditionally been excluded from regenera-tive treatment recommendations, and only a few studies reportresults in this patient population thus confirming the limited outcome regardless of the surgical technique used.

Microfracturing (MF) produces age-dependent results and a greater improvement in patients younger than 35 years, as observed by Steadman et al. [44]. Moreover, Kreuz et al. [41] reported poorer clinical results in patients over 40 years old. Marcacci et al. found a better clinical and functional outcome in younger patients treated with mosaicplasty procedure [45], so did Gudas et al. [46].

Similarly, even though cell-based regenerative approaches have been successfully tested in older patients, they offered poorer outcomes than those expected for younger ones. Kon et al. used MACT in non-OA patients over 40 years old [47], with a significant improvement in all scores at medium-term follow-up, but inferior results and higher failure rates with respect to younger study populations. Knutsen et al. reported better clin- ical outcomes in active and younger (<30 years) patients [48], who had either undergone MF or ACT. Krishnan et al used collagen-covered ACT for the treatment of 199 patients, with a negative correlation between clinical result and older age [49]. More recently, this age dependency was confirmed by de Windtet al. [50] when analysing microfracture or 1st or 2nd generation ACT at the knee, and reported a better clinical improvement in patients under 30 years old.

Although most authors agree that the outcomes of different cartilage procedures are age-dependent, there is no agreement on a precise age cutoff, and thresholds vary in different studies. This may be due to the fact that the biological stage of the joint may be as important as the patient age itself. With regards to this, a new classification was proposed for "Early" OA, to focus better on the status of the affected joint, by identifying patients before the "point of no return". This might be a key aspect for a positive clinical result, rather than age, as suggested also by the interesting results obtained by other authors in older patients affected by cartilage lesions [51]. This paves the way to success-fully defining a patient category with older age but who can still benefit from biological procedures to improve symptoms and delay further joint degeneration.

This study shows that this patient category may benefit from a biological reconstruction, even though the partial improve- ment documented underlines the complexity of the early OA environment, with degenerative processes that may have already affected the whole joint. Inflammatory, cellular or molecular fac- tors, are involved in OA development [27], and cytokine secre- tion [52] from degenerative tissues around the implant might cause dedifferentiation or apoptosis and impair the quality of the regenerative tissue [53]. Thus, an increased knowledge of the whole joint degenerative biochemical environment is crucial to develop integrated treatments, able to improve the obtainable clinical outcome by addressing both the osteochondral unit and the intra-articular homeostasis. In the meanwhile, until new integrated approaches have been developed, the indication of this cell-free osteochondral scaffold implantation for early OA joints should be limited and considered as a salvage procedure for compromised knees of young patients, otherwise doomed to more invasive and sacrificing procedures.

This study presents some weaknesses: first, it lacks histolog- ical evaluations. However, this is a preliminary study testing the effects of a scaffold-based regenerative treatment in a group of patients affected by Early OA, with most of the patients having a significant clinical benefit in the short-term after the implan- tation of this osteochondral substitute. Moreover, MRI findings showed the presence of several abnormal findings. However, the MOCART evaluation of the MRI scans showed stable results between 12 and 24 months of follow-up. Accordingly with most of the available Literature [54,55], we have found a lack of cor-relation between the clinical outcome and MOCART parameters: MRI is a useful and easy tool to evaluate the repair tissue, but its role in predicting or reflecting the clinical outcome is question-able. The available MRI scoring systems have been specifically focused on the chondral layer, thus they don't take sufficient account of the complexity of ostechondral regenerative proce- dures. Maybe new scoring tools or the use of CT scans might give a better evaluation of tissue quality and clinical significance.

A further weakness is the relatively high number of combined surgical procedures, that may confuse the evaluation of the scaf-fold implantation, thus making it difficult to determine whether the clinical benefit is related to osteochondral defect treatment or to the concurrent procedures. However, OA is a multifactorial disease and chondral lesions are frequently a combined feature in this pathology, thus this kind of patient is unlikely to be found with isolated lesions and the patients documented in this

study reflect those found in the clinical practice. The lack of a control group is another weak point. Finally, longer follow-up studies are needed to evaluate whether the positive outcomes obtained remain stable over time. However, despite the above-mentioned limitations, this study is the first one to focus on this specific patient population treated with this ostecochondral scaffold, and demonstrate that this approach may provide a clinical improvement for the treatment of cartilage defects in an early OA degenerative context. This is particularly relevant consider-ing that even small and isolated defects of the articular surface can lead to more extensive joint damage [56, 57] and accelerate the degenerative process of the entire joint. Thus, the surgical treatment of chondral lesions in early OA patients may be useful not only for pain relief, but also to avoid or delay further joint degeneration and the need for joint replacement. Nonetheless, we recommend giving patients realistic expectations when con-sidering this surgical approach for early OA joints.

Conclusion

The implantation of this cell-free osteochondral scaffold offered satisfactory results at short-term follow-up and may therefore be a possible treatment solution for patients affected by early OA of the knee. However, patients presented mainly a partial improvement and the overall results are lower than those previously reported using the same procedure in different group of patients. Until new integrated approaches are developed to address all the factors responsible for the altered environment of degenerated joints, the indication of this treatment for early OA joints should be limited and considered as a salvage procedure for compromised knees of young patients.

Acknowledgments

- F. Balboni: II Clinic Biomechanics Laboratory, Rizzoli Orthopaedic Institute, Bologna, Italy.
- E. Pignotti, K. Smith: Task Force, Rizzoli Orthopaedic Institute, Bologna, Italy.

The research leading to these results has received funding from the European Union's Seventh Framework Programme (FP/2007-2013) under grant agreement number 278807.

References

- Bijlsma JW, Berenbaum F, Lafeber FP. Osteoarthritis: an update with relevance for clinical practice. Lancet 2011;377:2115–26.
- [2] Madry H, Luyten FP, Facchini A. Biological aspects of early osteoarthritis. Knee Surg Sports Traumatol Arthrosc 2012;20:407–22.
- [3] Heijink A, Gomoll AH, Madry H, Drobnic M, Filardo G, Espregueira-Mendes J, et al. Biomechanical considerations in the pathogenesis of osteoarthritis of the knee. Knee Surg Sports Traumatol Arthrosc 2012;20:423–35.
- [4] Cicuttini F, Ding C, Wluka A, Davis S, Ebeling PR, Jones G. Association of cartilage defects with loss of knee cartilage in healthy, middle-age adults: a prospective study. Arthritis Rheum 2005;52:2033–9.
- Davies-Tuck ML, Wluka AE, Wang Y, Teichtahl AJ, Jones G, Ding C, et al. The natural history of cartilage defects in people with knee osteoarthritis. Osteoarthritis Cartilage 2008;16:337–42.
- [6] Pape D, Filardo G, Kon E, van Dijk CN, Madry H. Disease-specific clinical problems associated with the subchondral bone. Knee Surg Sports Traumatol Arthrosc 2010;18:448–62.
- [7] Marcacci M, Filardo G, Kon E. Treatment of cartilage lesions: what works and why? Injury 2013;44(Suppl 1):S11–5.
- [8] Filardo G, Kon E, Berruto M, Di Martino A, Patella S, Marcheggiani Muccioli GM, et al. Arthroscopic second generation autologous chondrocytes implantation associated with bone grafting for the treatment of knee osteochondritis disse- cans: Results at 6 years. Knee 2012;19:658–63.

- [9] Kon E, Mutini A, Arcangeli E, Delcogliano M, Filardo G, Nicoli Aldini N, et al. Novel nanostructured scaffold for osteochondral regeneration: pilot study in horses. J Tissue Eng Regen Med 2010;4:300–8.
- [10] Kon E, Delcogliano M, Filardo G, Altadonna G, Marcacci M. Novel nano- composite multi-layered biomaterial for the treatment of multifocal degener- ative cartilage lesions. Knee Surg Sports Traumatol Arthrosc 2009;17:1312–5.
- Luyten FP, Denti M, Filardo G, Kon E, Engebretsen L. Definition and classification of early osteoarthritis of the knee. Knee Surg Sports Traumatol Arthrosc 2012;20:401–6.
- [12] Filardo G, Kon E, Di Martino A, Busacca M, Altadonna G, Marcacci M. Treatment of knee osteochondritis dissecans with a cell-free biomimetic osteochondral scaffold: clinical and imaging evaluation at 2-year follow-up. Am J Sports Med 2013 41:1786–93.
- [13] ICRS. ICRS Cartilage Evaluation Package. 2000.
- [14] Kon E, Filardo G, Di Martino A, Busacca M, Moio A, Perdisa F, et al. Clinical Results and MRI Evolution of a Nano-Composite Multilayered Biomaterial for Osteochondral Regeneration at 5 Years. Am J Sports Med 2014;42:158–65.
- Marlovits S, Striessnig G, Resinger CT, Aldrian SM, Vecsei V, Imhof H, et al. Definition of pertinent parameters for the evaluation of articular cartilage repair tissue with high-resolution magnetic resonance imaging. Eur J Radiol 2004;52:310–9.
- [16] Filardo G, Andriolo L, Balboni F, Marcacci M, Kon E. Cartilage failures. Systematic literature review, critical survey analysis, and definition. Knee Surg Sports Traumatol Arthrosc 2014 [Epub ahead of print].
- [17] Brittberg M, Lindahl A, Nilsson A, Ohlsson C, Isaksson O, Peterson L. Treatment of deep cartilage defects in the knee with autologous chondrocyte transplanta- tion. New Engl J Med 1994;331:889-95.
- [18] Browne JE, Anderson AF, Arciero R, Mandelbaum B, Moseley JB, Jr., Micheli LJ, et al. Clinical outcome of autologous chondrocyte implantation at 5 years in US subjects. Clin Orthop Rel Res 2005:237–45.
- [19] Peterson L, Brittberg M, Kiviranta I, Akerlund EL, Lindahl A. Autologous chondrocyte transplantation. Biomechanics and long-term durability. Am J Sports Medicine 2002;30:2–12.
- Minas T, Gomoll AH, Solhpour S, Rosenberger R, Probst C, Bryant T. Autologous chondrocyte implantation for joint preservation in patients with early osteoarthritis. Clin Orthop Rel Res 2010;468:147–57.
- Filardo G, Vannini F, Marcacci M, Andriolo L, Ferruzzi A, Giannini S, et al. Matrix-assisted autologous chondrocyte transplantation for cartilage regener- ation in osteoarthritic knees: results and failures at midterm follow-up. Am J Sports Med 2013;41:95–100.
- Filardo G, Kon E, Di Martino A, Patella S, Altadonna G, Balboni F, et al. Second- generation arthroscopic autologous chondrocyte implantation for the treat- ment of degenerative cartilage lesions. Knee Surg Sports Traumatol Arthrosc 2012;20:1704–13.
- Buckwalter JA, Mankin HJ. Articular cartilage: degeneration and osteoarthritis, repair, regeneration, and transplantation. Instr Course Lect 1998;47:487–504.
- Buckwalter JA, Mankin HJ. Articular cartilage repair and transplantation. Arthritis Rheum 1998;41:1331–42.
- Ozsoy MH, Aydogdu S, Taskiran D, Sezak M, Hayran M, Oztop F, et al. The effects of early or late treatment of osteochondral defects on joint homoeo- stasis: an experimental study in rabbits. Knee Surg Sports Traumatol Arthrosc 2009;17:578–89.
- Rodrigo JJ, Steadman JR, Syftestad G, Benton H, Silliman J. Effects of human knee synovial fluid on chondrogenesis in vitro. Am J Knee Surg 1995;8:124–9.
- Saris DB, Dhert WJ, Verbout AJ. Joint homeostasis. The discrepancy between old and fresh defects in cartilage repair. J Bone Joint Surg Br 2003;85:1067–76.
- [28] Hollander AP, Dickinson SC, Sims TJ, Brun P, Cortivo R, Kon E, et al. Maturation of tissue engineered cartilage implanted in injured and osteoarthritic human knees. Tissue Eng

- 2006;12:1787–98.
- [29] Cavallo C, Desando G, Facchini A, Grigolo B. Chondrocytes from patients with osteoarthritis express typical extracellular matrix molecules once grown onto a three-dimensional hyaluronan-based scaffold. J Biomed Mater Res A 2010;93:86–95.
- [30] Tallheden T, Bengtsson C, Brantsing C, Sjogren-Jansson E, Carlsson L, Peterson L, et al. Proliferation and differentiation potential of chondrocytes from osteoar-thritic patients. Arthritis Res Ther 2005;7:R560–8.
- Ossendorf C, Kaps C, Kreuz PC, Burmester GR, Sittinger M, Erggelet C. Treatment of posttraumatic and focal osteoarthritic cartilage defects of the knee with autologous polymer-based three-dimensional chondrocyte grafts: 2-year clinical results. Arthritis Res Ther 2007;9:R41.
- [32] Kreuz PC, Muller S, Ossendorf C, Kaps C, Erggelet C. Treatment of focal degen- erative cartilage defects with polymer-based autologous chondrocyte grafts: four-year clinical results. Arthritis Res Ther 2009;11:R33.
- [33] Ferruzzi A, Buda R, Faldini C, Vannini F, Di Caprio F, Luciani D, et al. Autologous chondrocyte implantation in the knee joint: open compared with arthroscopic technique. Comparison at a minimum follow-up of five years. J Bone Joint Surg Am 2008;90(Suppl 4):90–101.
- [34] Filardo G, Kon E, Andriolo L, Di Matteo B, Balboni F, Marcacci M. Clinical profiling in cartilage regeneration: prognostic factors for midterm results of matrix-assisted autologous chondrocyte transplantation. Am J Sports Med 2014;42:898–905.
- Brix MO, Stelzeneder D, Chiari C, Koller U, Nehrer S, Dorotka R, et al. Treatment of Full-Thickness Chondral Defects With Hyalograft C in the Knee: Long-term Results. Am J Sports Med 2014;42:1426–32.
- [36] Gomoll AH, Madry H, Knutsen G, van Dijk N, Seil R, Brittberg M, et al. The sub-chondral bone in articular cartilage repair: current problems in the surgical management. Knee Surg Sports Traumatol Arthrosc 2010;18:434–47.
- Filardo G, Kon E, Perdisa F, Di Matteo B, Di Martino A, Iacono F, et al. Osteochondral scaffold reconstruction for complex knee lesions: a comparative evaluation. Knee 2013;20:570–6.
- Marcacci M, Zaffagnini S, Kon E, Marcheggiani Muccioli GM, Di Martino A, Di Matteo B, et al. Unicompartmental osteoarthritis: an integrated biomechanical
- [39] and biological approach as alternative to metal resurfacing. Knee Surg Sports Traumatol Arthrosc 2013;21:2509–17.
- [40] Kon E, Filardo G, Di Martino A, Busacca M, Moio A, Perdisa F, et al. Clinical results and MRI evolution of a nano-composite multilayered biomaterial for osteochondral regeneration at 5 years. Am J Sports Med 2014;42:158–65.
- [41] Kon E, Filardo G, Perdisa F, Di Martino A, Busacca M, Balboni F, et al. A one-step treatment for chondral and osteochondral knee defects: clinical results of a biomimetic scaffold implantation at 2 years of follow-up. J Mater Sci Mater Med 2014;25:2437–44.
- [42] Kreuz PC, Erggelet C, Steinwachs MR, Krause SJ, Lahm A, Niemeyer P, et al. Is microfracture of chondral defects in the knee associated with different results in patients aged 40 years or younger? Arthroscopy 2006;22:1180–6.
- Buckwalter JA, Roughley PJ, Rosenberg LC. Age-related changes in cartilage proteoglycans: quantitative electron microscopic studies. Microsc Res Tech 1994;28:398–408.
- Plaas AH, Sandy JD. Age-related decrease in the link-stability of proteoglycan aggregates formed by articular chondrocytes. Biochem J 1984;220:337–40.
- [45] Steadman JR, Briggs KK, Rodrigo JJ, Kocher MS, Gill TJ, Rodkey WG. Outcomes of microfracture for traumatic chondral defects of the knee: average 11-year follow-up. Arthroscopy 2003;19:477–84.
- [46] Marcacci M, Kon E, Zaffagnini S, Iacono F, Neri MP, Vascellari A, et al. Multiple osteochondral arthroscopic grafting (mosaicplasty) for cartilage defects of the knee: prospective study results at 2-year follow-up. Arthroscopy 2005;21:462–70.

- [47] Gudas R, Gudaite A, Pocius A, Gudiene A, Cekanauskas E, Monastyreckiene E, et al. Ten-year follow-up of a prospective, randomized clinical study of mosaic osteochondral autologous transplantation versus microfracture for the treat- ment of osteochondral defects in the knee joint of athletes. Am J Sports Med 2012;40:2499–508.
- ^[48] Kon E, Filardo G, Condello V, Collarile M, Di Martino A, Zorzi C, et al. Second- generation autologous chondrocyte implantation: results in patients older than 40 years. Am J Sports Med 2011;39:1668–75.
- [49] Knutsen G, Drogset JO, Engebretsen L, Grontvedt T, Isaksen V, Ludvigsen TC, et al. A randomized trial comparing autologous chondrocyte implantation with microfracture. Findings at five years. J Bone Joint Surg Am 2007;89:2105–12.
- [50] Krishnan SP, Skinner JA, Bartlett W, Carrington RW, Flanagan AM, Briggs TW, et al. Who is the ideal candidate for autologous chondrocyte implantation? J Bone Joint Surg Br 2006;88:61–4.
- [51] de Windt TS, Concaro S, Lindahl A, Saris DB, Brittberg M. Strategies for patient profiling in articular cartilage repair of the knee: a prospective cohort of patients treated by one experienced cartilage surgeon. Knee Surg Sports Traumatol Arthrosc 2012;20:2225–32.
- Miller BS, Steadman JR, Briggs KK, Rodrigo JJ, Rodkey WG. Patient satisfaction and outcome after microfracture of the degenerative knee. J Knee Surg 2004;17:13–7.
- Borzi RM, Mazzetti I, Marcu KB, Facchini A. Chemokines in cartilage degrada- tion. Clin Orthop Rel Res 2004:S53–61.
- [54] Leyh M, Seitz A, Durselen L, Schaumburger J, Ignatius A, Grifka J, et al. Subchondral bone influences chondrogenic differentiation and collagen production of human bone marrow-derived mesenchymal stem cells and articular chondrocytes. Arthritis Res Ther 2014;16:453.
- [55] Blackman AJ, Smith MV, Flanigan DC, Matava MJ, Wright RW, Brophy RH. Correlation between magnetic resonance imaging and clinical outcomes after knee cartilage repair: author's response. Am J Sports Med 2013;41:NP49–50.
- [56] de Windt TS, Welsch GH, Brittberg M, Vonk L, Marlovits S, Trattnig S, et al. Correlation between magnetic resonance imaging and clinical outcomes after knee cartilage repair: letter to the editor. Am J Sports Med 2013;41:NP48–50.
- [57] Gratz KR, Wong BL, Bae WC, Sah RL. The effects of focal articular defects on cartilage contact mechanics. J Orthop Res 2009;27:584–92.
- [58] Gratz KR, Wong BL, Bae WC, Sah RL. The effects of focal articular defects on intra-tissue strains in the surrounding and opposing cartilage. Biorheology 2008;45:193–207.