Bone Graft Wrapping with Cellulose Polymer Sheet in Posterior Spinal Fusion. A Technical Note

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Abstract

Background: Spinal fusion is one of the most frequent procedures for treating various spinal morbidity. Pseudarthrosis remains a critical complication despite the use of hardware for mechanical stability. The type and proper placement of the bone graft play a fundamental role in achieving solid union. The ideal bone graft material should provide osteogenicity, osteoinductivity, and osteoconductivity, an optimal biological reaction and no risk of transmission of diseases.

Methods: We describe a new technique of bone grafting in two patients who suffered from spinal stenosis. Local bone graft which obtained during decompression of the spine was mixed with bone marrow harvested from the posterior iliac crest. The mixture was wrapped in Surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) and given a cylindrical shape. Finally, the handmade cylinders were placed laterally to the rod of the instrumentation, onto the decorticated transverse processes.

Results: The patients were followed radiographically every three months. The x-rays verified proper placement of the graft onto the transverse processes in both patients. Solid fusion was reported in both sides of the first patient at three months and at six months for the second. At one year postoperatively, fusion status was still graded solid.

Conclusion: The aforementioned technique uses the advantages of a bone auto graft which has been enhanced by bone marrow components, avoiding donor site morbidity. Using Surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) we can adapt the graft to the desired size and shape and finally place it with accuracy onto the decorticated transverse processes. This is a promising technique concerning solid fusion and complications; however, it is a pilot study and needs more time and patients to obtain safe results.

Keywords: Bone autograft; Iliac crest bone graft; Local bone autograft; Postero-lateral spinal fusion

Introduction

Spinal fusion is one of the most frequent procedures for treating various spinal morbidities such as deformity, trauma, and degenerative disc disease with instability.

The biological processes in bone regeneration in spinal fusion procedures require three critical elements: an osteogenic potential that is capable of directly providing cells to the newly forming bone, osteoinductive factors that are able to cause the osteoblastic differentiation of osteoprogenitor stem cells, and osteoconductive scaffold that facilitates neovascularization and supports the in-growth of bone. Furthermore, low-strain mechanical environment must be consistently maintained for the duration of the healing response. Union is achieved when this cellular osteogenic response incorporates and replaces grafted bone with a new matrix that is mechanically rigid in relation to the host bone. The ideal bone graft material possesses all of these three properties along with an optimal biological reaction and without a risk of transmission of diseases.

Iliac crest autograft has been used for many years to achieve lumbar fusion [1-4]. The most frequent complications following fusion surgery are pseudarthrosis and donor site morbidity. Despite the use of rigid instrumentation, the rate of pseudarthrosis remains significant with iliac crest autograft [4-7]. Furthermore, donor site problems, including pain, paresthesias, hematoma and infection, have been reported in up to 50% of patients in some series [8-10].

The use of local autograft as an alternative in lumbar fusion surgery has been evaluated. It achieves a similar fusion rate in 1-level fusion with less morbidity compared to iliac crest. However, it seems that when multiple segments are fused, local bone grafting showed inferior results in fusion compared to iliac crest bone [11].

Major concern in posterolateral fusion still remains the carrier of the bone graft, which should have structural integrity, allowing it to resist compression from the surrounding paraspinal musculature and maintain a space where the posterolateral fusion can form [12-17].

Surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) is an oxidized cellulose polymer made fabric used for haemostasis in various surgeries, including spinal surgery. Recently, animal studies proposed a potential role of oxidized cellulose in new tissue regeneration [18]. Additionally, diced bone grafts wrapped in oxidized cellulose have proven useful in craniomaxillofacial region surgeries, as they combine the advantages of easier adaptation and molding [19].

We describe the details of a surgical technique where we used Surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) as a spacer between the bone graft and the paraspinal musculature.

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USA) to wrap the diced local autograft, which had been previously mixed with bone marrow (4 cc), and place it in the desired position over the transverse processes during instrumented posterolateral fusion of the lumbar spine. We report our primary results in two patients, a 67 and a 68 year old female and male respectively, with instrumented posterolateral fusion for degenerative lumbar spinal stenosis.

Materials and Methods

Case 1

A 67 year old female (S.M.) presented in our outpatient clinic complaining of severe pain to her lower back, continuous for approximately two months and claudication during a walking distance over 200 meters. She had a five year clinical history of lower back pain and she was diagnosed with disk degeneration and grade I spondylolisthesis of L4-L5. Within the last two months she visited the emergency department twice, and she was treated with a combination of anti-inflammatory and muscle relaxant drugs with no relief. She was advised to perform a magnetic resonance imaging (MRI) scan to her lumbar spine, and she was diagnosed with degenerative lumbar spinal stenosis and grade I spondylolisthesis of L4-L5 spine. An instrumented posterolateral fusion and decompression of the lumbar spine from L3 to L5 level was performed.

Case 2

A 68 year old male (M. B), had a previous history of 11 years lower back pain associated with intermittent sciatica pain and claudication. Lately, the patient suffered from increasing numbness in both legs and feet. At clinical examination he was found having impaired reflexes and associated weakness to both legs. A recent electromyogram showed impaired chronic function of L4 and L5 root. Radiographic and MRI scans confirmed the diagnosis of degenerative lumbar spinal stenosis and scoliosis with a left lumbar scoliotic curve of 20 degrees. Wide decompression from L2 to L5 level, instrumented posterolateral fusion from L1 to L5 and correction of the scoliotic curve was performed.

Surgical technique

The local autograft used for grafting was obtained during decompression of the spine. The cephalad and caudal spinous processes were totally ambulated and a wide decompression of the preoperatively marked levels was performed. The harvested pieces of the resected spinous processes, lamina, medial facet and part of the lateral facet as well as osteophytic projections were then cleaned of all soft tissues and bone bits were diced in to small pieces of maximum diameter of 5mm with the usage of a small roger.

The autologous local bone graft pieces were mixed with bone marrow (4 cc) harvested with a syringe from the posterior iliac crest and given a cylindrical shape sized 9 × 2 cm (Figure 1). Then they were wrapped in Surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) original gauze, which was previously cut in dimensions of 10.16 X 10.16 cm. The two edges of the surgicel were tied with an absorbing synthetic suture Vicryl Rapide 3-0, FS-2 (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) (Figure 2). Finally, the two handmade cylinders were placed laterally to the rod of the instrumentation, onto the decorticated transverse processes.

Results

The assessment of fusion was conducted radiographically every 3 months for the first year. Fusion was evaluated by two independent spinal surgeons who where blinded to each other and blinded to the study. Fusion grade was defined as solid fusion, indefinite and non-union based on evidence from plain radiographs.

Fusion was reported as solid in both sides of the first patient at three months postoperatively. Fusion in the second patient was considered as indefinite at three months (Figure 3A & 3B). However, at six months postoperatively there was evidence of solid fusion in both patients. At one year after surgery, fusion status was still graded solid in both patients.

Discussion

Obtaining a solid arthrodesis is the main objective in lumbar surgical procedures. Successful fusion depends on a number of surgical and host factors including the selection of a bone graft or bone graft substitute with adequate osteoconductive and osteoinductive properties.

Spine surgeons use instrumentation to maintain low-strain mechanical environment for the duration of the healing response. Iliac crest bone graft remains the standard to which bone graft substitutes and bone graft enhancers are compared, combining osteogenic, osteoinductive and osteoconductive properties. Nevertheless, published fusion rates for posterolateral fusion using iliac crest bone graft vary...
from 40% to 80% [20,21]. Furthermore, a number of complications associated with iliac crest bone harvest have been widely reported [9,22-27].

Due to these drawbacks a variety of bone grafts have been developed for use in posterolateral fusion in lumbar spine.

Allograft bone is a widely used grafting material, commonly regarded as a bone graft extender. Allografts have an osteoconductive scaffold with minimal osteoinductive factors; however, they are not able to provide osteogenic cells because of the processing that they undergo in order to decrease their antigenicity. Compared to autografts, the allografts are incorporated slower and less completely with decreased vascularization and osteoconduction [28]. Beyond the question of efficacy, the primary concern surrounding the use of allograft bone is the risk for disease transmission.

Demineralized bone matrix (DBM) is a class of bone graft materials made by processing allograft bone. Demineralization process leaves behind the small quantities of bone morphogenetic protein (BMP) found in allograft bone, and thus generates osteoinductive potential. However, DPMs lack structural strength and also there is concern about their osteogenic potential [29,30].

Cell-based strategies describes a variety of techniques by which cells with osteogenic potential are collected, concentrated, and applied as bone graft enhancers [31-34]. The most common target cells are bone marrow components and platelets. Bone marrow components have promising results in spinal fusion as is a well established source of mesenchymal stem cells (MSCs). There are animal models [35] and clinical studies [36] that have demonstrated efficacy in spinal fusion using bone marrow-derived mesenchymal stem cells (MSCs). On the other hand there is data that questions the efficacy of platelets in posterolateral fusion [37-39]. A drawback in the use of cell-based strategies is that these materials totally lack structural strength and can only be used as bone graft enhancers.

BMPs are members of the transforming growth factor-beta superfamily. By binding to specific receptors present on the surface of the osteogenic progenitor, intracellular cascades – which resemble endochondral ossification – are activated. Although, they have the greatest osteogenic potential, they lack osteoconductivity, diffuse away from the fusion site easily, and become inactivated in vivo when they used alone. Therefore, recombinant BMPs are combined with a carrier matrix that serves to retain the concentration and releases them consistently over time. Drawbacks to their use are complications such as, osteolysis, swelling/edema, heterotopic bone formation, antibody reaction, and their high cost.

Ceramic scaffolds (calcium phosphates) and a variety of new agents as calcium sulfate, bioactive glass, dual hydroxyapatite composite with porous and solid parts, poly and highly porous hydroxyapatite, have been tested in animal studies and have demonstrated to possess an osteoconductive ability [37,40-45]. Nevertheless, they lack osteogenic and osteoinductive activity, thus becoming effective only as vehicles of other bone grafts.

Local autografts can be used as an alternative to iliac crest autograft. They combine osteogenic, osteoinductive and osteoconductive properties, avoiding morbidity associated with the harvesting of iliac crest bone autograft. Concerns may be raised about the sufficient quantity [11] and the placement of the local autograft in cases of posterolateral fusion. Cammisa et al. [46] suggested that autogenous bone graft volumes of 10cc or less, in one-level fusion, were of limited value.

In our proposed technique the local bone graft, coming from spinous processes, lamina, posterior and medial portions of facet, as well as osteophytic projections, was cleaned of all soft tissues and bone bits were diced in to small pieces. Local autograft was then mixed with bone marrow (4cc) harvested with a syringe from both posterior iliac crests. The autologous bone graft pieces were given a cylindrical shape and they were wrapped in surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) original gauze. The use of surgicel (Ethicon, Johnson & Johnson Medical Ltd, Somerville, NJ, USA) aimed in taking advantage of the features of easy adaptation and molding [19], optimizing the connective tissue formation and giving support to the regenerating tissue [18].

The first results, using this technique, are promising as far as concern solid fusion and complications. However, this is a pilot study which presents a new technique of collection, preparation, and placement of local bone autograft mixed with bone marrow in posterolateral spine fusion, and needs more time and patients to obtain safe results.

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