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Clinical and radiographic outcomes of extreme lateral approach to interbody fusion with β-tricalcium phosphate and hydroxyapatite composite for lumbar degenerative conditions

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Abstract

Background: Historically, iliac crest bone graft has been used as the graft of choice for lumbar spine fusion procedures. Because fusion techniques have become less invasive, the demand for minimally disruptive grafting options has increased. This prospective study was performed to assess clinical and radiographic outcomes of patients treated with an iliac crest bone graft alternative and lateral lumbar interbody fusion.

Methods: Fifty degenerative lumbar patients were treated with the extreme lateral approach to interbody fusion and a β-tricalcium phosphate–hydroxyapatite graft with bone marrow aspirate (BMA) at 1 or 2 adjacent levels. BMA was collected from the iliac crest with a bone aspiration needle and applied to the FormaGraft (NuVasive, Inc., San Diego, California) in a 1:1 ratio. Radiolucent cages were filled with FormaGraft strips, granules, or blocks and implanted in a standard fashion. Clinical data were collected at baseline, 6 weeks, and 3, 6, and 12 months and included visual analog scale, Oswestry Disability Index, and Short Form 36 patient-reported assessments. Fusion assessments were made from neutral anteroposterior/lateral radiographs, lateral flexion/extension radiographs, and computed tomography images taken at least 12 months after surgery.

Results: Forty-four patients treated at 49 levels completed follow-up. The mean patient age was 54.7 ± 10.8 years, and mean body mass index was 30.8 ± 7.7 kg/m². Radiographic fusion was observed in 41 of 44 assessed levels (93.2%). Blood loss was less than 100 mL in 95.5% of patients. Of the patients, 93.2% spent 1 night or less in the hospital. By the 6-week follow-up, all clinical outcomes were significantly improved (P < .05). Improvements were maintained or increased throughout the course of follow-up.

Conclusions: This report shows that the technique of extreme lateral approach to interbody fusion in combination with FormaGraft and BMA in the interbody space is a safe and effective treatment option for interbody fusion of the lumbar spine when compared with other approaches and biologic options.

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Keywords: Obese; Spine; Lumbar fusion; β-Tricalcium phosphate; Synthetic; Graft; BMA

The clinical and radiographic outcomes after a lumbar interbody fusion procedure are influenced by a number of factors, including surgical approach, interbody implant, internal fixation, and application of biologics.

The extreme lateral approach to interbody fusion (XLIF) (NuVasive, Inc., San Diego, California) is a minimally disruptive technique for fusion of the anterior spine. The technique was first described in 2006¹ and has gained widespread acceptance within the spine community.

With decreased morbidity at the surgical site, there has been an increased demand for minimally disruptive grafting options as well. Traditionally, interbody fusion procedures have included the use of iliac crest bone graft (ICBG) to increase fusion success and healing. However, concerns about graft-site morbidity and resultant changes in clinical and financial outcomes have fed a demand for alternatives. Bone marrow aspirate (BMA) has the same osteogenic qualities as ICBG and can be collected from either the vertebral bodies or the iliac crest with a thin aspirate needle. The combination of BMA with osteoconductive materials such as hydroxyapatite (HAp) and β-tricalcium phosphate (β-TCP) can theoretically provide a minimally disruptive alternative to ICBG. HAp and β-TCP graft alternatives have been studied in a number of applications and have been
shown to perform as well as or better than iliac crest autograft in comparative studies,\textsuperscript{2–4} observational studies,\textsuperscript{5–8} and a number of animal studies.\textsuperscript{9,10}

**Methods**

**Study design**

The use of a $\beta$-TCP–HAp composite (FormaGraft; NuVasive, Inc.) with BMA as a substitute for iliac crest bone autograft was prospectively studied in 54 nonrandomized patients undergoing the XLIF procedure at 1 or 2 levels between February and November 2008. The study was approved and overseen by an institutional review board. Before enrollment, all patients gave their written consent for participation.

Study participation was limited to patients aged between 18 and 70 years with a primary diagnosis of internal disc disruption at 1 or 2 contiguous lumbar levels between L1 and L5 and persistent back or leg pain for at least 6 weeks. Patients were excluded if they had more than 1 previous failed fusion or active or latent infection of the disc or spine.

**Surgical protocol**

The procedure was performed by standard surgical technique as previously described,\textsuperscript{1} with optional internal fixation. BMA was collected from the iliac crest with a bone aspiration needle during the primary procedure. A maximum of 2 mL of BMA was collected per aspiration site, for a total of 15 to 20 mL on average. The FormaGraft was prepared by applying BMA to the graft in a 1:1 ratio by volume. Radiolucent cages were filled with synthetic graft strips, granules, or blocks and implanted in a standard fashion.

**Outcome measures**

Baseline demographics were collected at the preoperative visit. Surgical outcomes, including estimated blood loss, surgical time, method of graft application, and intraoperative complications, were captured during surgery.

Clinical outcome measures were collected at each visit and included complication assessments, work status, and visual analog scale (VAS), Oswestry Disability Index (ODI), and Short Form 36 patient-reported assessments. Outcome measures were completed by patients preoperatively and postoperatively at 6 weeks and 3, 6, and 12 months.

Anteroposterior and neutral lateral radiographs were collected at every visit. Flexion and extension lateral radiographs were collected preoperatively and postoperatively at 3, 6, and 12 months. Computed tomography (CT) scans were performed after the 12-month visit to assess fusion. Radiographic assessment of interbody fusion was conducted by an independent board-certified orthopedic surgeon using a modification of the Lenke grading system,\textsuperscript{11} where grade I and grade II levels were classified as fused and grade III levels were classified as unfused. In addition to the Lenke scale, a secondary scale was used to describe the volume of bone present in the disc space. The Lenke scale and volume scales are defined in Table 1.

**Results**

**Patient demographics**

A total of 55 levels were treated in 50 patients at a single center. Two patients died of causes unrelated to their spine condition or procedure, and 4 patients were lost to follow-up. At the end of the study, 44 patients (31 women and 13 men) had completed a clinical evaluation and/or radiographic evaluation at least 12 months after surgery: 36 had both clinical and radiographic records, 5 had CT scans without 12-month clinical outcomes, and 3 had 12-month clinical outcomes without radiographs. Baseline and procedure data are reported for all 44 patients, clinical data are limited to those patients who completed a 12-month clinical assessment, and the fusion assessment is limited to those patients with CT evaluations at least 12 months after surgery.

At baseline, the mean patient age was 54.7 ± 10.8 years (range, 24–70 years) and mean body mass index was 30.8 ± 7.7 kg/m\(^2\). Significant comorbidities included smoking (n = 21, 42.0%), obesity (n = 24, 48.0%), and diabetes mellitus (n = 4, 8.0%). Before surgery, 13 patients (26.0%) were working, 17 (34.0%) were retired voluntarily, 6 (12.0%) were retired because of illness, 7 (14.0%) were medically disabled, and 2 (4.0%) were homemakers.

Indications for surgery included 1 or more of the following: spondylolisthesis (n = 26, 52.0%), degenerative disc disease (n = 22, 44.0%), stenosis (n = 22, 44.0%), lateral listhesis (n = 8, 16.0%), adjacent segment disease (n = 7, 14.0%), instability (n = 7, 14.0%), scoliosis (n = 7, 14.0%), herniated nucleus pulposus (n = 4, 8.0%), and post-laminectomy instability (n = 2, 4.0%). On average, patients were treated at 1.1 levels. L4-5 was included in 28 procedures (63.6%) and accounted for over half of levels treated. L1-2 was included in 2 procedures (4.5%), L2-3 in 5

| Table 1 | Radiographic evaluation scales |
|---------|--------------------------------|
| **Lenke scale** | Description |
| Grade I | Fused with remodeling and trabeculae |
| Grade II | Graft intact, not fully remodeled and incorporated throughout; no lucencies |
| Grade III | Graft intact but definite lucency at top or bottom of graft |
| Grade IV | Definitely not fused with resorption of bone graft and with collapse |

| Volume scale | Description |
|-------------|-------------|
| Grade I | 0%–25% of available space occupied by bone |
| Grade II | 26%–50% of available space occupied by bone |
| Grade III | >50% of available space occupied by bone |
(11.4%), and L3-4 in 14 (31.8%). Six patients were also treated at L5-S1 with a trans-sacral approach (TranS1 Inc., Wilmington, North Carolina); these levels are not included in the fusion results of this study.

Operative data

Intraoperatively, 22 patients (50.0%) lost less than 50 mL of blood, 20 (45.5%) lost between 50 and 100 mL, and 2 (4.5%) lost greater than 100 mL. The operative time for patients ranged from 40 to 138 minutes (mean, 66.0 ± 27.4 minutes). All levels were treated with polyetheretherketone cages in the interdisc space. Supplemental posterior internal fixation included unilateral posterior pedicle screw fixation (n = 26, 59.1%) or bilateral posterior pedicle screw fixation (n = 7, 15.9%). In 11 patients (25%) posterior pedicle screws were not used; rather, they were treated with supplemental lateral fixation.

A combination of β-TCP–HAp strips, granules, and blocks was used with BMA; patients were treated with strips alone (n = 22, 50.0%), both strips and granules (n = 20, 45.5%), or blocks alone (n = 2, 4.5%). Length of hospitalization ranged from 0 to 4 days, with 90.7% of patients spending 1 night or less in the hospital. There were no complications noted during surgery or hospital stay.

Patient-reported clinical outcomes

Mean baseline scores were as follows: Visual Analogue Scale (VAS) for back pain, 8.2 ± 1.8 cm; VAS for leg pain, 7.9 ± 2.0 cm; and ODI, 50.9% ± 15.2%. By the 6-week follow-up, all clinical outcomes were significantly improved (P < .05). Improvements were maintained or increased throughout the course of follow-up. At 12 months, the mean VAS score for back pain was 4.8 ± 3.1 cm and the mean VAS score for leg pain was 3.7 ± 3.1 cm. The mean ODI was 33.1% ± 19.6%, and the mean Short Form 36 score was 53.3 ± 19.5 (Figs. 1 and 2).

Early complications—defined as those that occurred up to 6 weeks after surgery—included 1 readmission to the hospital for pain and swelling within 24 hours of discharge, as well as 1 revision for residual stenosis. Late complications—defined as those that occurred more than 6 weeks after surgery—included 1 incident of residual stenosis at the operative level that required surgical intervention, as well as 1 patient who underwent excision of a subcutaneous lesion believed to be residual graft material 12 months after the index procedure, performed by another surgeon. There were no complications reported related to BMA collection or implant subsidence.

Radiographic results

CT scans used to assess fusion were collected a mean of 17.3 months (range, 12–31 months) after surgery. At the time of radiographic review, 41 of the 44 levels assessed (93.2%) were fused (Fig. 3). A modification of the Lenke interbody fusion scores yielded the following: grade I, 68.1% (n = 30); grade II, 25.0% (n = 11); grade III, 6.8% (n = 3); and grade IV, 0% (n = 0). Volume assessments were as follows: grade I, 0% (n = 0); grade II, 6.8% (n = 3); and grade III, 93.2% (n = 41).
Discussion

All patients in this series were treated with BMA from the iliac crest. Although this application requires multiple aspiration sites in the iliac crest, harvesting BMA is quicker and far less invasive than removing ICBG. None of the patients in this series had any complications at the site of BMA collection, and none required revisions related to graft subsidence or nonunion.

In 2009 Rihn et al. reported on a cohort of 130 Transforaminal Lumbar Interbody Fusion (TLIF) patients treated with either recombinant BMP or ICBG. Despite reasonable fusion success (96.4% in cohort with no significant difference between groups), both patient groups had considerable complications, many of which were directly attributable to the graft used. In the recombinant human BMP group, 5.8% of patients had symptomatic vertebral osteolysis and 2.3% had ectopic bone formation; 30.3% of ICBG patients had documented donor-site morbidity. The only graft-related complication in this cohort is presumed presence of residual subcutaneous graft material—at the time of excision, no ectopic ossification was reported by the operating surgeon.

In our study successful fusion was achieved in 93.2% of levels, with abundant new bone growth. There were no revisions for segmental nonunions in this cohort. Recently reported lumbar fusion rates have ranged from 91% to 100% for XLIF and 90% to 100% for posterior lumbar fusion (PLIF)/TLIF with no apparent difference between minimally invasive and traditional open techniques.

In 2010 Schuster et al. reported on infection and concluded that case-specific factors including blood loss and case length influence the incidence of postoperative infection. In 2009 Rihn et al. reported the combined autograft harvest site/lumbar wound infection rate of single-level TLIF with ICBG at 9.2%. In this series, there was 1 report of wound swelling that did not require surgical treatment and there was no incidence of postoperative infection. Through the use of a minimal incision and the elimination of an additional surgical site for ICBG harvesting, the lateral approach with β-TCP-HAp may have effectively reduced the risk of infection by reducing surgical times and blood loss.

Conclusion

This report shows that the XLIF technique in combination with FormaGraft and BMA in the interbody space is a safe and effective treatment option for interbody fusion of the lumbar spine when compared with reports of other approaches and biologic options.

References

1. Ozgun BM, Aryan HE, Pimenta L, Taylor WR. Extreme lateral interbody fusion (XLIF): a novel surgical technique for anterior lumbar interbody fusion. Spine J 2006;6:435–43.
2. Bansal S, Chauhan V, Sharma S, Maheshwari R, Juyal A, Raghuvanshi S. Evaluation of hydroxyapatite and beta-tricalcium phosphate mixed with bone marrow aspirate as a bone graft substitute for posterolateral spinal fusion. Indian J Orthop 2009;43:234–9.
3. Dai LY, Jiang LS. Single-level instrumented posterolateral fusion of lumbar spine with beta-tricalcium phosphate versus autograft: a prospective, randomized study with 3-year follow-up. Spine (Phila Pa 1976) 2008;33:1299–304.
4. Lerner T, Bullmann V, Schulte TL, Schneider M, Liljenqvist U. A level-1 pilot study to evaluate of ultraporous beta-tricalcium phosphate as a graft extender in the posterior correction of adolescent idiopathic scoliosis. Eur Spine J 2009;18:170–9.
5. Dai LY, Jiang LS. Anterior cervical fusion with interbody cage containing beta-tricalcium phosphate augmented with plate fixation: a prospective randomized study with 2-year follow-up. Eur Spine J 2008;17:698–705.
6. Epstein NE. A preliminary study of the efficacy of Beta tricalcium phosphate as a bone expander for instrumented posterolateral lumbar fusions. J Spinal Disord Tech 2006;19:424–9.
7. Epstein NE. An analysis of noninstrumented posterolateral lumbar fusions performed in predominantly geriatric patients using lamina autograft and beta tricalcium phosphate. Spine J 2008;8:882–7.
8. Epstein NE. Fusion rates and SF-36 outcomes after multilevel laminectomy and noninstrumented lumbar fusions in a predominantly geriatric population. J Spinal Disord Tech 2008;21:159–64.
9. Becker S, Maissen O, Ponomarev I, et al. Osteopromotion with a plasmagmanumisinate on a beta-TCP ceramic. J Mater Sci Mater Med 2008;19:659–65.
10. Walsh WR, Vizesi F, Cornwall GB, Bell D, Olivier R, Yu Y. Posterolateral spinal fusion in a rabbit model using a collagen-mineral composite bone graft substitute. Eur Spine J 2009;18:1610–20.
11. Bridwell KH, Lenke G, McEnery KW, Baldus C, Blanke K. Anterior fresh frozen structural allografts in the thoracic and lumbar spine. Do they work if combined with posterior fusion and instrumentation in adult patients with kyphosis or anterior column defects? Spine (Phila Pa 1976) 1995;20:1410–8.
12. Mroz TE, Wang JC, Hashimoto R, et al. Complications related to osteobiologics use in spine surgery: a systematic review. Spine (Phila Pa 1976) 2010;35:S86–104.
13. Rihn JA, Patel R, Makda J, et al. Complications associated with single-level transforaminal lumbar interbody fusion. Spine J 2009;9:623–9.
14. Burak MO, Vijay A, Erin N, et al. Two-year clinical and radiographic success of minimally invasive lateral transposa approach for the treatment of degenerative lumbar conditions. SAS J 2010;4:41–6.
15. Oliveira L, Murchi L, Coutinho E, et al. The use of rh-BMP2 in standalone extreme lateral interbody fusion (XLIF): clinical and radiological results after 24 months follow-up. World Spinal Column J 2010;1:19–25.
16. Rodgers WB, Cox CS, Gerber EJ. Early complications of extreme lateral interbody fusion in the obese. *J Spinal Disord Tech* 2010;23:393–7.

17. Rodgers WB, Gerber EJ, Patterson JR. Fusion after minimally disruptive ALIF: analysis of XLIF by computed tomography. *Spine Arthroplasty Soc J* 2010;4:37–40.

18. Dhall SS, Wang MY, Mummaneni PV. Clinical and radiographic comparison of mini-open transforaminal lumbar interbody fusion with open transforaminal lumbar interbody fusion in 42 patients with long-term follow-up. *J Neurosurg Spine* 2008;9:560–5.

19. Gill K, Blumenthal SL. Posterior lumbar interbody fusion. A 2-year follow-up of 238 patients. *Acta Orthop Scand Suppl* 1993;251:108–10.

20. Hackenberg L, Halm H, Bullmann V, Vieth V, Schneider M, Liljenqvist U. Transforaminal lumbar interbody fusion: a safe technique with satisfactory three to five year results. *Eur Spine J* 2005;14:551–8.

21. Lauber S, Schulte TL, Liljenqvist U, et al. Clinical and radiologic 2–4-year results of transforaminal lumbar interbody fusion in degenerative and isthmic spondylolisthesis grades 1 and 2. *Spine (Phila Pa 1976)* 2006;31:1693–8.

22. Schuster JM, Rechtine G, Norvell DC, et al. The influence of perioperative risk factors and therapeutic interventions on infection rates after spine surgery: a systematic review. *Spine (Phila Pa 1976)* 2010;35:S125–37.