

ORIGINAL ARTICLE

Comparison between unilateral curved and bilateral straight percutaneous vertebral augmentation in the treatment of osteoporotic vertebral compression fractures: A meta-analysis

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Osteoporotic vertebral compression fractures (OVCFs) are common in the aging population, and there are approximately 1.7 million cases of OVCF in the United States (US) and Europe annually.^[1] These fractures often lead to low back pain, spinal deformity, neurological dysfunction, and even impaired quality of life.^[2] Percutaneous vertebral augmentation (PVA), including percutaneous kyphoplasty (PKP) and percutaneous vertebroplasty (PVP), is a minimally invasive interventional technique and have been proven to be a successful surgical procedure for patients with OVCFs to provide rapid pain control,

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ABSTRACT

Objectives: The aim of this meta-analysis was to compare the efficacy and safety of unilateral curved and bilateral straight percutaneous vertebral augmentation (PVA) in the treatment of osteoporotic vertebral compression fractures (OVCFs).

Materials and methods: We performed a comprehensive literature search from electronic databases including Springer, Web of Science, PubMed, Cochrane Library databases and ScienceDirect up to July 2022. Three randomized-controlled trials (RCTs) and one retrospective study which met the inclusion criteria were analyzed.

Results: There were significant differences in the operative time, injected bone cement volume, bone cement leakage rate and X-ray frequency between the bilateral straight PVA and unilateral curved PVA. No significant differences were found regarding postoperative Cobb angle, Visual Analog Scale or Oswestry Disability Index between the two groups.

Conclusion: Compared to bilateral straight PVA, unilateral curved PVA may decrease operative time, injected bone cement volume, bone cement leakage rate, and X-ray frequency in the treatment of OVCFs. However, the Cobb angle, pain, and clinical scores are comparable. Due to the limited quality and data of the evidence currently available, more high-quality RCTs are required.

Keywords: Meta-analysis, osteoporotic vertebral fractures, percutaneous, vertebral augmentation.

correct deformity, and improve postoperative function. $\ensuremath{^{[3]}}$

Using conventional straight PVA, puncture approaches include unipedicular puncture and bipedicular puncture.^[4] Traditionally, the standard technique for PVA is a bipedicular approach and

it has been shown to be associated with uniform cement distribution in the vertebral body.^[5] However, bipedicular puncture approach can increase operating time, surgical trauma to the patients, risk of bone cement leakage and X-ray exposure.^[6] During unipedicular percutaneous puncture approach, bone cement can often only fill the ipsilateral vertebral bone, leaving the contralateral vertebral bone poorly filled. The non-uniform distribution of the bone cement is likely to increase the risk of re-collapse of the non-augmented contralateral vertebral bone, particularly during lateral bending.^[7] Moreover, the unipedicular puncture requires a more aggressive, lateral-to-medial approach as compared to the bipedicular approach, which increases the risk of injury to paravertebral nerves or vessels.[8]

In recent years, an improved PVA with curved device has been introduced and applied through unipedicular puncture, achieving uniform distribution of bone cement in the vertebral body.^[9] To date, several studies^[8,10-12] have compared unilateral curved and bilateral straight PVA in treating OVCFs. However, whether unilateral curved PVA is superior to bilateral straight PVA still remains controversial. In the present study, we, therefore, performed a meta-analysis with a large sample to compare the efficacy and safety of unilateral curved and bilateral straight PVA in treating HPVA in treating OVCFs.

MATERIALS AND METHODS

Search strategy

This meta-analysis was performed according to Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines. We performed a comprehensive literature search from electronic databases such as Springer, Web of Science, PubMed, Cochrane Library databases and ScienceDirect up to July 2022. We also checked the references of the identified articles to find other relevant articles. The language of identified articles was not restricted. The keywords used for the search strategy included: "osteoporotic vertebral compression fractures", "percutaneous", "vertebral augmentation" and "curved".

Inclusion criteria

Studies were selected for inclusion if they met the following criteria: (*i*) the sample included OVCFs patients treated with PVA; (*ii*) the test group was unilateral curved PVA and the control group was bilateral straight PVA; (*iii*) the clinical outcomes included operative time, X-ray frequency, injected bone cement volume, postoperative functional score, the Visual Analog Scale (VAS) score, Cobb angle, cement leakage or other complications; and (*iv*) the study was a published comparative trial, including randomized-controlled trials (RCTs) and non-RCTs. Two independent reviewers assessed the eligibility of identified articles. A third reviewer resolved any disagreement between the reviewers.

Exclusion criteria

Studies were excluded for following reasons: (*i*) duplicate articles or articles including the same patients, content and results; (*ii*) theoretical research, case reports, meta-analyses, systematic reviews, expert comments, economic analyses and conference reports; and (*iii*) studies with non-relevant outcome.

Data extraction

Data extraction was performed independently from the included articles by two reviewers. The following information was extracted: the first author's name, the publication year, country conducted in, the size of the sample, intervention, the comparable baselines, the follow-up time and the computed endpoints in each study. Endpoints include operative time, X-ray frequency, injected bone cement volume, postoperative functional score, the VAS score, Cobb angle, bone cement leakage or other complications. Other relevant data were also extracted from the included studies. If there were incomplete data, we contacted the corresponding author through e-mail for details.

Quality assessment

The methodological quality of the RCTs were assessed with a modification of the generic evaluation tool described in the Cochrane Handbook for Systematic Reviews of Interventions.^[13] The methodological quality of non-RCTs was evaluated by the methodological index for non-randomized studies (MINORS).^[14] Two authors independently performed the methodological quality assessment. Disagreements in methodological assessment were solved by discussion, and a third reviewer was consulted if necessary.

Data analysis and statistical analysis

Statistical analysis was performed using the RevMan version 5.1 software (The Cochrane Collaboration, Oxford, UK). The I^2 values and p values were used to estimate the level of heterogeneity. When $I^2 <50\%$, p>0.1, heterogeneity could be accepted and the fixed-effects model was used for data analysis. Otherwise, significant heterogeneity was considered, and a random-effects model was used for the data analysis. Subgroup analysis was performed to investigate the sources of significant heterogeneity. For continuous variables, mean differences (MDs) and 95% confidence intervals (CIs) were calculated. For dichotomous outcomes, odds ratios (ORs) and 95% CIs were calculated.

RESULTS

Search results

A total of 58 studies were retrieved from the selected data base search. No additional study was identified through other sources. After carefully reviewing the titles and abstract, 54 studies were excluded. Finally, three RCTs and one non-RCTs were included for data extraction and meta-analysis. The detailed search process is summarized in Figure 1.

Characteristics of the included studies

General information of included studies is summarized in Table I. The baseline characteristics of two groups in all studies were comparable.

Risk of bias assessment

The methodological quality of the RCTs is shown in Figure 2. The inclusion and exclusion criteria were clearly stated in all RCTs. All RCTs reported randomized sequence generation. None of RCTs reported allocation concealment and blinding method. Unclear bias was not found due to incomplete outcome data or selective outcomes. The MINORS of the non-RCTs was 20 (Table II). The non-RCTs did not prospectively perform prospective calculation of the sample size and data collection.

Outcomes of the meta-analysis

Operative time

Operative time was recorded in four studies. Figure 3 shows that operative time in the curved group was lower than that in the straight group (MD= -13.51, 95% CI: -17.29 to -9.733; p<0.00001).

X-ray frequency

X-ray frequency was reported in three studies. Figure 4 depicts that X-ray frequency in the curved group was less than that in the straight group (MD= -8.57, 95% CI: -10.34 to -6.81; p<0.00001).

Injected bone cement volume

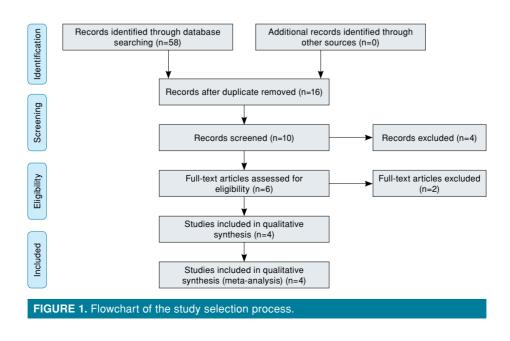
Injected bone cement volume was reported in four studies. Figure 5 shows that injected bone cement volume in the curved group was less than that in the straight group (MD= -1.13, 95% CI: -1.63 to -0.62; p<0.0001).

Postoperative Oswestry Disability Index (ODI)

Postoperative ODI was documented in three studies. Figure 6 shows that postoperative ODI in the curved group was similar to that in the straight group (MD= -0.78, 95% CI: -2.28 to 0.72; p=0.31).

Postoperative VAS

Postoperative VAS scores were assessed in four studies. Figure 7 depicts that postoperative VAS in the



trials

					TABLE I					
				Characteri	Characteristics of included studies	d studies				
Authors	Year	Study design	PVA type	Group	Cases (n)	Female (n)	Mean age (year)	Preoperative VAS	Preoperative ODI	Follow-up (months)
		TOD		Curved	30	18	71.8±11.2	8.0±1.1	I	ო
	2013		L > L	Straight	22	12	69.8±12.1	8.3±0.7	·	ო
	1000	TOD		Curved	25	17	70.7±6.8	7.9±0.6	40.1±3.5	ო
uerig ei al.	2021	j L	L > L	Straight	31	21	70.4±6.6	7.9±0.5	40.1±4.7	ო
	1000	ЦСТ		Curved	36	26	75.6±6.1	7.6±1.3	70.8±10.0	9
wang er al.	1202		L Z L	Straight	36	24	76.5±6.2	7.4±1.0	72.6±9.4	9
7home of ol 8	0100	300		Curved	29	26	70.7±7.5	8.3±1.0	75.0±11.1	12
ଯାଏଥାର କା ଲା ^ଲ	2013	0	L > L	Straight	75	63	73.8±8.2	8.2±0.8	73.4±11.7	12
PVA: Percutaneous vertebral augmentation; RCT: Randomized-controlled trial; RCS: Retrospective, controlled study; PVP: Percutaneous vertebroplasty; PKP: Percutaneous kyphoplasty; VAS: Visual Analog Scale; ODI: Oswestry Disability Index.	bral augmer Index.	ntation; RCT: Random	ized-controlled trial; I	RCS: Retrospective	e, controlled study;	PVP: Percutaneou	us vertebroplasty; Pl	<p: k)<="" p="" percutaneous=""></p:>	phoplasty; VAS: Visu	al Analog Scale;

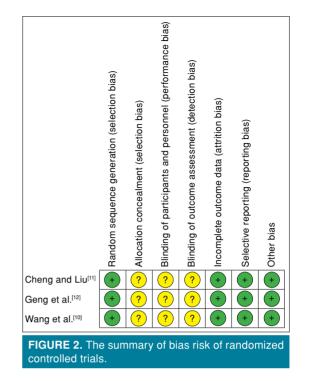


TABLE II	
tv assessment for non-randomi	zec

Quali

Quality assessment for non-randomized trials	Zhong et al. ^[8] 2019
A clearly stated aim	2
Inclusion of consecutive patients	2
Prospective data collection	0
Endpoints appropriate to the aim of the study	2
Unbiased assessment of the study endpoint	2
A follow-up period appropriate to the aims of study	2
Less than 5% loss to follow-up	2
Prospective calculation of the sample size	0
An adequate control group	2
Contemporary groups	2
Baseline equivalence of groups	2
Adequate statistical analyses	2
Total score	20

curved group was similar to that in the straight group (MD= -0.09, 95% CI: -0.32 to 0.13; p=0.43).

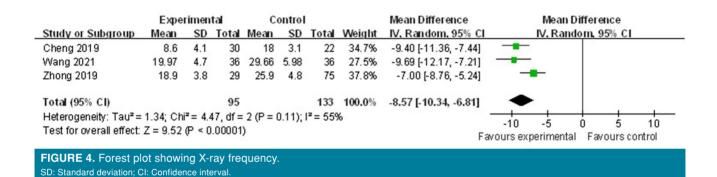
Postoperative Cobb angle

Postoperative Cobb angle was available in two studies. Figure 8 shows that postoperative Cobb angle in the curved group was similar to that in the straight group (MD= 0.17, 95% CI: -2.47 to 2.82; p=0.90).

Unilateral curved and bilateral straight PVA in treating OVCFs

	Expe	rimen	ta	C	ontrol			Mean Difference	Mean Di	fference
Study or Subgroup	Mean	SD	Tota	Mean	SD	Tota	Weight	IV, Random, 95% C	I IV, Rando	m, 95% Cl
Cheng 2019	21.9	8.9	30	37.9	6.9	22	22.7%	-16.00 [-20.30, -11.70]		
Geng 2021	38.47	3.59	25	55.42	5.72	31	28.2%	-16.95 [-19.41, -14.49]		
Wang 2021	39.3	7.87	36	48.19	9	36	23.9%	-8.89 [-12.80, -4.98]		
Zhong 2019	29.2	8	29	41	8.2	75	25.3%	-11.80 [-15.25, -8.35]		
Total (95% CI)			120			164	100.0%	-13.51 [-17.29, -9.73]	-	
Heterogeneity: Tau ² :	= 11.62; C	hi² = 1	4.45, d	f = 3 (P	= 0.00	2); I ² =	79%			
Test for overall effect									-20 -10 (Favours experimental) 10 20 Favourscontrol

FIGURE 3. Forest plot showing operative time. SD: Standard deviation; CI: Confidence interval.



	Expe	erimen	ta	C	ontrol	i -		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Tota	Mean	SD	Tota	Weight	IV, Random, 95% Cl	IV, Random, 95% CI
Cheng 2019	4.6	1.2	30	6.3	1.6	22	17.2%	-1.70 [-2.49, -0.91]	
Geng 2021	3.54	0.35	25	5.06	0.37	31	28.9%	-1.52 [-1.71, -1.33]	
Wang 2021	3.84	0.55	36	4.78	0.67	36	27.5%	-0.94 [-1.22, -0.66]	-
Zhong 2019 (L)	4.3	1.5	3	6.5	2	9	4.6%	-2.20 [-4.34, -0.06]	
Zhong 2019 (T)	4.5	2.1	2	4.2	2.2	5	1.9%	0.30 [-3.19, 3.79]	
Zhong 2019 (TL)	5.3	1.4	30	5.5	1.8	68	19.8%	-0.20 [-0.86, 0.46]	
Total (95% CI)			126			171	100.0%	-1.13 [-1.63, -0.62]	•
Heterogeneity: Tau ² =	0.22; Ch	ni² = 24	4.57, df	= 5 (P =	= 0.000)2); l ² =	80%		
Test for overall effect: 2	Z = 4.40	(P < 0	.0001)					Favo	-4 -2 0 2 4 ours experimental Favours control

FIGURE 5. Forest plot showing injected bone cement volume. SD: Standard deviation; CI: Confidence interval.

	Expe	rimen	tal	C	ontrol			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Tota	Weight	IV, Fixed, 95% (CI IV, Fixed, 95% CI
Geng 2021	19	4.7	25	20.3	4.7	31	36.7%	-1.30 [-3.78, 1.18]]
Wang 2021	23.52	4.45	36	23.97	4.43	36	53.5%	-0.45 [-2.50, 1.60]	
Zhong 2019	28.7	11.5	29	29.3	10.3	75	9.8%	-0.60 [-5.39, 4.19]	i <u> </u>
Total (95% CI)			90			142	100.0%	-0.78 [-2.28, 0.72]	
Heterogeneity: Chi2 =	0.27, df =	= 2 (P =	= 0.87)	; I ² = 0%	,				-4 -2 0 2 4
Test for overall effect	Z = 1.01	(P = 0	.31)						-4 -2 0 2 4 Favours experimental Favours control

FIGURE 6. Forest plot showing postoperative ODI. SD: Standard deviation; CI: Confidence interval.

	Expe	rimen	tal	С	Control Mean Difference				Mean Difference		
Study or Subgroup	Mean	SD	Tota	Mean	SD	Tota	Weight	IV, Random, 95% Cl	IV, Random, 95% CI		
Cheng 2019	1.2	0.5	30	1.7	0.8	22	20.0%	-0.50 [-0.88, -0.12]			
Geng 2021	2.4	0.5	25	2.5	0.4	31	29.8%	-0.10 [-0.34, 0.14]			
Wang 2021	1.33	0.47	36	1.27	0.45	36	32.3%	0.06 [-0.15, 0.27]			
Zhong 2019	1.9	1	29	1.8	0.9	75	17.9%	0.10 [-0.32, 0.52]			
Total (95% CI)			120			164	100.0%	-0.09 [-0.32, 0.13]			
Heterogeneity: Tau ² =	0.03; Ch	i ² = 7.0)7, df =	3 (P = 1	0.07);1	² = 589	6				
Test for overall effect:	Z = 0.80	(P = 0	.43)					F	-1 -0.5 0 0.5 1 avours experimental Favours control		

FIGURE 7. Forest plot showing postoperative VAS. SD: Standard deviation; CI: Confidence interval; VAS: Visual Analog Scale



SD: Standard deviation; CI: Confidence interval.

	Experime	ental	Contr	ol		Odds Ratio	Odds Rati	>
Study or Subgroup	Events	Tota	Events	Tota	Weight	M-H, Fixed, 95% CI	M-H, Fixed, 95	% CI
Cheng 2019	3	34	6	32	16.3%	0.42 [0.10, 1.84]		
Geng 2021	2	25	6	31	14.3%	0.36 [0.07, 1.98]		
Wang 2021	3	36	8	36	21.2%	0.32 [0.08, 1.32]		
Zhong 2019	8	35	36	82	48.1%	0.38 [0.15, 0.93]		
Total (95% CI)		130		181	100.0%	0.37 [0.20, 0.69]	•	
Total events	16		56					
Heterogeneity: Chi ² = 0	0.07, df = 3	(P = 0.9)	99); l ² = 0'	%		L		
Test for overall effect: 2	Z = 3.10 (P	= 0.002	2)			0.0 Favou		10 100 ourscontrol

FIGURE 9. Forest plot showing bone cement leakage rate. Cl: Confidence interval.

Bone cement leakage rate

Bone cement leakage rate was reported in four studies, of which 16 out of 130 in the curved group and 56 out of 181 in the straight group. Figure 9 depicts that bone cement leakage rate in the curved group was lower than that in the straight group (OR=0.37, 95% CI: 0.20 to 0.69; p=0.002).

Other outcomes

Zhong et al.^[8] reported that five and two patients developed fragile vertebral refractures at non-adjacent and adjacent levels, respectively (one patient developed adjacent level refractures twice), in the bipedicular vertebroplasty group. In contrast, none of the patients developed refractures at either non-adjacent or adjacent levels in the curved vertebroplasty group during the 36-month period. However, limited data could not be extracted for meta-analysis.

DISCUSSION

Our meta-analysis included three RCTs and one retrospective study. All included studies were small simple size and had relatively short follow-ups. The objective of our meta-analysis was to compare the efficacy and safety of unilateral curved and bilateral straight PVA in treating OVCFs. In the current study, we found that, compared to bilateral straight PVA, unilateral curved PVA was associated with less operative time, injected bone cement volume and X-ray frequency and lower bone cement leakage rate. To the best of our knowledge, the present study is the first quantitative meta-analysis to evaluate the efficacy and safety of unilateral curved PVA in treating OVCFs.^[15]

Bone cement leakage is the most frequent complication of PVA that has not been completely avoided. Previous studies have reported that the incidence of bone cement leakage ranges from 4.8 to 39%.^[16] Although the majority of patients was asymptomatic, bone cement leakage may lead to neurological impairment, pulmonary embolism and even death in certain severe cases.^[17] In PVA, the causes of bone cement leakage include the fracture of the perivertebral wall or endplate, injected bone cement volume and the pressure of bone cement perfusion.^[18] Cheng and Liu^[11] conducted a RCT which reported that unilateral percutaneous curved vertebroplasty was associated with less bone cement injection volume and bone cement leakage rate than bilateral PVP. Another randomized study led by Wang et al.^[10] included 72 elderly patients with OVCF. They concluded that bone cement injection volume was lower in unilateral percutaneous curved kyphoplasty group than that in conventional bilateral PKP group. In total, 16 of 130 cases in the curved group and 56 of 181 cases in the straight group reported the occurrence of bone cement leakage. The present meta-analysis showed that unilateral curved PVA reduced the risk of bone cement leakage and injected bone cement volume compared to bilateral straight PVA. Thus, unilateral curved PVA can significantly decrease bone cement leakage rate.

In PVA, repeated fluoroscopy is mandatory and helps to confirm the needle location. Both surgeons and patients may face a risk of dermatitis, cataract, and cancer, due to significant doses of radiation exposure from repeated fluoroscopy.^[19] Wang et al.^[10] compared OVCF patients receiving unilateral curved PKP with traditional bipedicular PKP, and the results showed that X-ray frequency was significantly less in unilateral curved PKP group. Zhong et al.^[8] in a retrospective study of 104 OVCFs treated with unilateral percutaneous curved vertebroplasty or bilateral PVP, found that percutaneous curved vertebroplasty could reduce fluoroscopy frequency during operation. In the current study, X-ray frequency and operation time were significantly lower in the curved group. The results of the present study are in line with these previous results.

The VAS score is the most commonly used tool for the evaluation of perioperative period. Although PVA was reported significant pain relief after PVA, residual back pain is not rare and greatly decreased patient satisfaction.^[20] Several studies^[20,21] have reported that the percentage of patients who experience unrelieved back pain after PVP ranges from 5 to 20%. In a retrospective analysis of 1,316 cases with PVP, Yang et al.[22] considered that sufficient bone cement injection with satisfactory distribution could induce a better analgesic effect. In the current study, postoperative VAS was not significantly different between the two groups. Thus, unilateral curved PVA can induce the same analgesic effect as analgesic effect as bilateral straight PVA.

Nonetheless, the limitations of present meta-analysis should be noted. First, the meta-analysis is limited to only four articles published, and the number of patients included in this meta-analysis is relatively small. Second, methodological weakness of prospective calculation of the sample size exists in non-RCTs and allocation concealment and blind method in RCTs may decrease the level of evidence. Third, postoperative height of fractured vertebrae and recurrent fractures were incomplete and we were unable to conduct a meta-analysis on these parameters.

In conclusion, compared to bilateral straight PVA, unilateral curved PVA may decrease operative time, injected bone cement volume, bone cement leakage rate, and X-ray frequency in the treatment of OVCFs. Due to the limited quality and data of the evidence currently available, more high-quality RCTs are required.

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Ethics Committee Approval: No ethical approval was required, as all data in this meta-analysis were derived from previously published research. The study was conducted in accordance with the principles of the Declaration of Helsinki.

Data Sharing Statement: The data that support the findings of this study are available from the corresponding author upon reasonable request.

Author Contributions: All authors contributed equally to the article.

Conflict of Interest: The authors declared no conflicts of interest with respect to the authorship and/or publication of this article.

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