Comparison of Magnetic Resonance Imaging Findings in Anterior Cruciate Ligament Grafts With and Without Autologous Platelet-Derived Growth Factors

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Purpose: To determine whether the use of platelet-rich plasma gel (PRPG) affects magnetic resonance imaging (MRI) findings in the anterior cruciate ligament (ACL) graft during the first year after reconstruction. Methods: A prospective single-blinded study of 50 ACL reconstructions in 50 patients was performed. In group A (study group) PRPG was added to the graft with a standardized technique, and in group B (control group) no PRPG was added. An MRI study was performed postoperatively between 3 and 9 months in group A and between 3 and 12 months in group B. The imaging analysis was performed in a blind protocol by the same radiologist. Results: The mean heterogeneity score value at the time of MRI, assigned by the radiologist, was 1.14 in group A and 3.25 in group B. Both groups were comparable in terms of sex and age (P < .05). The mean time to obtain a completely homogeneous intra-articular segment in group A (PRPG added) was 177 days after surgery, and it was 369 days in group B. Using the quadratic predictive model, these findings show that group A (PRPG added) needed only 48% of the time group B required to achieve the same MRI image (P < .001). Conclusions: ACL reconstruction with the use of PRPG achieves complete homogeneous grafts assessed by MRI, in 179 days compared with 369 days for ACL reconstruction without PRPG. This represents a time shortening of 48% with respect to ACL reconstruction without PRPG. Level of Evidence: Level III, case-control study.

Rupture of the anterior cruciate ligament (ACL) is an injury commonly observed in sports medicine. Return to professional sports occurs at around 6 to 7 months, depending on the sport practiced. In sports medicine this time period is often very long for the athlete; thus methods have been sought to shorten the biological time required for the graft to acquire biomechanical properties similar to the original ACL.

The clinical results of ACL reconstruction and time to return to sports could be improved if the graft healing process is enhanced. In a classic publication on this topic in 1982, Arnoczky and Tarvin described the behavior of the graft used in ACL reconstruction in dogs, describing 3 stages in the process of graft metaplasia: incorporation, neoligament formation, and remodeling.

Various authors have tried to study the behavior of the graft in clinical trials, with histology or imaging studies, which experienced a significant boost with the appearance of magnetic resonance imaging (MRI).

In 1995, in a prospective clinical study that relied on second-look arthroscopy to perform a histologic and MRI assessment of the graft at 6, 9, and 12 months of postoperative evolution, we described...
how the patellar tendon graft used in human ACL reconstruction is incorporated. We concluded that the graft maturation takes a long time: 12 months to achieve histology similar to a normal ACL. At 12 months, the MRI study of the graft was homogeneous and hyperintense, without swelling in the bone tunnels. The correlation of the histology with MRI was of great help in establishing a reliable imaging pattern, which allowed us to noninvasively verify the graft healing process.

Weiler et al.\textsuperscript{12} report correlations between biomechanical properties and vascularity of an ACL graft and MRI in a sheep model.

Clinical applications of autologous platelet-rich plasma gel (PRPG) include maxillofacial surgery, treatment of bone fractures, and tendon repair, reporting excellent outcomes.\textsuperscript{13-16} Platelets contain different growth factors that facilitate healing. PRPG is a fraction of plasma volume with a platelet concentration above baseline (whole blood). Platelet concentrates contain an enormous amount of activated platelet-derived growth factors (PDGFs).\textsuperscript{17-23}

Platelets contain PDGFs, transforming growth factors (TGFs), insulin-like growth factors, epidermal growth factors, vascular endothelial growth factors, and fibroblast growth factors. These factors are involved in the majority of biological remodeling processes in the body. In the specific case of ACL graft, PDGFs, fibroblast growth factor 1, and the various types of TGF-\(\beta\) are responsible for accelerating the healing process, as well as increasing the tensile strength of the graft.\textsuperscript{24-30}

Only 2 articles have shown an enhancing effect of treatment with PRPG on the tendon or ligament in humans. In a human study Orrego et al.\textsuperscript{31} showed an enhancing effect over the graft maturation process as evaluated by MRI signal intensity, without showing a significant effect on the osteoligamentous interface or tunnel widening evolution. In human tenocyte cultures, de Mos et al.\textsuperscript{32} showed that PRPG stimulates cell proliferation and collagen production.

Currently, it is practically impossible to perform human clinical trials of biomechanical or histologic assessments of the graft’s behavior in ACL reconstruction. For this reason, we decided to practice an indirect and noninvasive assessment in our patients, using MRI. The purpose of our investigation was to study MRI findings in the ACL graft when PRPG was added during surgery, thus allowing future studies correlating MRI findings with histology and ultimate load and strength. We hypothesized that PRPG has a positive effect on cell proliferation and collagen production in the human tendon and plays a key role in the remodeling and repair processes of the graft used in ACL reconstruction.

**METHODS**

**Study Design**

This is a prospective and single-blinded study performed between June 2005 and December 2006. The inclusion criteria were sport athletes of both gender between 18 and 35 years old with an isolated ACL tear shown by MRI. Exclusion criteria were previous ACL revision surgery, chronic or systemic disease under treatment, and previous or current treatment for malignant disease. These pathologies can modify the biologic behavior of the graft. Fifty consecutive patients met the inclusion criteria.

The type of graft used was determined according to our institution’s protocol, and it depended on the type of sports the patient practiced. Bone–patellar tendon–bone (BPTB) autograft was used in rugby and soccer players, whether hamstring autograft was used in players who practiced skiing, hockey, tae kwon do, and volleyball. One of the surgeons (R.Y.) did not use PRPG, and the other (F.R.) did. Two groups were established: Group A included 25 patients (18 men and 7 women), with a mean age of 30 years (range, 18 to 33 years), with ACL reconstruction plus PRPG; 15 of these patients underwent reconstruction with BPTB. Group B included 25 patients (21 men and 4 women), with a mean age of 32 years (range, 18 to 35 years), with ACL reconstruction without PRPG; 10 of these patients underwent reconstruction with hamstring autografts. Both groups (A and B) followed the same rehabilitation protocol.

**Surgical Technique**

In the case of BPTB autograft, fixation was performed with metallic interference screws. Hamstring autograft fixation was performed with metallic or bioabsorbable cross-pin femoral fixation using the Trans-Fix technique (Arthrex, Naples, FL) in the distal femur and a Delta-type bioabsorbable screw with a metallic staple in the proximal tibia (Arthrex).

In group A PRPG was administered by an application technique developed to allow standardization of the dose of concentrate used and avoidance of its loss when the graft goes through the bone tunnels.\textsuperscript{33,34} The autologous platelet concentrate was obtained from the GPS system of Biomet (Warsaw, IN). This procedure is done aseptically in the same operating room. Pre-
operatively, 60 mL of autologous blood is obtained and centrifuged at 3,200 rpm for 15 minutes. In the case of BPTB graft, after adaptation of the bone plugs for them to fit through the tunnels, the femoral bone segment and the intra-articular segment are wrapped with a bioabsorbable synthetic gelatin called Gelfoam (Pfizer, New York, NY) and secured to it with a No. 3-0 Vicryl suture (Ethicon, Somerville, NJ). In the case of hamstring tendon graft, it is prepared in the usual manner with removal of the remnant muscle tissue. At each end, a 3-cm-long braid with FiberWire (Arthrex) is made, and the tendon’s thickness and length are measured. Under moderate tension, a piece of Gelfoam is placed between the portion of the tendons that will be located in the femoral tunnel and the intra-articular segment. This is sutured to the adjacent tendon with No. 3-0 Vicryl. The Gelfoam acts as a sponge that maintains the platelet concentrate dose in direct contact with the graft used (Fig 1). A total volume of 5 mL of platelet-rich plasma, activated at the moment of inoculation on the graft, is added homogeneously so as to completely cover the graft, waiting until it forms a clot (Fig 2). The dose administered was determined based only on our criteria. We do not know the ideal dose, and at the time of this study, nothing about the ideal dose had been published. The formed clot adheres to the graft, because of the presence of the sutured and compressed Gelfoam. This allows the graft to hold a precise amount of PRPG and, even more importantly, avoids the loss of PRPG when the graft goes through the bone tunnels (Fig 3).

![Figure 1. Hamstring graft preparation with Gelfoam and PRPG.](image1)

![Figure 2. The Gelfoam acts as a sponge that maintains the PRPG dose in direct contact with the graft.](image2)

### Imaging Assessment

The imaging protocol was standardized and similar in both groups. Included were a series of MRI scans focused to study the intra-articular segment of the graft. The study of the femoral and tibial parts of the graft was not considered because its maturation process occurs first, compared with the intra-articular part. This was performed with a T1 and T2 sequence (repetition time, 4,020 milliseconds; echo time, 105 milliseconds) with a 1.5-T Siemens Magnetom MRI Scanner (Siemens AG, Erlangen, Germany). Slices of 2 mm in thickness, in the oblique parasagittal view, between 10° and 15°, centered on the intercondylar region, with the knee flexed at 9° to 10°,11 were obtained. Patients in group A had MRI performed at 3, 4, 5, 6, 7, 8, and 9 months postoperatively so as to build a homogenization curve of the graft, according to the statistic quadratic predictive model, and supported by this study’s hypothesis that the use of PRPG accelerates the graft homogenization time. The control group had MRI performed at 6, 7, 8, 9, 10, 11, and 12 months, with the assumption that before 6 months, homogenization is not present.

The imaging analysis was done by the same radiologist, experienced in musculoskeletal studies, blinded to the time of reconstruction and to PRPG application to the graft.

The radiologist divided the intra-articular segment of the graft into 3 segments: proximal, medial, and distal. To each segment, he assigned a score according to the degree of heterogeneity observed. Therefore a score of 0 was assigned to an absolutely homogeneous
segment (Fig 4): 1, slightly heterogeneous; 2, moderately heterogeneous; and 3, severely heterogeneous (Fig 5). A sum of the scores for each segment was
obtained for each patient, which was compared statistically between the 2 groups and correlated with the time at which the MRI study was done.

Statistical Analysis

For the statistical analysis, data were analyzed with the SPSS data analysis program (SPSS, Chicago, IL). This program was used to work with quadratic statistics, graphs, data, and descriptive indicators. To determine whether the 2 groups were comparable in terms of number, age, and sex, an F test and Student t test were used.

The quadratic predictive model was used for data analysis to determine, through a linear relation, the extrapolated midpoint that predicted the time when both groups had completely homogeneous grafts.

RESULTS

The mean heterogeneity score value at the time of MRI, assigned by the radiologist, was 1.14 in group A and 3.25 in group B. Both groups were comparable in terms of sex and age (P > .05). The mean time to obtain a completely homogeneous intra-articular segment in group A (PRPG added) was 177 days after surgery, and it was 369 days in group B. Using the quadratic predictive model, the percentage of time that group A (PRPG added) needed to achieve the same MRI aspect as group B was 48% (Fig 6). This fact is even more evident when we compared only the BPTB graft cases in both groups: a homogeneous graft was obtained in 109 days in patients with PRPG versus 363 days in the control group, that is, one third the time as that for control group (Fig 7). In the comparative analysis of those patients in whom BPTB graft was used, we observed an even shorter time required for the graft’s homogenization when PRPG was used.
but this can only be considered a trend, because the sample’s number was too small to draw conclusions with statistical significance (β type error).

The latter fact, despite the large difference between the groups, merely shows a statistical trend, because it lacks statistical significance. On the other hand, to certify the findings of the quadratic model, in both groups only the patients who fully completed the requirements of a return to sports without restrictions, with normal functional sport-specific and isokinetic evaluations, were selected. The mean time to obtain an MRI score of 0, that is, a completely homogeneous graft, was compared between the groups (12 in the PRPG group and 6 in the control group). It was determined that the mean time in days to obtain a homogeneous graft was 179 days and 362 days in the PRPG and control groups, respectively (Fig 8). This finding indicates a decrease by half of the time (49.4%) in the group with PRPG versus the control group (P < .001). Once again, modifying the data analysis, the homogenization time of the intra-articular segment of the ACL graft evaluated with specific MRI slices is halved when growth factors, obtained through a standardized autologous platelet concentrate method, are used.

**DISCUSSION**

For elite athletes, recovery from ACL injury must reach a level close to normal and occur in the shortest time possible so as not to affect the future athletic performance. In the last decade great advances have occurred in ACL reconstruction surgery, considerably improving the outcomes. This is because of the development of more anatomic reconstruction techniques, stronger and more stable methods of attachment over time, accelerated rehabilitation protocols, better technical training, and increased expertise of surgical teams. However, reinjury in these athletes, attributed to trauma in early periods of sports reintegration, clearly indicates that the biological period of maturation and metaplasia needed by the graft used in the reconstruction is not affected by the described advances. For the patellar tendon graft, this period is on average 9 to 12 months. Therefore, during the last years the focus of research has been on advancements in the basic sciences relating to the study of function and capacity for repair, as well as development of growth factors and tissue. Yasuda et al., analyzing the effect of growth factors applied to grafts in dog models, indicated that TGF-β and epidermal growth factor act by increasing the collagen and fibroblast synthesis by 40% in the graft. Anderson et al. indicated that the presence of TGF-β1, TGF-β2, TGF-β3, and TGF-1 growth factors directly influences the graft by improving the scarring rate and increasing the tensile force resistance by 65%. In another interesting experimental study, Weiler et al. indicate that the application of autologous PDGFs applied to the graft during surgery was capable of changing its natural evolution, improving tensile strength and resistance, increasing the maturation rate, and improving collagen quality. The results in our study showed a significant shortening of the biological maturation time of the graft, by at least 48%. Our results show that when PRPG is used, the time required by the graft to achieve complete homogenization, as assessed by MRI, is statistically shortened.
This is very important for the graft’s biological maturation. This means that the graft used with PRPG could undergo its complete process in half the time it naturally requires. We are performing a follow-up of all of our operated athletes to see what happened regarding reinjury, but the follow-up time is still short.

The use of the gelatin (Gelfoam) could affect the magnetic resonance image or analysis, but because it is absorbable, it may already be biodegraded at the time of imaging assessment. There are no studies in the literature describing local changes related to bone or tendon grafts. The early changes seen in the intra-articular segment of the graft with the use of PRPG suggest some effect on it. There are no published studies that relate the quality of the MRI signal intensity with histology or strength of grafts in human models. However, investigations by Weiler et al.12 in sheep models showed that there is a correlation among the homogeneity of the graft on MRI, maturation, and strength, similar to the native ACL.

Much field to cover still remains. The current application of autologous PDGFs22-25 does not allow us to specifically isolate the factors related to the process. We are most likely applying a mixture of factors that apparently do not participate in or influence the healing process of these tissues.4-6,24-26,30,35,36 It is also not clear to us whether isolated application at the time of surgery is enough or whether it would be even more effective to repeat application of these factors during the postoperative recovery and rehabilitation process. Which are the growth factors that are actually needed in ACL reconstruction? Is the quantity we are applying adequate? Is it important to maintain the interaction and balance between all of the growth factors present in the platelet concentrate? How long does their effect last? We still do not have the answers to these questions, and further studies are required.

CONCLUSIONS

ACL reconstruction with the use of PRPG achieves completely homogeneous grafts, assessed by MRI, in 179 days compared with 369 days for ACL reconstruction without PRPG. This represents a time shortening of 48% with respect to ACL reconstruction without PRPG.

REFERENCES


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