Percutaneous vertebroplasty for vertebral compression fractures with and without intravertebral clefts

K.-Y. Ha, J.-S. Lee, K.-W. Kim, J.-S. Chon

From The Catholic University and Dongshin General Hospital, Seoul, Korea

We present the clinical and radiological results of percutaneous vertebroplasty in the treatment of 58 vertebral compression fractures in 51 patients at a minimum follow-up of two years. Group 1 consisted of 39 patients, in whom there was no associated intravertebral cleft, whilst group 2 comprised 12 patients with an intravertebral cleft. The Oswestry disability index (ODI) and visual analogue scale (VAS) scores were recorded prospectively. The radiological evidence of kyphotic deformity, vertebral height, leakage of cement and bone resorption around the cement were studied retrospectively, both before and after operation and at the final follow-up.

The ODI and VAS scores in both groups decreased after treatment, but the mean score in group 2 was higher than that in group 1 (p = 0.02 (ODI), p = 0.02 (VAS)). There was a greater initial correction of the kyphosis in group 2 than in group 1, although the difference was not statistically significant. However, loss of correction was greater in group 2. Leakage of cement was seen in 24 (41.4%) of 58 vertebrae (group 1, 32.6% (15 of 46); group 2, 75% (9 of 12)), mainly of type B through the basal vertebral vein in group 1 and of type C through the cortical defect in group 2. Resorption of bone around the cement was seen in three vertebrae in group 2 and in one in group 1. There were seven adjacent vertebral fractures in group 1 and one in group 2.

Percutaneous vertebroplasty is an effective treatment for osteoporotic compression fractures with or without an intravertebral cleft. Nonetheless, higher rates of complications related to the cement must be recognised in patients in the presence of an intravertebral cleft.

Percutaneous vertebroplasty was introduced by Galibert et al1 and has been used to treat patients with osteoporotic vertebral compression fractures, vertebral metastatic cancer, myeloma and haemangioma, all of which cause severe pain. A success rate of 90% to 95% has been claimed for managing osteoporotic vertebral compression fractures with this treatment.2,3 Percutaneous vertebroplasty has also been used in Kummell’s disease, in which an intravertebral cleft occurs in the presence of an osteoporotic compression fracture. Jang, Kim and Lee4 and Lane et al5 reported excellent results in this situation. The intravertebral clefts, first described by Maldague, Noel and Malghem,6 are considered to represent avascular necrosis of the vertebral body and are not thought to be associated with acute osteoporotic compression fractures.7 Kummell’s disease causes rapid collapse of the vertebra, loss of lordosis due to necrosis of the anterior vertebral body and fibrosis around the bone necrosis.7 As a result, there is a large void in the anterosuperior portion of the vertebral body. Lane et al5 and McKiernan and Faciszewski8 found that the filling patterns of polymethylmethacrylate (PMMA) in percutaneous vertebroplasty were different in cleft vertebrae and uncleft vertebrae.

We have compared the clinical and radiological results of percutaneous vertebroplasty in patients with osteoporotic vertebral compression fractures with or without an intravertebral cleft.

Patients and Methods

Between July 2000 and October 2002 we treated 54 patients with persistent pain due to osteoporotic vertebral compression fractures by percutaneous vertebroplasty. Two were lost to follow-up and one died, leaving 51 consecutive patients with 58 fractures who were followed up for more than two years. They were separated into two groups. Group 1 consisted of 39 patients with 46 involved vertebrae, without an associated intravertebral cleft and
group 2 comprised 12 patients with 12 vertebral compression fractures, with an intravertebral cleft. The male-to-female ratio in group 1 and group 2 was 6:33 and 1:11, respectively. The mean age for all patients at the time of treatment was 70.9 years (64 to 79), with a mean age of 71.1 years for group 1 and 62.5 years for group 2.

Of the 46 vertebral compression fractures in group 1, 32 were at T12 to L2, compared with nine at the same sites in group 2 (Fig. 1). The fractures selected for percutaneous vertebroplasty were determined from plain radiographs, bone scans, MRI and physical examination for the location of pain in patients who had no improvement in pain after three weeks of conservative treatment. From the bone scan, fracture segments showing increased uptake of isotope were selected. All patients underwent MRI and the vertebrae with intravertebral fluid or air shadows in the sagittal plane were selected and their location matched to the site of pain, in order to determine which segment should be treated. If severe vertebral compression fractures seen on plain radiographs did not show increased uptake of isotope, they were thought to represent old injuries.

Percutaneous vertebroplasty was undertaken under local anaesthesia, with the spine extended as described by Jensen et al.\(^3\) A bone-biopsy needle was inserted into the anterior third of the vertebra, through a transpedicular approach, to inject the liquid bone cement under radiological control using an image intensifier. Injection continued until the vertebra was opacified, or leakage into the paraspinal veins was seen. After finishing the injection, the same procedure was repeated through the opposite pedicle. PMMA cement (Depuy International Ltd, Leeds, United Kingdom) mixed with barium sulphate contrast was used in all patients. The mean quantity of bone cement in groups 1 and 2 was 6.2 ml and 5.4 ml, respectively. Using dual-energy x-ray absorptiometry (Hologic QDR-4500A, Bedford, Massachusetts), the bone mineral density (BMD) of the lumbar spine and proximal femur was checked in all patients who underwent percutaneous vertebroplasty. The T-score for the BMD was less than -2.5 in all patients, confirming the diagnosis of osteoporosis. The mean follow-up was for 26.3 months (24 to 36).

In all patients, plain radiographs were obtained on the day immediately before the percutaneous vertebroplasty, one day after treatment and at the final follow-up. CT was also performed one to three days after treatment. The kyphotic angle, the vertebral height, resorption of bone around the injected cement and fractures of the adjacent vertebrae were assessed from the plain films (Fig. 2). The vertebral height was measured using the method of McKiernan, Faciszewski and Jensen.\(^9\) For pre- and post-operative comparison and for the final follow-up, we calculated the ratio of the height of the anterior and posterior aspects of the vertebral body and that of the middle and the posterior vertebral height.

Based on the post-operative CT scans, leakage of any cement was classified into three types: type B (leakage through the basal vertebral vein), type S (leakage through the segmental vein) and type C (leakage through the site of cortical bone loss).\(^10,11\) A visual analogue scale for pain (VAS) and the Oswestry Disability Index (ODI)\(^12\) were
applied pre-operatively, three months after treatment and at the final follow-up, to compare the severity of pain and its influence on daily life.

**Statistical analysis.** The paired t-test, Fisher’s exact test and the Pearson chi-squared test were used for statistical analysis. The statistical significance before and after the operation and at the final follow-up was tested by repeated-measures analysis of variance (ANOVA). The results were considered to be significant if p < 0.05.

**Results**

**Clinical findings.** In group 1, the mean pre-operative ODI score of 75.1 had decreased to 38.7 by three months after the operation. The difference was statistically significant (repeated-measures ANOVA, p < 0.0001). The mean score at the final follow-up was 37.3, but this further change was not statistically significant.

In group 2, the mean pre-operative ODI score of 79.9 had decreased to 48.5 by three months after the operation. The difference was statistically significant (repeated-measures ANOVA, p < 0.0001). The mean score at the final follow-up had increased to 55.4, but this difference was not statistically significant (Fig. 3).

In group 1, the mean pre-operative VAS score was 8.3 and had decreased to 4.2 after the operation. The difference was statistically significant (repeated-measures ANOVA, p < 0.0001). The mean score at the final follow-up was 3.5, but this change was not statistically significant.

In group 2, the mean pre-operative VAS score was 9.4, which decreased to 4.5 after the operation. The difference was statistically significant (repeated-measures ANOVA, p < 0.0001). The mean score at the final follow-up was 5.9, which was not statistically significant from the result at three months (Fig. 4).

At the 5% level of significance, the ODI and VAS of group 2 were significantly higher than those of group 1 (repeated-measures ANOVA, p = 0.02).

**Radiological findings.** In group 1, the mean kyphosis measured 12.0° (SD 5.8) pre-operatively, 11.2° (SD 5.5) post-operatively, and 11.2° (SD 5.6) at the final follow-up, representing a correction of 6.7%. Group 2 had a mean kyphosis of 15.5° (SD 6.9) pre-operatively, 12.0° (SD 4.0) post-operatively, and 12.7° (SD 4.9) at the final follow-up. A correction of 18.1% was achieved, but this was not fully maintained. The difference between the two groups was not statistically significant.

In group 1, the mean pre-operative anterior and posterior vertebral height ratio measured 70% (SD 1.5), the post-operative ratio was 72% (SD 2.2) and the ratio was 71.6% (SD 3.7) at the final follow-up. By contrast, in group 2, the mean pre-operative ratio was 60% (SD 3.2), the post-operative ratio 66% (SD 4.8) and at final follow-up, was 64.4% (SD 4.7). Group 2 had a larger increase in anterior height than group 1, although the difference was not statistically significant.

In group 1, the mean pre-operative middle and posterior vertebral height ratio measured 60% (SD 3.8). Post-operatively it was 68% (SD 2.6) and at final follow-up, 67.6% (SD 1.4). This compared with group 2, which had a mean pre-operative ratio of 55% (SD 3.1), a post-operative ratio of 59% (SD 5.4) and a ratio of 58% (SD 3.5) at final follow-up. Group 2 showed a larger increase in middle height than group 1, although the difference was not statistically significant.

**Complications.** Those complications directly related to the percutaneous vertebroplasty occurred in 31 of 58 vertebrae (53.4%), but there were no neurological symptoms caused by the treatment and no infections. The Pearson chi-
squared test was used for the statistical analysis of complications.

Leakage of bone cement was observed in 24 of 58 vertebrae (41.4%). In group 1, leakage was seen in 15 of 46 vertebrae (32.6%) and in 13 of these was classified as type B. In group 2, leakage was seen in nine of the 12 vertebrae (75%) and seven were classified as type C. More leakage was observed in group 2 than in group 1 (p < 0.05).

Bone resorption around the cement was observed in one of 46 vertebrae (2.2%) in group 1 and three of 12 (25%) in group 2 (Fig. 5). This difference was significant (p = 0.023).

Fractures adjacent to the lesion occurred in five of the 51 patients (9.8%). There were seven in four patients in group 1 and one in group 2. There was no statistical significance between the groups.

**Discussion**

Studies on vertebroplasty have focused mainly on its advantages, the method of operation or the results of its application to osteoporotic compression fractures. Our study compared the results of percutaneous vertebroplasty in vertebral compression fractures with or without intravertebral clefts at a minimum follow-up of two years.

The ODI and VAS scores decreased in both groups after treatment. However, group 2 showed higher scores than group 1, suggesting that patients with an intravertebral cleft experienced less reduction of pain and more inconvenience in daily life after the operation than those without a cleft. Nevertheless, percutaneous vertebroplasty provided satisfactory results in these patients. The ODI and VAS scores at the final follow-up, when compared with those immediately after operation, showed a slight decrease in patients with intravertebral clefts, whereas they increased slightly in those without a cleft (group 1). However, the difference between patients with and without a cleft was not statistically significant. This suggested that the treatment provided a comparatively well-maintained improvement.

In group 2, four of nine vertebrae with leakage of cement, three with bone resorption around the injected cement and one with a fracture in the adjacent vertebra showed an increase in the ODI and VAS scores at the final follow-up compared with those after three months. Similarly, in group 1, ten of 15 vertebrae with leakage of cement, one with bone resorption around the injected cement and seven with fractures at the adjacent vertebrae showed an increase in the ODI and VAS scores at the final follow-up, compared with those after three months.

Our patients were representative of those with vertebral compression fractures in general, with most involved vertebrae being located at the thoracolumbar junction. The extent of the correction of the kyphosis in our series was consistent with that of Michael et al. We found that the correction was partially lost at the final follow-up, in particular in patients with an intravertebral cleft. This agrees with the results of McKiernan, Jensen and Faciszewski, who reported that posture had a greater influence on the extent of the kyphosis in patients with a cleft than in those without. McKiernan et al compared the pre-operative plain radiographs taken in the upright position with those taken prone, noting that the correction of the kyphotic deformity seemed to be more significant in the latter position. However, in our study all the radiographs were obtained with the patients standing. Of note was the finding that the ratio of vertebral height increased in both groups after operation but had decreased by the final follow-up.

Leakage of bone cement from the vertebra on to the dura mater has been reported in 30% to 70% of percutaneous vertebroplasties. Leakage of bone cement was observed in 24 of 58 vertebrae (41.4%). In group 1, leakage was seen in 15 of 46 vertebrae (32.6%) and in 13 of these was classified as type B. In group 2, leakage was seen in nine of the 12 vertebrae (75%) and seven were classified as type C. More leakage was observed in group 2 than in group 1 (p < 0.05).

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the presence of a cleft and was consistent with the findings of Yeom et al., who found that in the absence of a cleft, the leakage of bone cement was mainly of type B. However, they reported that type B leakage made up about 40% of the total, which was much lower than our incidence of 86.7%.

Bone resorption around the injected bone cement was found more often in vertebrae with a cleft than in those without. McKiernan and Faciszewski reported the patterns of opacification after percutaneous vertebroplasty and found that persistently mobile, clefted vertebrae filled as a confluent reservoir for cement, with uniform opacity and sharp radiological margins. In the absence of a cleft, the cement was interspersed throughout the trabecular space in a more even manner. We found that in vertebral compression fractures with an intravertebral cleft, the cement filled a large gap in the anterosuperior region of the vertebral body. Therefore, the load on the vertebra was thought to be transferred to the bone cement, which was harder than the surrounding bone, thus creating a stress-shielding effect. This possibly caused more bone resorption in patients with a cleft than in those without.

Our results indicate that percutaneous vertebroplasty can be used to treat osteoporotic vertebral compression fractures with or without an intravertebral cleft. However, there is a greater likelihood of complications if an intravertebral cleft is present.

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No benefits in any form have been received or will be received from a commercial party related directly or indirectly to the subject of this article.

References