Clinical Observation of Two Bone Cement Distribution Modes After Percutaneous Vertebroplasty for Osteoporotic Vertebral Compression Fractures

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Research Article

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Abstract

Background: Current findings suggest that percutaneous vertebroplasty (PVP) is a suitable therapeutic approach for osteoporotic vertebral compression fractures (OVCFs). The present retrospective study aimed to investigate the differences in clinical efficacy and related complications between the two bone cement distribution modes.

Methods: We retrospectively reviewed the medical records of the patients with single-segment OVCFs who underwent bilateral percutaneous vertebroplasty. Patients were divided into blocky and spongy group according to the type of postoperative bone cement distribution. Clinical efficacy and related complications was compared between the two bone cement distribution modes on 24h after the operation and last follow-up.

RESULTS: The mean follow-up time was 17.54 months. The VAS and ODI after operation improved significantly in both two groups. The VAS and ODI in the spongy group was significantly lower than that in the blocky group, 24h postoperatively, and at the last follow-up. There were 42 cases (12.8%) of adjacent vertebral fractures, 26 cases (19.8%) in the blocky group and 16 cases (8.1%) in the spongy group. There were 57 cases (17.3%) of bone cement leakage, 18 cases (13.7%) in blocky group and 39 cases (19.7%) in the spongy group. At 24 hour postoperatively and at the last follow-up, local kyphosis and anterior vertebral height were significantly corrected in both groups, but gradually decreased over time, and the degree of correction was significantly higher in the spongy group than in the block group. Loss of local kyphosis and loss of vertebral body height were also less severe in the spongy group at the last follow-up.

Conclusions: Compared with blocky group, spongy group can better maintain the height of the vertebral body, correct local kyphosis, reduce the risk of the vertebral body recompression, long-term pain and restore functions.

Introduction

With the aging of the social population, incidence of osteoporosis is constantly increasing seriously affecting life quality of elderly patients[1]. Osteoporotic Vertebral Compression Fractures (OVCFs), one of the most common complications of osteoporosis, often occur in low energy damage or the absence of a clear trauma history, which primarily results in persistent back pain, local vertebral kyphosis, a reduced quality of life as well as increased mortality[2, 3]. The prevalence increases with age, with an estimated 140 000 new vertebral fractures per year in the United States[4, 5]. Percutaneous vertebroplasty can provide instant pain relief and stabilize the fractured vertebral body through the minimally invasive injection of polymethylmethacrylate (PMMA) bone cement, which has been widely used in OVCFs treatment[6, 7]. However, the risks related with vertebroplasty are not uncommon such as kyphosis, loss of height and adjacent segment fracture, and may be related to the type of bone cement distribution[6-9]. However, few studies have reported the relationship between bone
cement distribution and imaging changes and clinical outcomes following PVP. Therefore, the present study aimed to investigate the differences in vertebral height correction, local vertebral kyphosis correction, pain relief, functional recovery, etc between the two bone cement distribution modes.

**Methods**

**General data**

The clinical data of patients with single-segment OVCFs who underwent bilateral percutaneous vertebroplasty from April 2016 to April 2019 were retrospectively reviewed. The inclusion criterias are as follows: 1. the patient had obvious thoracolumbar and back pain, and limited physical activity, especially in cases of turning over or getting up. 2. T score ≤ −2.5 at spine/hip at Dual energy X-ray absorptiometry (DXA). 3. The signal change of the lumbar fracture by lumbar magnetic resonance imaging (MRI) suggesting a hyperintense T2 signal and a hypointense T1 signal, or a whole-body bone scan performed a active bone metabolism. Exclusion criterias are as follows: 1. Patients with OVCFs caused by tumor, infection, or tuberculosis. 2. Patients had coagulation dysfunction, combined systemic disease, and inability to tolerate the procedure. 3. Systemic or local infection. 4. Spinal cord compression and obvious neural symptoms such as numbness and/or muscle weakness. 5. Incomplete follow-up data. We eventually recruited 392 patients, including 60 males and 269 females, mean age 68.21±10.48 years, and mean follow-up time 17.54 months. The present study was approved by the medical ethics committee of the Peoples Hospital of Ningxia Hui Autonomous Region. All included patients signed an informed consent.

**Surgical method**

The patient was placed in the prone position, the abdomen was vacated, and the fractured vertebrae were located under C-arm fluoroscopic guidance. The puncture needle was inserted into the vertebral body via bilateral arch pathways. The tip of the puncture needle was located in the anterior middle third of the vertebral body on the lateral view, and the anterior-posterior view was located between the inner edge of the ipsilateral pedicle and the vertebral body midline. The working channel is established and the high-viscosity cement is slowly injected under C-arm fluoroscopy until the bone cement approaches the posterior wall of the vertebral body where leakage may occur, and the working channel is slowly withdrawn after the cement has hardened. The whole procedure was done with the assistance of C-arm fluoroscopy. All patients were given oral calcium and vitamin D postoperatively and an intravenous infusion of Zoledronic acid (Aclasta, 100ml/5mg) once a year thereafter for 3 years. Patients were reviewed on the postoperative 24h for anteroposterior and lateral radiographs and discharged 2 to 3 days after surgery. X-ray films of injured vertebra were reviewed periodically after surgery.

**Grouping method**
Anteroposterior and lateral radiographs were taken 24 hours after surgery and divided into blocky group (Figure 1) and spongy groups (Figure 2) according to the difference in the distribution of bone cement in the vertebral body on the X-ray images after PVP treatment.

**Evaluation method**

Gender, nationality, age, diabetes history, hypertension history, fracture history, body mass index (BMI), Bone mineral density (BMD), fracture segment, follow up time, bone cement volume, operation duration, blood loss, adjacent vertebral fracture and bone cement leakage were documented. The Oswestry Disability Index (ODI) and Visual Analog Scale (VAS) were recorded to assess the clinical outcomes before surgery, 24h after surgery and at the last follow-up. The anterior vertebral height (AVH) and local kyphotic angle (LKA, Cobb’s method) of the fractured vertebral body were measured before surgery, 24h after surgery and at the last follow-up. AVH change was defined as postoperative AVH - preoperative AVH. The anterior vertebral height ratio (AVHR) was defined as the height of the anterior wall of the compressed vertebral body / (the height of the anterior wall of the upper vertebral body + the height of the anterior wall of the lower vertebral body)×2(Figure 3). The anterior vertebral height recovery ratio (AVHRR) was defined as postoperative AVHR - preoperative AVHR. The anterior vertebral height loss ratio (AVHLR) was defined as postoperative AVHR-last follow-up AVHR. Cobb angle was defined as the angle formed by the upper and lower endplates of the fractured vertebral body(Figure 3). Local kyphotic angle change was defined as last follow-up Cobb angle – postoperative Cobb angle.

**Statistical analysis**

The data were statistically analyzed using SPSS 22.0 software (SPSS, Inc., USA). Categorical variables were expressed as rates, and the chi-square test was used for comparison between groups. Continuous variables were expressed as mean ± standard deviation, and independent samples t-test or analysis of variance (ANOVA) was used for comparison between groups. Differences were defined as statistically significant at P<0.05.

**Results**

All patients received PVP treatment for single level OVCF and finished mean 17.54 months of follow-up (range from 11 to 36 months). There was no significant difference in gender, nationality, age, comorbidities (diabetes history, hypertension history, fracture history), BMI, BMD, fracture segment, follow up time, bone cement volume, operation duration and blood loss, as depicted in Table 1.

There were no statistically significant differences in the VAS and ODI between the two groups before the operation, but a significant reduction in pain was reported in the blocky and spongy group after surgery (Table 2). The VAS and ODI in the spongy group was significantly lower than that in the blocky group, 24h postoperatively, and at the last follow-up (Table 2).
There were 42 cases (12.8%) of adjacent vertebral fractures, 26 cases (19.8%) in blocky group and 16 cases (8.1%) in spongy group. The incidence of postoperative adjacent vertebral fractures in spongy group was significantly lower than that in blocky group, and the difference was statistically significant (Table 2). There were 57 cases (17.3%) of bone cement leakage, 18 cases (13.7%) in blocky group and 39 cases (19.7%) in spongy group (Table 2). There were no clinical symptoms between two groups, and the difference was not statistically significant.

Local kyphotic angle change was significantly smaller in spongy group (Table 2). AVHHR in spongy group is higher than that in blocky group, it was significant difference in AVHRR between the two groups (Table 2). At the last follow-up, AVHLR in spongy group was significantly lower than that in group A. AVH and AVHR were all significantly restored at 24h and last follow-up after surgery, compared with the preoperative data in groups (Table 2). Similarly, Cobb angle improved significantly after surgery for both groups (Table 2). At the 24h and last follow-up, AVH, AVHR, AVH change in spongy group were significantly higher than that in blocky group, while Cobb angle was significantly lower (Table 2).

Discussion

In 1987, Galibert first used vertebroplasty to treat C2 vertebral hemangioma. Studies have been reported on PVP for OVCFs in 1988 and 1994, respectively. Since then, the technique has been developed and refined, and it has gradually become the main methods for treating OVCFs due to its simplicity and efficacy. PVP strengthens the vertebral body by injecting bone cement into the fractured vertebral body to restore the height, strength, and stiffness of the vertebral body, while correct the local kyphosis and produce a thermal effect on the nociceptive nerves around the vertebral body to rapidly relieve pain symptoms. However, this method is affected by many factors, such as BMD, the amount, distribution, and leakage of bone cement, etc.

A biomechanical study of 120 vertebrae from 10 osteoporotic female cadavers found, on average 16.2% and 29.8% of the vertebral cement filling is required to restore strength and stiffness, respectively, and it was no correlation between the recovery of vertebral strength and stiffness and the percentage volume of bone cement filling [18]. Liebschner [19] et al. performed a single lumbar PVP finite element analysis study, which found that only a small amount of bone cement (14% volume) was required to restore the stiffness of the fractured vertebrae to pre-injury levels, and that a larger volume of bone cement did not provide greater benefit, as the increase in vertebral strength with bone cement resulted in asymmetric distribution of bone cement and strength imbalance on both sides of the vertebral body. Related studies also confirm the above-mentioned view [20, 21]. In addition, the correlation between bone cement volume and surgical outcome is small, and an increase in cement volume may increase the risk of cement leakage [22]. Bone cement injection volume is a one-sided indicator of the benefit of bone cement and does not reflect the distribution of bone cement within the vertebral body. Therefore, it is important to study the distribution of bone cement and the clinical outcome and prognosis of PVP. However, few studies have reported the effect of bone cement distribution on radiographic and functional recovery after PVP treatment.
Compared to PVP, PKP results in poorer cement distribution and a greater likelihood of postoperative vertebral height loss [23]. The main reason is that the expansion of the balloon compresses the more lax cancellous bone in the cone, thus creating a "cavity" at the balloon site, and the injected bone cement tends to be distributed in this low-pressure cavity without dispersing into the surrounding bone, making it difficult to bind tightly to the cancellous bone. Therefore, this blocky distribution of bone cement has been shown to be an important factor in vertebral body height loss. The subjects selected in this study were all post-PVP patients, excluding the influence of the surgical approach on the results.

A study by Chen [24] et al. reported that the symmetric distribution of bone cement is closely related to the stiffness of the vertebral body, and in unilateral vertebroplasty, the bone cement is often confined to the ipsilateral side of the vertebral body and cannot effectively diffuse across the midline; therefore, the end result can be a significant reduction in the stiffness of the vertebral body on the unreinforced side of the bone cement compared to the reinforced side. The biomechanical imbalance can exacerbate the pressure load on the spine, resulting in effects that are difficult to reverse, such as loss of height of the fractured vertebral body, disc degeneration in adjacent segments, and even fracture of the vertebral body in adjacent segments [25, 26]. Therefore, all subjects included in this study received bilateral arch root PVP surgery, further reducing the detrimental effects of asymmetric distribution of bone cement in the coronal plane. The majority of females in this study indicated that postmenopausal women are more likely to have osteoporosis in patients.

Spinal imaging changes such as vertebral body height and Cobb angle are often used as indicators to assess the efficacy of PVP[27]. Yan[28] et al. reported that PVP was able to restore vertebral body height and correct kyphosis. The results of the present study showed that PVP was able to significantly restore anterior vertebral body height and reduce the Cobb angle without considering the cement distribution, and if the cement was spongy in the vertebral body, it could better maintain the height of the vertebral body and reduce the risk of postoperative vertebral body height loss as well as local kyphosis. There are numerous factors that contribute to enhanced postoperative vertebral body height loss and increased kyphosis that do not require exposure to a traumatic event and may be related to the severity of osteoporosis, daily activity level, and cement distribution[25]. However, bone cement distribution is an important factor contributing to vertebral height loss[9]. He[23] et al. reported that the incidence of vertebral height loss was higher in the uninterlocked solid pattern of bone cement distribution than in the interlocked solid pattern. Furthermore, Yu[8] et al. also confirmed that the comparatively diffused pattern of bone cement distribution has better medium and long-term clinical outcomes, compared to the solid lump distribution pattern. The spongy bone cement distribution allows cancellous bone and bone cement to more fully interlock and increases vertebral strength and stiffness with greater homogenization, thus reducing the risk of vertebral height loss after PVP[29]. Our study also found a loss of vertebral body height and local kyphosis over time in the blocky and spongy group postoperatively, which is consistent with previous findings[30, 31]. In contrast, the blocky group showed more pronounced changes on imaging than the spongy group, which was related to the distribution of bone cement.
Our study is consistent with previous studies reporting\cite{32, 33} that PVP significantly relieves short-term pain and restores function, regardless of the bone cement distribution pattern. In long-term follow-up, it was also found that VAS and ODI scores were significantly lower in the spongy group than in the blocky group, suggesting that spongy bone cement distribution has better analgesic and functional recovery effects. The cause of persistent lower back pain is mainly related to insufficient filling of the bone cement\cite{34}. The bone cement does not bind effectively to the fractured vertebrae, and the low strength and stiffness of the vertebrae are not sufficient to provide effective support, resulting in a continuous loss of height\cite{35, 36}. Therefore, we speculate that the spongy distribution of the bone cement allows greater contact with the cancellous bone within the vertebral body, which can adequately immobilize the fractured fragment, increasing spinal stability and reducing micromovement of the trabeculae, thus reducing pain and achieving functional recovery\cite{34, 37}.

Liebschner\cite{19} et al. reported that the recovery of vertebral body strength was closely related to the distribution of bone cement. The strength and stiffness of the vertebral body after bone cement strengthening are significantly higher than the adjacent vertebrae, and the inhomogeneous distribution of bone cement makes the strength and stiffness of the vertebral body asymmetrical in all areas, and all of these factors tend to increase the risk of fracture of the adjacent vertebral body after PVP\cite{24, 38, 39}. Therefore, numerous studies have confirmed that homogeneous distribution of bone cement within cancellous bone can reduce stress concentration and thus reduce the risk of fracture in adjacent vertebrae\cite{29, 40}. The spongy group has a spongy and homogeneous distribution of bone cement, which can fill the cancellous bone better and reduce the concentrated stress between adjacent vertebrae. Our results confirmed that the incidence of adjacent vertebral fractures was significantly lower in the blocky group than spongy. Therefore, achieving good distribution of bone cement within cancellous bone is crucial in reducing the risk of adjacent vertebral fractures.

There is an association between the distribution of bone cement and cement leakage\cite{29}. The overall bone cement leakage rate in our findings was consistent with the results reported in previous studies\cite{41}, and most patients were clinically asymptomatic. The rate of bone cement leakage was lower in the blocky group than spongy, but the difference was not statistically significant. This may be due to the diffuse distribution of the spongy cement, which is more widely distributed than the blocky group and more likely to leak through the broken bone cortex or endplate to the intervertebral disc or paravertebral area\cite{40}. Therefore, the surgeon should carefully analyze the imaging data preoperatively and should suspend the procedure in case of intraoperative cement leakage.

**Limitations:** This study currently has some limitations. First, our study is a retrospective study with a relatively small sample size, which may result in some bias. Second, the grouping method in our study differs from previous studies, in which it may lead to subjective bias in the results. Therefore, multicenter, prospective studies with larger samples are needed to further elucidate the relationship between intravertebral bone cement distribution and clinical outcomes of PVP.

**Conclusion**
Both groups of bone cement distribution have good immediate analgesic effect. However, compared with the blocky group, the spongy group could better maintain the height of the vertebral body, correct local kyphotic, improve function, reduce the risk of postoperative adjacent vertebral fractures, and it was more effective.

**Abbreviations**

OVCFs=Osteoporotic Vertebral Compression Fractures; PVP=Percutaneous vertebroplasty; PMMA=Polymethylmethacrylate; DXA=Dual energy X-ray absorptiometry; MRI=Magnetic resonance imaging; BMD=Bone mineral density; BMI=body mass index; ODI=Oswestry Disability Index; VAS=Visual Analog Scale;

AVH=anterior vertebral heigh; LKA=local kyphotic angle; AVHR=anterior vertebral height ratio; AVHRR=anterior vertebral height recovery ratio; AVHLR=anterior vertebral height loss ratio

**Declarations**

**Ethics approval and consent to participate**

This study was approved by the institutional review boards/Ethics Committees of People's Hospital of Ningxia Hui Autonomous Region, and was conducted in compliance with the ethical principles of the Helsinki Declaration of 1975. Written informed consent was obtained from the patients or their family members.

**Consent for publication**

Not applicable.

**Availability of data and materials**

Please contact the corresponding author for data requests.

**Competing interests**

No potential conflict of interest relevant to this article was reported.

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**Authors’contributions**
All authors made substantive intellectual contributions in this study to qualify as authors. L QJ, L XX and C LJ designed this study. L QJ, L XX, W YB, F XM, G T, H XH and Jiang Xiaocheng participated in collecting and analyzing raw materials. An initial draft of the manuscript was written by L QJ and L XX. G DG, L JH, and C LJ re-drafted parts of the manuscript and provided helpful advice on the final revision. All authors were involved in writing the manuscript. All authors read and approved the final manuscript.

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References

1. Compston JE, McClung MR, Leslie WD: Osteoporosis. LANCET 2019, 393(10169):364–376.
2. Lu X, Yang J, Zhu Z, Lv X, Wu J, Huang J, Yu L, Wen Z, Luo J, Wang Y: Changes of the adjacent discs and vertebrae in patients with osteoporotic vertebral compression fractures treated with or without bone cement augmentation. SPINE J 2020, 20(7):1048–1055.
3. Kendler DL, Bauer DC, Davison KS, Dian L, Hanley DA, Harris ST, McClung MR, Miller PD, Schousboe JT, Yuen CK et al: Vertebral Fractures: Clinical Importance and Management. AM J MED 2016, 129(2):221.
4. Chen X, Guo W, Li Q, Ou Z, Lao Z, Liu Y, Zhu C, Han Z, Chu X, Cai D: Is Unilateral Percutaneous Kyphoplasty Superior to Bilateral Percutaneous Kyphoplasty for Osteoporotic Vertebral Compression Fractures? Evidence from a Systematic Review of Discordant Meta-Analyses. PAIN PHYSICIAN 2018, 21(4):327–336.
5. Yang EZ, Xu JG, Huang GZ, Xiao WZ, Liu XK, Zeng BF, Lian XF: Percutaneous Vertebroplasty Versus Conservative Treatment in Aged Patients With Acute Osteoporotic Vertebral Compression Fractures: A Prospective Randomized Controlled Clinical Study. Spine (Phila Pa 1976) 2016, 41(8):653–660.
6. Zhu RS, Kan SL, Ning GZ, Chen LX, Cao ZG, Jiang ZH, Zhang XL, Hu W: Which is the best treatment of osteoporotic vertebral compression fractures: balloon kyphoplasty, percutaneous vertebroplasty, or non-surgical treatment? A Bayesian network meta-analysis. Osteoporos Int 2019, 30(2):287–298.
7. Firanescu CE, de Vries J, Lodder P, Venmans A, Schoemaker MC, Smeets AJ, Donga E, Juttmann JR, Klazen C, Elgersma O et al: Vertebroplasty versus sham procedure for painful acute osteoporotic vertebral compression fractures (VERTOS IV): randomised sham controlled clinical trial. BMJ 2018, 361:k1551.

8. Yu W, Xiao X, Zhang J, Li Z, Wang X, Tang F, Jiang X, Zhong Y: Cement Distribution Patterns in Osteoporotic Vertebral Compression Fractures with Intravertebral Cleft: Effect on Therapeutic Efficacy. WORLD NEUROSURG 2019, 123:e408-e415.

9. Lin J, Qian L, Jiang C, Chen X, Feng F, Lao L: Bone cement distribution is a potential predictor to the reconstructive effects of unilateral percutaneous kyphoplasty in OVCFs: a retrospective study. J ORTHOP SURG RES 2018, 13(1):140.

10. Li Z, Xu Y, Xu W, Zhu X, Chen Y: The Correlation Between the Diffusion Coefficient of Bone Cement and Efficacy in Percutaneous Vertebroplasty. ORTHOPEDICS 2021, 44(1):e95-e100.

11. Lv B, Ji P, Fan X, Yuan J, Xu T, Yao X, Huang A, Zou T: Clinical Efficacy of Different Bone Cement Distribution Patterns in Percutaneous Kyphoplasty: A Retrospective Study. PAIN PHYSICIAN 2020, 23(4):E409-E416.

12. Galibert P, Deramond H, Rosat P, Le Gars D: [Preliminary note on the treatment of vertebral angioma by percutaneous acrylic vertebroplasty]. NEUROCHIRURGIE 1987, 33(2):166–168.

13. Duquesnal J, Bascoulergu Y, Leclercq R. Percutaneous injection methacrylate int he vertebral body for the treatment of various disease. Radiology, 1988, 25:369–37.

14. Jensen ME, Evans AJ, Mathis JM, Kallmes DF, Cloft HJ, Dion JE: Percutaneous polymethylmethacrylate vertebroplasty in the treatment of osteoporotic vertebral body compression fractures: technical aspects. AJNR Am J Neuroradiol 1997, 18(10):1897–1904.

15. Karmakar A, Acharya S, Biswas D, Sau A: Evaluation of Percutaneous Vertebroplasty for Management of Symptomatic Osteoporotic Compression Fracture. J Clin Diagn Res 2017, 11(8):C7-C10.

16. Xie L, Zhao ZG, Zhang SJ, Hu YB: Percutaneous vertebroplasty versus conservative treatment for osteoporotic vertebral compression fractures: An updated meta-analysis of prospective randomized controlled trials. INT J SURG 2017, 47:25–32.

17. Yang JS, Liu JJ, Chu L, Li J, Chen C, Chen H, Liu P, Yan L, Liu TJ, Hao DJ: Causes of Residual Back Pain at Early Stage After Percutaneous Vertebroplasty: A Retrospective Analysis of 1,316 Cases. PAIN PHYSICIAN 2019, 22(5):E495-E503.

18. Molloy S, Mathis JM, Belkoff SM: The effect of vertebral body percentage fill on mechanical behavior during percutaneous vertebroplasty. Spine (Phila Pa 1976) 2003, 28(14):1549–1554.

19. Liebschner MA, Rosenberg WS, Keaveny TM: Effects of bone cement volume and distribution on vertebral stiffness after vertebroplasty. Spine (Phila Pa 1976) 2001, 26(14):1547–1554.

20. Belkoff SM, Mathis JM, Jasper LE, Deramond H: The biomechanics of vertebroplasty. The effect of cement volume on mechanical behavior. Spine (Phila Pa 1976) 2001, 26(14):1537–1541.
21. Nieuwenhuijse MJ, Bollen L, van Erkel AR, Dijkstra PD: Optimal intravertebral cement volume in percutaneous vertebroplasty for painful osteoporotic vertebral compression fractures. Spine (Phila Pa 1976) 2012, 37(20):1747–1755.

22. Fu Z, Hu X, Wu Y, Zhou Z: Is There a Dose-Response Relationship of Cement Volume With Cement Leakage and Pain Relief After Vertebroplasty? Dose Response 2016, 14(4):714835597.

23. He D, Lou C, Yu W, Zhu K, Wu Z, Liu F, Chen M, Zheng L, Chen Z, Fan S: Cement Distribution Patterns Are Associated with Recompression in Cemented Vertebrae After Percutaneous Vertebroplasty: A Retrospective Study. WORLD NEUROSURG 2018, 120:e1-e7.

24. Chen B, Li Y, Xie D, Yang X, Zheng Z: Comparison of unipedicular and bipedicular kyphoplasty on the stiffness and biomechanical balance of compression fractured vertebrae. EUR SPINE J 2011, 20(8):1272–1280.

25. Yu W, Liang, Yao Z, Qiu T, Ye L, Huang X, Jiang X: Risk factors for recollapse of the augmented vertebrae after percutaneous verteoplasty for osteoporotic vertebral fractures with intravertebral vacuum cleft. Medicine (Baltimore) 2017, 96(2):e5675.

26. Lee JH, Lee DO, Lee JH, Lee HS: Comparison of radiological and clinical results of balloon kyphoplasty according to anterior height loss in the osteoporotic vertebral fracture. SPINE J 2014, 14(10):2281–2289.

27. Wang Y, Liu H, Pi B, Yang H, Qian Z, Zhu X: Clinical evaluation of percutaneous kyphoplasty in the treatment of osteolytic and osteoblastic metastatic vertebral lesions. INT J SURG 2016, 30:161–165.

28. Yan L, Jiang R, He B, Liu T, Hao D: A comparison between unilateral transverse process-pedicle and bilateral puncture techniques in percutaneous kyphoplasty. Spine (Phila Pa 1976) 2014, 39(26 Spec No.):B19-B26.

29. He S, Zhang Y, Lv N, Wang S, Wang Y, Wu S, He F, Chen A, Qian Z, Chen J: The effect of bone cement distribution on clinical efficacy after percutaneous kyphoplasty for osteoporotic vertebral compression fractures. Medicine (Baltimore) 2019, 98(50):e18217.

30. Huang S, Zhu X, Xiao D, Zhuang J, Liang G, Liang C, Zheng X, Ke Y, Chang Y: Therapeutic effect of percutaneous kyphoplasty combined with anti-osteoporosis drug on postmenopausal women with osteoporotic vertebral fracture and analysis of postoperative bone cement leakage risk factors: a retrospective cohort study. J ORTHOP SURG RES 2019, 14(1):452.

31. Patel A, Petrone B, Carter KR: Percutaneous Vertebroplasty And Kyphoplasty. 2021.

32. Watts NB, Harris ST, Genant HK: Treatment of painful osteoporotic vertebral fractures with percutaneous vertebroplasty or kyphoplasty. Osteoporos Int 2001, 12(6):429–437.

33. Tan L, Wen B, Guo Z, Chen Z: The effect of bone cement distribution on the outcome of percutaneous Vertebroplasty: a case cohort study. BMC Musculoskelet Disord 2020, 21(1):541.

34. Ye LQ, Liang, Jiang XB, Yao ZS, Lu H, Qiu T, Yu WB, Mo L, Zhang SC, Jin DX: Risk Factors for the Occurrence of Insufficient Cement Distribution in the Fractured Area after Percutaneous Vertebroplasty in Osteoporotic Vertebral Compression Fractures. PAIN PHYSICIAN 2018, 21(1):E33-E42.
35. Niu J, Zhou H, Meng Q, Shi J, Meng B, Yang H: Factors affecting recompression of augmented vertebrae after successful percutaneous balloon kyphoplasty: a retrospective analysis. ACTA RADIOL 2015, 56(11):1380–1387.

36. Kim YY, Rhyu KW: Recompression of vertebral body after balloon kyphoplasty for osteoporotic vertebral compression fracture. EUR SPINE J 2010, 19(11):1907–1912.

37. Steinmann J, Tingey CT, Cruz G, Dai Q: Biomechanical comparison of unipedicular versus bipedicular kyphoplasty. Spine (Phila Pa 1976) 2005, 30(2):201–205.

38. Furtado N, Oakland RJ, Wilcox RK, Hall RM: A biomechanical investigation of vertebroplasty in osteoporotic compression fractures and in prophylactic vertebral reinforcement. Spine (Phila Pa 1976) 2007, 32(17):E480-E487.

39. Chevalier Y, Pahr D, Charlebois M, Heini P, Schneider E, Zysset P: Cement distribution, volume, and compliance in vertebroplasty: some answers from an anatomy-based nonlinear finite element study. Spine (Phila Pa 1976) 2008, 33(16):1722–1730.

40. Chen JB, Xiao YP, Chen D, Chang JZ, Li T: Clinical observation of two bone cement distribution modes of percutaneous vertebroplasty in the treatment of thoracolumbar Kummell's disease. J ORTHOP SURG RES 2020, 15(1):250.

41. Xiang GH, Tong MJ, Lou C, Zhu SP, Guo WJ, Ke CR: The Role of Unilateral Balloon Kyphoplasty for the Treatment of Patients with OVCFS: A Systematic Review and Meta-Analysis. PAIN PHYSICIAN 2018, 21(3):209–218.

Tables

Table 1. Comparison of baseline characteristics of patients between two groups.
|                              | Blocky group | Spongy group | t/χ²-value | P-value |
|------------------------------|--------------|--------------|------------|---------|
| **Gender**                   |              |              |            |         |
| Male                         | 22①16.8%    | 38①19.2%    | 0.304      | 0.581   |
| Female                       | 109①83.2%   | 160①80.8%   |            |         |
| **Nationality**              |              |              | 0.117      | 0.732   |
| Han                          | 118①90.1%   | 176①88.9%   |            |         |
| Hui                          | 13①9.9%     | 22①11.1%    |            |         |
| **Age(years)**               |              |              | 3.968      | 0.265   |
| <60                          | 4①3.1%      | 12①6.1%     |            |         |
| 60~70                        | 47①35.9%    | 71①35.9%    |            |         |
| 70~80                        | 54①41.2%    | 89①44.9%    |            |         |
| >80                          | 26①19.8%    | 26①13.1%    |            |         |
| **Diabetes history(%)**      | 9.2%         | 8.6%         | 0.032      | 0.857   |
| No(cases)                    | 119          | 181          |            |         |
| Yes(cases)                   | 1            | 17           |            |         |
| **Hypertension history(%)**  | 48.9%        | 44.9%        | 0.483      | 0.487   |
| No(cases)                    | 67           | 109          |            |         |
| Yes(cases)                   | 64           | 89           |            |         |
| **Fracture history(%)**      | 16.8%        | 15.7%        | 0.075      | 0.784   |
| No(cases)                    | 109          | 167          |            |         |
| Yes(cases)                   | 22           | 31           |            |         |
| **BMI(kg/m²)**               | 24.25±3.61   | 23.65±3.42   | 1.537      | 0.125   |
| **Bone mineral density (T score)** | -3.23±0.85 | -3.26±0.90 | 0.260 | 0.795 |
| **Fracture segment**         |              |              | 3.544      | 0.170   |
| Thoracic (T1~9)              | 10①7.6%     | 20①10.1%    |            |         |
| Thoracolumbar (T10~L2)       | 78①59.5%    | 97①49.0%    |            |         |
| Lumbar (L3~5)                | 43①32.8%    | 81①40.9%    |            |         |
| **Follow up time(months)**   | 17.83±7.33   | 17.35±6.54   | 0.618      | 0.537   |
| **Bone cement volume(mL)**   | 4.15±1.16    | 4.11±1.11    | 0.264      | 0.792   |
|                          | Group A       | Group B       | p-value | Significance |
|--------------------------|---------------|---------------|---------|--------------|
| Operation duration (mins)| 42.6±12.04    | 41.73±11.12   | 0.67    | 0.503        |
| Blood loss (mL)          | 11.17±3.66    | 11.07±3.58    | 0.239   | 0.811        |

Table 2 Analysis of outcome between two groups
|                          | Blocky group | Spongy group | T-value | P-value |
|--------------------------|--------------|--------------|---------|---------|
| **VAS**                  |              |              |         |         |
| pre-op                   | 6.29±0.90    | 6.28±1.05    | 0.060   | 0.948   |
| 24h post-op              | 2.54±0.91    | 2.12±0.79    | 4.355   | <0.001  |
| last follow-up           | 2.13±0.738   | 1.81±0.67    | 4.090   | <0.001  |
| **ODI**                  |              |              |         |         |
| pre-op                   | 63.63±11.94  | 61.02±11.79  | 1.958   | 0.051   |
| 24h post-op              | 35.88±6.72   | 33.84±6.38   | 2.773   | 0.006   |
| last follow-up           | 26.78±5.53   | 23.2±7.59    | 4.941   | <0.001  |
| **AVH(mm)**              |              |              |         |         |
| pre-op                   | 14.60±2.96   | 15.22±3.16   | -1.785  | 0.075   |
| 24h post-op              | 16.72±3.12   | 17.62±3.19   | -2.505  | 0.013   |
| last follow-up           | 16.23±3.13   | 17.14±3.25   | -2.545  | 0.011   |
| **Cobb(%)**              |              |              |         |         |
| pre-op                   | 26.71±6.91   | 27.08±7.55   | -0.465  | 0.642   |
| 24h post-op              | 17.94±3.51   | 15.85±4.03   | 4.847   | <0.001  |
| last follow-up           | 24.31±4.92   | 21.04±4.59   | 6.142   | <0.001  |
| **AVHR(%)**              |              |              |         |         |
| 24h post-op              | 54.95±10.61  | 58.17±10.55  | -2.703  | 0.007   |
| last follow-up           | 46.58±10.96  | 50.55±10.84  | -3.237  | 0.001   |
| **AVH change(mm)**       |              |              |         |         |
| 24h post-op              | 2.13±0.73    | 2.40±0.69    | -3.427  | 0.001   |
| last follow-up           | 1.63±1.078   | 1.93±1.08    | -2.466  | 0.014   |
| **AVHLR(%)**             | 8.37±2.97    | 7.62±2.96    | 2.245   | 0.025   |
| **AVHRR(%)**             | 6.99±2.42    | 7.94±2.35    | -3.544  | <0.001  |
| **Local kyphotic angle change(°)** | 6.37±4.27 | 5.20±3.81 | 2.553 | 0.011 |
| **Adjacent vertebral fractures(%)** | 19.8% | 8.1% | 9.802 | 0.002 |
| No(cases)                | 105          | 182          |         |         |
| Yes(cases)               | 26           | 16           |         |         |
## Figures

### Figure 1

Distribution characteristics of blocky bone cement. A Anteroposterior X-ray film of local solid distribution pattern in the blocky group. B Lateral X-ray film of local solid distribution pattern in the blocky group.

| Bone cement leakage (%) | 13.7% | 19.7% | 1.953 | 0.162 |
|-------------------------|-------|-------|--------|--------|
| No(cases)               | 113   | 151   |        |        |
| Yes(cases)              | 18    | 39    |        |        |
Figure 2

Distribution characteristics of spongy bone cement. A Anteroposterior X-ray film of diffuse distribution pattern in the spongy group. B Lateral X-ray film of diffuse distribution pattern in the spongy group.

Figure 3
Radiographic evaluation of compressed vertebrae. A The anterior vertebral height ratio (AVHR) was defined as the height of the anterior wall of the compressed vertebral body (b) / (the height of the anterior wall of the upper vertebral body (a) + the height of the anterior wall of the lower vertebral body (c)) \times 2. B Cobb angle was defined as the angle formed by the upper endplates (Line a) and lower endplates (Line b) of the fractured vertebral body.