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ORIGINAL RESEARCH

A Comparative Study on the Effects of Postoperative ¹²⁵I Brachytherapy and Irradiation After Surgical Decompression and Stabilization for Metastatic Spinal Cancers

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Purpose: The study aims to investigate and compare the efficacy and safety of intraoperative ¹²⁵I implantation and postoperative irradiation after surgical decompression and stabilization in the treatment of patients with metastatic epidural spinal cord compression (MESCC).

Methods: The study retrospectively enrolled 122 MESCC patients treated with surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy (the brachytherapy group) or postoperative radiotherapy (the irradiation group). Operation time, intraoperative blood loss, pain relief, postoperative ambulatory status, postoperative survival outcome, complications, and length of hospitalization were collected and compared between the two groups. Ten potential risk factors were analyzed for postoperative survival outcome.

Results: No significant difference was found in baseline characteristics between the two groups (P>0.05). Postoperative VAS score was significantly decreased, as compared with preoperative scores in both groups (P<0.001). The VAS in the brachytherapy group was significantly lower than that in the irradiation group at postoperative 1 month, 3 months, and 6 months (P<0.05). The postoperative ambulatory rates were 90.0% (54/60) in the brachytherapy group and 83.9% (52/62) in the irradiation group (P=0.32). The median overall survival time was similar between the two groups (7.43 months vs 7.27 months, P=0.37). Of all patients in the brachytherapy group, 25.0% (15/60) of patients suffered from complications, while 46.8% (29/62) of patients had complications in the irradiation group (P=0.0086). According to the multiple Cox regression, primary sites (P=0.038), ECOG performance status (P=0.014), and visceral metastases (P=0.0016) showed significance for postoperative survival outcome.

Conclusion: Surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy is a relatively safe and useful method in MESCC patients.

Keywords: metastatic epidural spinal cord compression, ¹²⁵I brachytherapy, radiotherapy, surgical decompression and spine stabilization, prognosis

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Introduction

Metastatic epidural spinal cord compression (MESCC), a common complication of malignant tumors, occurs when malignant tumors metastasize to the vertebra or epidural space and consequently causes spinal cord compression.^{1,2} MESCC requires interdisciplinary cooperation,^{1,2} including decompressive surgery, minimally invasive

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technique, radiotherapy, embolization, and systematic internal medicine. Recently, increasing researchers revealed that direct decompressive and stabilized surgery followed by radiotherapy was superior to radiotherapy alone in terms of postoperative ambulatory status, pain relief, and even survival prognosis.^{3–5} Therefore, surgical decompression and spine stabilization followed by postoperative radiotherapy has become increasing popular for MESCC patients.

However, patients treated with radiation had plenty of acute or subacute toxicities, subsequently requiring additional interventions, impacting patient's quality of life, and burdening medical costs. Radiation therapy was known to impair wound healing regardless of the fractionation of radiation.⁶ It was reported that 53.9% to 63.0% radiation-treated patients with spine metastasis suffered from toxicities,^{7–9} such as gastrointestinal toxicity and even spinal cord damage. Although stereotactic body radiation therapy could provide better pain relief and local control of disease with a relatively low toxicity rate (32.1% to 38.5%), it was not widely used due to high total gross charge.^{7–9}

¹²⁵I seed implantation is a booming method which vields encouraging clinical results in the treatment of spine metastasis. Percutaneous vertebroplasty (PVP) combined with ¹²⁵I implantation was superior to radiotherapy alone in terms of pain relief and patient's quality of life in patients with spine metastasis.^{10,11} Decompressive surgery combining with ¹²⁵I brachytherapy seed implantation facilitates rapid decompression, immediate metastatic radiation, and accurate tumor assessment, which has become a new modality for MESCC patients.^{12,13} However, the available studies were subject to small and retrospective samples, simple observations, and indistinct endpoints. To our knowledge, there is no report comparing the efficacy and safety of surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy and with postoperative radiotherapy in the treatment of MESCC patients.

Therefore, we speculated ¹²⁵I brachytherapy could obtain better pain relief and less radiation-related complications. The study aims to investigate and compare the efficacy and safety of surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy and postoperative radiotherapy in MESCC patients.

Methods

Patients

We retrospectively enrolled 122 patients with MESCC admitted to our departments between January 2017 and

December 2019. Patients were included if they had (1) an age of more than 18 years, (2) MRI evidence of MESCC, (3) clear diagnosis of malignant tumor, (4) severe back pain or neurological deficit due to MESCC, and (5) treated with surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy or combined with postoperative radiotherapy. Patients were excluded if they had (1) intradural metastases, (2) vertebrae metastatic lesions previously treated by surgery or radiotherapy, (3) pathological fracture in the lower limbs, (4) too poor health to undergo surgery, and (5) paralysis more than 42 hours. A study profile is shown in Figure 1. Patients were divided into two groups: patients in the brachytherapy group were treated with surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy; patients in the postoperative irradiation group were treated with surgical decompression and pedicle stabilization combined with postoperative radiotherapy. The study was approved by the Medical Research Ethics Board of the Fifth Medical Center of Chinese PLA General Hospital. Follow-up records were obtained with the patient's informed consent. The study was conducted following the Declaration of Helsinki.

Procedures

Patients in the brachytherapy group were treated with surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy. Patient's involved vertebrae were performed with a 64-slice CT scan before surgery. The scan thickness was 0.625 mm. The images were imported into 3D treatment planning system (3D-TPS). The dose of ¹²⁵I seeds was calculated according to the shape and size of lesions. The size of ¹²⁵I is 0.8 mm×4.2 mm, which is contained by titanium alloy, the half-life is 60.2 days, and the activity is 0.6~0.9 mCi per seed. Besides, the images of CT were saved in DICOM format. These files were imported into MIMICS 10.01 software (Materialise Company, Belgium) to rebuild the 3D model which was then exported as STL format. UG Imageware 12.0 (EDS. USA) was used to open the 3D model and design the appropriate position and direction of seeds insertion into the involved vertebra.

After the successful induction of general anesthesia, intubation was inserted with the help of a laryngoscope. The patient was placed in the prone position, routinely disinfected and draped in a sterile manner. G-arm fluoroscopy was used to locate the involved vertebra. Incisions were made in the center of the involved vertebra, and then the subcutaneous tissue and fascia were stripped off. The



Figure I The study profile. SCC indicates spinal cord compression; MESCC indicates metastatic epidural spinal cord compression.

lamina of the involved vertebra was exposed using a detacher. After the pedicles of the adjacent noninvolved vertebras were placed with pedicle screws, a spinous rongeur was used to bite off the spinous process and the surrounding soft tissue, and a lamina rongeur was used to remove the lamina of the involved vertebra. Palliative tumor resection of the intra-vertebra was performed. Surgeons inserted two 18G puncture needles into the front of the involved vertebrae. G-arm fluoroscopy was used to confirm the guide needle was in an ideal position. The radioactive ¹²⁵I seeds were placed in the gun and were gradually implanted into the metastatic lesions. Finally, the G-arm fluoroscopy was performed to confirm that both pedicle screws and radioactive ¹²⁵I seeds were in suitable positions. A drainage tube was placed under the incision. After the installation of the connecting rod and fixation of the end caps, the incision was sutured layer by layer. Figures 2 and 3 show a case report.

For patients in the irradiation group, surgical decompression and spine stabilization were performed within 48 hours after randomization. Apart from the implantation of radioactive ¹²⁵I seeds, the other procedures were similar to the brachytherapy group. Conventional radiation was administrated after wound healing, about 2 to 4 weeks after surgery, and the dose was 8 Gray in a single fraction.

Outcome Assessments

Patient's baseline characteristics, including age, gender, primary cancer sites, preoperative ambulatory status, Eastern Cooperative Oncology Group (ECOG) performance status, visceral metastases, bone metastasis at cancer diagnosis, extra-spinal bone metastases, and time developing motor deficits, were recorded and compared between the two groups. Operation time, intraoperative blood loss, pain relief, postoperative ambulatory status,



Figure 2 A 47-year-old female diagnosed with lung cancer treated with surgical decompression and pedicle stabilization combined with 125 I brachytherapy seeds implantation. (**A**) Preoperative lateral X-ray showed T6 and T8 vertebral collapse. (**B**) Preoperative anteroposterior X-ray showed T6 and T8 vertebral collapse. (**C**) Preoperative lateral CT showed T6 and T8 bone destruction. (**D**) Preoperative lateral MRI showed T6 and T8 spine metastases. (**E**) Preoperative transverse CT showed bone destruction at T8. (**F**) Preoperative transverse MRI showed deformation of the dural sac due to metastatic tumor.



Figure 3 (A) Postoperative lateral X-ray showed ¹²⁵I brachytherapy seeds implantation and pedicle stabilization with screws. (B) Postoperative anteroposterior X-ray showed ¹²⁵I brachytherapy seeds implantation and pedicle stabilization with screws. (C) Anterior bone scan at postoperative 10 months. (D) Posterior bone scan at postoperative 19 months. (E) Anterior bone scan at postoperative 19 months. (F) Posterior bone scan at postoperative 19 months. The above bone scan indicated no disease progression at 19 months after surgery.

postoperative survival time, length of hospitalization, and complications were analyzed between the two groups.

Operation time was defined as the time interval between skin incision and suture. Intraoperative blood loss was recorded during surgery. Pain relief was evaluated using Visual analogue score (VAS). VAS ranged from 0 representing no pain to 10 representing the maximum pain in life. VAS was collected before surgery and at the time of discharge, 1 month, 3 months, and 6 months after surgery. Postoperative ambulatory status was evaluated using Frankel scores before surgery and 4 weeks after surgery. In detail, Frankel A to C was non-ambulatory,

while Frankel D and E were ambulatory. If a patient could ambulate with or without aid, the patient was regarded as having the ability to walk.³ Survival time was defined as the time interval between surgery and death data or the last follow up. Patients who were alive at the last follow-up were censored in the overall survival analysis. Length of hospitalization was defined as the time interval between the data of hospitalization and discharge. Complications were classified into surgery-related complications, such as operation site infection, wound dehiscence, cerebrospinal fluid leakage, and epidural hematoma, radiation-related complications, such as abdominal complaints (diarrhea, abdominal pain, nausea, and vomiting) and skin complaints (itching and painful skin), and systematic complications, such as pneumonia, pulmonary embolism, stroke, septicemia, intestinal bleeding, and multiple organ failure.

Screening Predictors for Survival Outcome

The Cox regression model was used to screen the ten potential risk factors for postoperative survival outcome. The ten factors included age (years), gender (female vs male), primary cancer sites (slow growth vs moderate growth vs rapid growth, conformed to previous studies¹⁴), preoperative ambulatory status (ambulatory vs not ambulatory), ECOG performance status (1–2 vs 3–4), visceral metastases (No vs Yes), bone metastasis at cancer diagnosis (No vs Yes), extra-spinal bone metastases (No vs Yes), time developing motor deficits (\leq 14 days vs >14 days, conformed to previous studies¹⁴), and treatment strategies (the brachytherapy group vs the irradiation group).

Statistical Analysis

The Chi-square test and *t*-test were performed to analyze the baseline characteristics between the brachytherapy and irradiation group. Analysis of the VAS between the two groups was performed via repeated measures of the correlated variance model, supplemented by Wilcoxon ranksum test. Postoperative ambulatory status between the two groups was compared by the Chi-square test. Postoperative overall survival prognosis was evaluated using the Kaplan–Meier method and Log rank test. Complications were assessed by the Chi-square test and the hospitalization length was measured by Wilcoxon rank test. The univariate and multivariate analyses of postoperative survival outcome were estimated by the simple and multiple Cox proportional hazards regression models, respectively. A P value of 0.05 or less was considered statistically significant. Statistical analysis was performed using SAS 9.2 software.

Results

Patient's Characteristics

The brachytherapy group included 60 patients and the irradiation group had 62 patients. The mean age was 58.3±11.0 years old in the brachytherapy group and 57.3±14.4 years old in the irradiation group (P=0.66, Table 1). There were 31 female patients in the brachytherapy group and 34 female patients in the irradiation group (P=0.73). The majority of cancer type was rapid growth cancer, which accounted for 51.7% (31/60) in the brachytherapy group and 51.6% (32/62) in the irradiation group (P=0.13). In detail, 27 patients had lung cancer, 15 patients had breast cancer, 4 patients had liver cancer, 3 patients had prostate cancer, and 11 patients had other cancers in the brachytherapy group. In the irradiation group, 28 patients had lung cancer, 14 patients had breast cancer, 4 patients had liver cancer, 6 patients had prostate cancer, and 10 patients had other cancers. Most of the patients had unsatisfactory performance status in both groups (P=0.25). The distribution of visceral metastases, bone metastasis at cancer diagnosis, and extra-spinal bone metastases were also similar between the two groups (P>0.05).

Pain Relief

The mean pain score in 24-hour period was 7.93±1.44 in the brachytherapy group before surgery. At the time of discharge, one month, three months, and six months after the operation, mean scores decreased to 2.72±1.12 (P<0.001), 1.75±0.78 (P<0.001), 2.35±0.83 (P<0.001), and 3.39±1.00 (P<0.001), respectively (Table 2). Similar decreases in VAS were observed in the irradiation group. The mean pain score was 7.69±1.13 in the irradiation group before surgery, and the mean scores decreased to 3.02±1.50 (P<0.001) at the time of discharge, 2.95±1.31 (P<0.001) at 1 month, 3.53±1.77 (P<0.001) at three months, and 4.00 ± 1.42 at six months (P<0.001) after surgery. VAS was similar between the two groups at the time of discharge $(2.72\pm1.12 \text{ vs } 3.02\pm1.50,$ P=0.20). However, VAS in the brachytherapy group was significantly lower than that in the irradiation group one month, three months, and six months after surgery (P<0.05, Figure 4).

Function Outcome

The postoperative ambulatory rates were 90.0% (54/60) in the brachytherapy group and 83.9% (52/62) in the

Characteristics	The Brachytherapy Group (n=60)	The Irradiation Group (n=62)	P-values	
Age (mean, years)	58.3±11.0	57.3±14.4	0.66	
Gender				
Female	31	34	0.73	
Male	29	28		
Primary cancer sites				
Slow growth	15	7	0.13	
Moderate growth	14	23		
Rapid growth	31	32		
Preoperative ambulatory status				
Ambulatory	42	39	0.41	
Not Ambulatory	18	23		
ECOG performance status				
I-2	21	28	0.25	
3-4	39	34		
Visceral metastases				
No	37	29	0.10	
Yes	23	33		
Bone metastasis at cancer diagnosis				
No	33	33	0.84	
Yes	27	29		
Extraspinal bone metastases				
No	33	34	0.99	
Yes	27	28		
Time developing motor deficits				
≦ I4 days	31	30	0.72	
>14 days	29	32		

Table	I Patient	's Charad	teristics	of the	Brachy	therapy	and	Irradiation	Group

Abbreviations: ECOG, Eastern Cooperative Oncology Group; VAS, visual analogue score; MESCC, metastatic epidural spinal cord compression.

irradiation group (P=0.32, Table 2). In detail, in the brachytherapy group, 95.2% (40/42) patients who had the ability to walk before surgery sustained ambulation after surgery, and 92.3% (36/39) patients continued to ambulate in the irradiation group. Regarding patients who were unable to walk before surgery, 77.8% (14/18) of patients regained ambulation in the brachytherapy group and 69.6% (16/23) of patients in the irradiation group.

Complications and Hospitalization

Of all the patients in the brachytherapy group, 25.0% (15/ 60) suffered from complications and 46.8% (29/62) patients had complications in the irradiation group (P=0.0086, Table 2). In detail, patients in the brachytherapy group had a lower rate of radiation-related complications as compared with patients in the irradiation group (8.3% vs 38.7%, P<0.001). The rates of surgery-related (P=0.35) and systematic complications (P=0.82) were similar between the two groups. In the brachytherapy group, eight patients had surgery-related complications, five patients had radiation-related complications, and six patients had systematic complications. In detail, three patients with surgery-related complications also had radiation-related complications. One patient with surgeryrelated complication also had systematic complication. In the irradiation group, five patients had surgery-related complications, 24 patients had radiation-related complications, and seven patients had systematic complications. In detail, three patients with surgery-related complications also had radiation-related complications, four patients with radiation-related complications also had systematic complications, and two patients with surgery-related complications also had systematic complications. Five patients received radiotherapy three to eight months after surgery

Characteristics	The Brachytherapy Group	The Irradiation Group	P-values	
	Patients (n=60)	Patients (n=62)		
Operation time (mean, min)	206±36	190±34	0.017	
Intraoperative blood loss (mean, mL)	562±117	420±94	<0.001	
Preoperative VAS (mean)	7.93±1.44 ^a	7.69±1.13 ^b	0.32	
Postoperative VAS at the time of discharge	2.72±1.12 ^a	3.02±1.50 ^b	0.20	
Postoperative VAS at 1 months	1.75±0.78 ^a	2.95±1.31 ^b	<0.001	
Postoperative VAS at 3 months	2.35±0.83 ^a	3.53±1.77 ^b	<0.001	
Postoperative VAS at 6 months	3.39±1.00 ^a	4.00±1.42 ^b	0.039	
Postoperative ambulatory status				
Ambulatory	54	52	0.32	
Not ambulatory	6	10		
Postoperative survival time (median, months)	7.27	7.43	0.37	
Complications				
Yes	15	29	0.0086	
No	47	33		
Surgery-related complications				
Yes	8	5	0.35	
No	52	57		
Radiation-related complications				
Yes	5	24	<0.001	
No	55	38		
Systematic complications				
Yes	6	7	0.82	
No	54	55		
The length of hospitalization (mean, days)	21.2±5.8	19.7±6.2	0.17	

Table 2 The Comparison of Intraoperative Variables and Postoperative Clinical Outcomes Between the Brachytherapy and Irradiatic	'n
Group	

Notes: VAS, visual analogue score; ^aAll P < 0.001, as compared with the preoperative VAS in the brachytherapy group; ^bAll P < 0.001, as compared with the preoperative VAS in the irradiation group.

due to disease progression, and radiotherapy-related complication occurred in one patient. Regarding hospitalization, the mean length was 21.2 ± 5.8 days in the brachytherapy group and 19.7 ± 6.2 days in the irradiation group (P=0.17).

Survival Prognosis

The median overall survival time was 7.27 months (95% CI: 5.70–8.20 months) in the brachytherapy group and 7.43 months (95% CI: 6.43–10.73 months) in the irradiation group (P=0.37, Log rank test, Figure 5). In the brachytherapy group, the 6 and 12 months overall survival rates were 60.7% and 24.6%, respectively. In the irradiation group, the 6 and 12 months overall survival rates were 65.2% and 29.7%, respectively.

In the simple analysis of the ten factors, primary cancer sites (HR=1.55; 95% CI: 1.19–2.03, P=0.001), ECOG

performance status (HR=2.19; 95% CI: 1.43–3.34, P<0.001), and visceral metastases (HR=1.73; 95% CI: 1.17–2.55, P=0.006) were significantly associated with postoperative overall survival time, but other seven factors were not significant (Table 3). According to the multiple Cox regression analysis, primary sites (HR=1.38; 95% CI: 1.02–1.87, P=0.038), ECOG performance status (HR=1.82; 95% CI: 1.13–2.93, P=0.014), and visceral metastases (HR=1.89; 95% CI: 1.27–2.79, P=0.0016) maintained significance.

Discussion

In the present study, we retrospectively analyzed 146 MESCC patients and enrolled 122 patients. Of all the included patients, 60 patients were treated with surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy and 62 were treated with surgical



Figure 4 The mean VAS score in a 24-hour period before surgery and at the time of discharge, 1 month, three months, and six months after surgery in both groups. "pre-" indicates pre-operation; "disch-" indicates discharge; "m" indicates months; *Indicates P<0.05; ***Indicates P<0.01.



Figure 5 The postoperative overall survival curves for the two groups (P=0.37, Log rank test).

decompression and pedicle stabilization combined with postoperative radiotherapy. Although ¹²⁵I brachytherapy required more operation time and suffer from more intraoperative blood loss, it could obtain relatively better pain relief and less radiation-related complications as compared with conventional radiotherapy in the treatment of MESCC patients. The overall survival outcome and length of hospitalization were similar. Notably, the study was the first article with a large sample investigating the efficacy and safety of surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy seed implantation in the treatment of MESCC patients.

Radioactive particle brachytherapy is a precision radiation therapy which already has a history of more than one hundred years. This modality was initially performed to treat patients with prostate cancer and obtained satisfactory efficacy.¹⁵ Brachytherapy resulted in an improved biochemical relapse-free survival as compared with external beam radiotherapy alone in prostate cancer patients,¹⁶ and less acute rectal toxicity and better prostate-specific antigen control.¹⁷ So far, this modality has also been used to treat various cancer types, namely, lung cancer, oral tongue cancer, and squamous cell anal carcinoma.^{18–20}

¹²⁵I seed implantation is a booming method which yields encouraging clinical results in the treatment of spine metastasis in other studies.^{10–13} Yang et al¹⁰ compared the clinical efficacy between the combination of percutaneous vertebroplasty (PVP) and ¹²⁵I implantation and regular radiotherapy alone for the treatment of spinal osteoplastic metastasis. The combination method was superior to radiotherapy alone in terms of pain relief and patient's quality of life. One week later, the VAS in the combined group dropped from 8.73 to 3.73, but the radiotherapy group did not show a significant difference, from 8.34 to 8.43. After one month, 2.54 in the combined group versus 5.46 in the radiotherapy group. After six months,

Factors	Patients (n)	The Simple Cox Regression		The Multiple Cox Regression		
		HR (95% CI)	P-values	HR (95% CI)	P-values	
Age	122	1.01(0.99–1.03)	0.15	Not included		
Gender Female Male	65 57	1.03 (0.70–1.52)	0.87	Not included		
Primary cancer sites Slow growth Moderate growth Rapid growth	22 37 63	1.55 (1.19–2.03)	0.001	1.38 (1.02–1.87)	0.038	
Preoperative ambulatory status Ambulatory Not Ambulatory	81 41	1.32 (0.88–1.99)	0.18	Not included		
ECOG performance status I–2 3–4	49 73	2.19 (1.43–3.34)	<0.001	1.82 (1.13–2.93)	0.014	
Visceral metastases No Yes	66 56	1.73 (1.17–2.55)	0.006	1.89 (1.27–2.79)	0.0016	
Bone metastasis at cancer diagnosis No Yes	66 56	0.96 (0.65–1.41)	0.83	Not included		
Extraspinal bone metastases No Yes	67 55	0.79 (0.53–1.16)	0.22	Not included		
Time developing motor deficits ≤14 days >14 days	61 61	1.39 (0.95–2.04)	0.09	Not included		
Groups The brachytherapy group The irradiation group	60 62	0.84 (0.57–1.24)	0.37	Not included		

Abbreviations: MESCC, metastatic epidural spinal cord compression; ECOG, Eastern Cooperative Oncology Group; HR, hazard ratio; Cl, confidence interval.

1.25 versus 3.54. After 12 months, 1.32 versus 4.73. A similar decrease was shown in the score of life quality. Li et al¹¹ found PVP combined with multi-needle implantation of ¹²⁵I seeds in patients with osteolytic spinal metastasis appeared to be better than PVP combined with single-needle implantation of ¹²⁵I seeds in terms of clinical benefit rate and disease control rate because multi-needle implantation had greater bone cement injection volume and the more ideal distribution of the ¹²⁵I seeds. Decompressive surgery combining with precise ¹²⁵I brachytherapy seed implantation facilitates rapid decompression, immediate metastatic radiation, and accurate tumor assessment, which has become a new modality for

MESCC patients.^{12,13} Rogers et al¹³ reported that surgical resection and ¹²⁵I brachytherapy was well tolerated and resulted in durable local control and ambulatory function outcome in patients with MESCC. Thirty patients were included in the study. The 2- and 3-year local control rate was 87.4% and 72.9%, respectively. After surgery, 84% of patients had either normal or improve ambulatory status. The morbidity was restricted to 4 postoperative events. Qian et al¹² concluded that pedicle fixation combined with ¹²⁵I brachytherapy could effectively relieve short-term pain and improve patient's quality of life after retrospectively analyzing seven metastatic thoracolumbar tumor patients. However, the available studies were

subject to small retrospectively samples, simple observations, and indistinct endpoints.

In the study, the postoperative VAS score was significantly lower compared with the preoperative scores in both groups, which suggested that both methods could significantly improve the patient's pain relief. Importantly, the VAS in the brachytherapy group was significantly lower than that in the irradiation group at 1 month, 3 months, and 6 months after surgery, which indicated that surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy had better pain relief as compared with surgical decompression and pedicle stabilization combined with postoperative radiotherapy. The ambulatory outcome was relatively better in the brachytherapy group, as compared with the patients in the irradiation group (90.0% vs 83.9%, P=0.32), but it did not reach the significance. Regarding survival prognosis, the median overall survival time was 7.27 months in the brachytherapy group and 7.43 months in the irradiation group, which showed no significance (P=0.37), and median overall survival time of 3 to 8 months was reported in other investigations.-^{3,14,21} Moreover, patients in the brachytherapy group were less likely to develop complications as compared with patients in the irradiation group (25.0% vs 46.8%, P=0.0086). In detail, patients in the brachytherapy group had a lower rate of radiation-related complications as compared with patients in the irradiation group (8.3% vs 38.7%, P<0.001). However, the rates of surgery-related complications and systematic complications were similar between the two groups (P=0.35 and P=0.82). These results were not surprised that the brachytherapy group had less radiation-related complications as compared with the irradiation group. Although the irradiation group had less operation time and intraoperative blood loss, the mean length of hospitalization length was similar in both groups. The above-mentioned results indicated surgical decompression and pedicle stabilization combined with ¹²⁵I brachytherapy was superior to surgical decompression and pedicle stabilization combined with postoperative radiotherapy in terms of pain relief and radiation-related complications. This method has the following advantages: (1) it could relieve the mechanical pain caused by metastatic compression and receive radiation therapy immediately after surgery; (2) the target dose was high enough to better kill biological pain caused by cancer cells; (3) the particles were targeted and matched to the metastatic lesion shape so it did not increase serious complications such as spinal cord injury.

According to the multiple Cox regression analysis, primary cancer sites, ECOG performance status, and visceral metastases were significant, which indicated that the three variables were the independent predictors for postoperative survival outcome. Bollen et al²² performed a systematic review with a focus on prognostic factors associating with survival in patients with spine metastasis. The authors concluded that the prognostic factors most frequently associated with survival were the primary tumor and performance status, and the prognostic factors most frequently not associated with survival were age, gender, number of spine metastases, location of the spine metastatic lesions, and the presence of a pathologic fracture after investigating 43 different prognostic factors.

We acknowledge surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy have some strengths and weaknesses. The strengths are as follows. First, this method can achieve adequate decompression, and radiation immediately after implanting radioactive seeds. In contrast, patients who are treated with decompressive surgery cannot receive regular radiotherapy until the surgical incision is healed, about 2 to 4 weeks after surgery. During this period, the local disease may lose control due to the progression of metastatic cancer. Second, the brachytherapy is conformal radiotherapy and can be precisely implanted. The extent of radiation can be matched to the metastatic tumor. The target dose is high, while the radiation dose of the surrounding normal soft-tissue is relatively low, thus enhancing the efficacy and decreasing the risk of radiationrelated complications, such as injury to the spinal cord. Third, precise ¹²⁵I brachytherapy seed can release ray for long periods, continuously inhibiting the growth of tumor cells and preventing the occurrence of local disease. Four, the half-life period of ¹²⁵I is relatively short, which also makes it easy to protect their family members from radiation.

The weaknesses were as follows. For one thing, the halflife period of ¹²⁵I is 60.2 days, indicating that it spends about 60.2 days for the radioactive intensity decreasing to its half value. The radioactive effects on metastatic tumor lesions would also decrease with the reduction of the radioactivity. Thus, to realize better local control of disease, it is necessary to supplement the radioactivity with an additional method. For another thing, the indictor of ¹²⁵I dose is unclear. The excessive doses may result in severe radioactive damage, while insufficient doses cannot prevent the early occurrence of local disease. Therefore, further studies should investigate the relationship between the radioactive dose and corresponding effects. Besides, although this study had a relatively large sample, it was a retrospective study in a single center. Therefore, the external validation of the study needs further investigation. A prospective multicenter study with a larger sample was warranted.

Conclusion

In conclusion, surgical decompression and spine stabilization combined with ¹²⁵I brachytherapy is a relatively safe and useful method in MESCC patients.

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Disclosure

The authors report no conflicts of interest for this work.

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