Aim of the study: The purpose of this study was to evaluate the efficacy and safety of applying CyberKnife (CK) radiosurgery in patients with spinal metastases.

Material and methods: Twenty-eight patients with vertebral metastases treated using the CK system were included in the study. Eleven patients suffered from pain, and in 1 case neurological symptoms were observed. The remaining patients were free from clinical symptoms of metastatic disease. The doses applied ranged between 8 and 40 Gy delivered in 4 fractions of 8–15 Gy.

Results: In the first follow-up evaluation (mean 4.5 months after the treatment), pain was stable in 5 of 8 evaluable cases and in 3 regression occurred. The last follow-up examination (mean 11 months after stereotactic radiosurgery) revealed stable ailments in 3 of 6 evaluable cases, improvement in 3 and new complaints in another 4 patients. In 17 patients imaging studies were conducted after a mean time of 11 months after CK treatment. Stabilization was confirmed in 11, regression in 4 and progression in 2 cases. Median overall survival was 20.6 months. Median progression-free survival was 12.6 months. No side effects attributable to the treatment were observed, but during follow-up transient or permanent deterioration in neurological status as a consequence of disease progression was diagnosed in 4 patients. Delivery time of a single fraction ranged between 0.5 and 1.5 hours.

**Conclusions:** Robotic stereotactic radiosurgery as part of multimodality therapy for metastatic spinal tumours is safe and effective. Because of long irradiation times, this kind of treatment is not suitable for patients in poor general condition.

**Key words:** radiosurgery, stereotactic body radiotherapy, vertebral metastases, CyberKnife, robotic radiosurgery.

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# Evaluation of efficacy and safety of robotic stereotactic body radiosurgery and hypofractionated stereotactic radiotherapy for vertebral metastases

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#### Introduction

Malignant neoplasms were confirmed in 10 million people worldwide in the year 2000. According to the World Health Organization's (WHO) estimation, in 2015 the number of newly diagnosed malignancies will rise to 15 million [1, 2]. Prevalence of metastatic spread at the time of diagnosis is varied, depending on histopathological type of primary tumour and quality of medical care. Vertebral metastases identified in 30–50% of patients with systemic cancer are detected at higher rates in patients with primary tumours in the breast, lung, prostate and kidney [1, 3, 4]. Moreover, spinal metastases are the most common vertebral neoplasms, diagnosed 20 times more frequently than primary tumours in this localization [1, 5]. Diagnosed in all age groups, most often at age 40 to 70 years, spinal metastatic lesions are described mainly in males [1, 6]. The most common localization is the thoracic part of the spinal column (70%) [1, 7]. Pain, pathological fractures and neurological deficiencies are the main symptoms caused by the growing tumour mass. Pain is the most common sign, affecting almost 90% of patients, while motor or sensory dysfunctions are observed in 35-75% of cases at the time of diagnosis [1, 7, 8]. Early detection and appropriate treatment is crucial in minimizing the consequences of imminent disability. Median survival in patients with metastatic spread to the spinal column is usually estimated on average as 7 months [1, 8]. The short life expectancy in these patients explains the widespread provision of palliative treatment instead of radical therapy [1, 2]. In selected cases of oligometastatic disease however, radical treatment of a metastatic lesion can result in long-term survival, which implies a search for both efficient methods of identification of patients with long life expectancy and effective methods of treatment. These lesions are often surgically inaccessible and respond poorly to chemotherapy, which is especially true in the case of bone metastases. The modern approach in the management of vertebral metastases complies with structural integrity of the involved spine, the patient's general condition and individual prognosis [3]. Conventional radiotherapy (RT) of vertebral metastases, first conducted in the 1950s, is regarded as a gold standard in treatment of painful vertebral metastases [1]. Commonly used schedules – 8 Gy in 1 fraction, 20 Gy in 5 fractions and 30 Gy in 10 fractions – are considered equivalent, and their application results in a similar percentage of surviving patients, satisfactory pain control, and diminished risk of pathological fractures and spinal cord compression [1, 9, 10]. Nevertheless, low irradiation doses reduce the probability of satisfactory local control [1, 9, 10]. The median pain response is most often short, 328 contemporary oncology

with the reported range of duration of 3–6 months [11]. The main factor limiting the use of conventional radiotherapy in treatment of spinal metastases is localization in the immediate proximity of the spinal cord. Radiation-induced injury may manifest as myelopathy and acute spinal cord oedema. Radiotherapy of multiple vertebral metastases requires application of doses exceeding the dose of tolerance for the spinal cord; therefore suboptimal treatment is executed [1].

The limitations of conventional irradiation triggered application of stereotactic radiotherapy in patients suffering from various neoplasms, from malignant and benign primary tumours to spinal metastases [1, 3, 11–39]. This technique provides precise distribution of a high dose of radiation in 1 to 5 fractions. A steep dose gradient allows for protection of surrounding radiosensitive normal structures (minimizing acute and late toxicities of treatment). Furthermore, stereotactic radiosurgery (SRS) is usually performed in an out-patient fashion. This contributes to reducing the costs of the treatment and improvement of general comfort of patients and their families. Many studies have documented the CyberKnife (CK) system's ability

Table 1. Patient characteristics

Age	median range	60 30–84
Sex	female male	10 18
Zubrod performance status	0 1 2 3–4	11 15 2 0
Primary tumour	prostate breast lung medullary thyroid carcinoma others	10 6 5 2
Metastases apart from spinal column	present absent	12 16
Pain at qualification	present absent	11 17
Neurologic signs	present absent	1 27
Decompressive surgery	yes no	1 27
Chemotherapy	yes no	1 27
Hormone therapy for breast cancer	yes no	4 2
Hormone therapy for prostate cancer	yes no	9 1
Spinal metastatic lesions	isolated multiple	20 8
Localization	cervical spine thoracic spine lumbar spine sacrum	2 17 13 2

to effectively palliate spinal pain with satisfactory local control. The CK system integrates the high precision and advantages of fractionating treatment. These features are crucial in optimal management of tumours located in the vicinity of the spinal cord.

Significant improvement in general condition and pain control is usually more noticeable after SRS than after conventional RT. In fact, pain is the most common though not the only indication for SRS. Radiosurgery is applied in order to prevent or reverse neurological dysfunction and to control local progression of the metastatic lesions, as well [1]. Furthermore, this procedure is considered as the sole treatment modality for patients who are not candidates for surgical resection due to comorbidities, lack of consent for surgery or characteristics of the spinal tumour. Application of stereotactic irradiation enables treatment of recurrent tumours after ineffective usage of conventional irradiation. Spinal metastases, which are regarded as radioresistant in case of applying doses normally tolerated by the spinal cord, may be successfully and safely treated using SRS, as well [14].

The purpose of our study was to perform a comprehensive evaluation of the efficacy and safety of applying CK radiosurgery in patients with spinal metastases in terms of pain control as well as radiological and metabolic response where applicable.

### Material and methods

Twenty-eight patients (33 treatments) with vertebral metastases treated using the CK system were included in the study. The largest subset consisted of patients with prostate cancer (10 cases). The remaining diagnoses were: breast cancer (6 patients), lung cancer (5 patients), medullary thyroid carcinoma (2 patients), renal cancer (1 patient), melanoma (1 patient), renal and prostate cancer (1 patient), rectal cancer (1 patient), and PEComa (1 patient). The mean and median age was 58, and 60, respectively, and ranged between 30 and 84 years. At the time of radiosurgical treatment, distant metastases to the brain, solid organs or bones were confirmed in 12 cases. Bisphosphonates were applied in 5/28 patients before SRS. Five patients with breast cancer and 9 patients with prostate cancer were receiving hormonal treatment at the time of stereotactic radiotherapy. In 6 cases of prostate cancer, spinal column was the only location of metastatic spread. The mean and median prostate-specific antigen (PSA) level before radiosurgery was 39.6 ng/ml and 9.8 ng/ml, respectively. The doses applied ranged between 8 and 40 Gy delivered in 1-3 fractions of 4-12 Gy. The main location of the treated lesions was the thoracic part of the vertebral column (17 patients). Most patients suffered from one isolated metastatic tumour in the vertebral column (20 patients). The detailed patient characteristics are shown in Table 1.

# Treatment planning

All patients were immobilized with vacuum-lock cushions. Before the SRS procedure, high resolution MRI and CT scans were performed in every patient. Data from imaging

studies were transferred to the treatment planning system where target volumes and critical organs were outlined. In every case the Xsight Spine system was used to track the target volume. The applied doses ranged from 8 to 40 Gy (median 16 Gy) in 1–4 fractions (median 2) of 8–15 Gy. Reference isodose was 75–89% (median 80%). Estimated treatment time per fraction ranged from 33 to 101 minutes (mean 56 minutes). A sample treatment plan with pre- and post-radiosurgery imaging data is shown in Figure 1.

### Data analysis

Medical records were analyzed to assess pain control and radiological response to the treatment. In prostate cancer patients the biochemical response was also assessed by analysis of PSA levels before and after the treatment. Progression-free survival and overall survival were calculated with the Kaplan-Meier method.

#### Results

Clinical examination that appraised efficacy of the radiosurgery was performed in 25 cases. Remaining patients were not able to complete scheduled visits or preferred further treatment in centres near their place of residence.

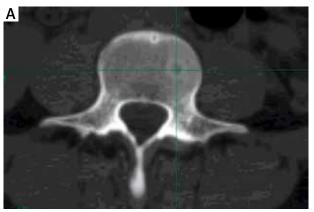
Pain as the main initial complaint in the evaluated population was confirmed in 11 cases. Neurological deficits were described in 1 case. In most cases the patients were free from clinical symptoms of metastatic tumour.

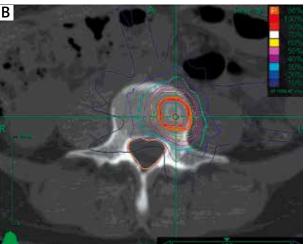
General status was assessed with the Zubrod scale. Almost all patients were in very good or good general condition (Zubrod 0 in 11 cases and Zubrod 1 in 15 cases). In only 2 patients the general status was described as satisfactory (Zubrod 2).

# Pain control and neurological deficits

The first scheduled visit was conducted a mean of 4.5 months (median 2 months) after SRS. From the group of 11 suffering from vertebral pain before SRS, 8 patients attended the first control visit. Pain was stable in 5 cases (5/8; 62.5%). Three patients reported improvement (3/8; 37.5%). The final follow-up examination was performed a mean of 11 months after radiosurgical treatment. Six patients from the group of 11 with spinal pain before SRS were assessed. Ailments were stable in 3 cases (3/6; 50%) and another 3 patients reported improvement (3/6; 50%). Progression, defined as the occurrence of pain in patients previously free from any complaints, was noted in 4 cases. The remaining patients were free from pain until the last evaluated follow-up visit.

Neurological status was assessed retrospectively by a neurologist, based on written information in medical records. The patient with neurological deficits (lower limb paresis) described before SRS was lost to follow-up. The first follow-up visit revealed deterioration in neurological status (as a consequence of progression of the metastatic spread) in 4 patients (disturbances in skin sensation and lower limb paresis). Decline was transient in 1 case (foot paresis). In 2 cases of lower limb paresis, ailments were present in the final assessment. One patient did not attend the scheduled visit.



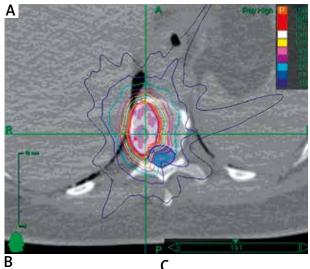




**Fig. 1. A)** CT image of a case of a 30-year-old female suffering from thyroid cancer with a painful metastatic lesion of L4. **B)** Image from the treatment planning system with the dose distribution. The patient received 16 Gy in 2 fractions using CK stereotactic radiosurgery. The reference isodose (orange) tightly encompasses the planning target volume (red). The spinal cord is spared by appropriate shaping of the isodoses. **C)** Computed tomography performed 2 years after CK treatment presents stable disease. This patient experienced satisfactory pain control as well

After SRS treatment, pharmacotherapy with bisphosphonates was continued in 3 cases (data were not available for 2 patients unable to attend the follow-up visit). In another 4 patients, bisphosphonates were newly applied after SRS completion.

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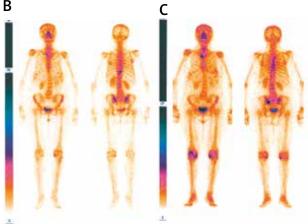


Fig. 2. A) A representative radiosurgical plan of a 65-year-old patient with a breast cancer metastasis of Th10. Figure represents optimal coverage of the target volume (purple line) with maximal spinal cord (blue line) sparing. B) Metastatic lesion of Th10 on bone scintigraphy performed before CK treatment. C) Regression of the metastatic lesion of Th10 confirmed on bone scintigraphy conducted 19 months after radiosurgery

### Prostate-specific antigen level

After CK treatment, PSA levels were quantified during the first and last control. Mean and median values were 51 ng/ml and 7.5 ng/ml (first visit) and 489.4 ng/ml and 19.5 ng/ml (last visit), respectively. After SRS, 7 patients continued hormone therapy, and hormonal treatment was replaced with a different scheme in 6 patients. In 1 case testicular irradiation was conducted, and in 1 case hormone therapy was discontinued.

### Local control

In 17 patients imaging studies were conducted to assess the efficacy of SRS treatment at a mean of 11 months after SRS (final follow-up visit). In 11 cases stabilization was confirmed. Regression was described in 4 patients (figure 2). In 2 patients progression was diagnosed.

# Adjuvant and salvage treatment after stereotactic radiosurgery

Hormonal treatment was continued in 7 of 9 patients with prostate cancer and 3 of 4 patients with breast can-

cer. Nine patients were qualified for palliative chemotherapy. Three received palliative radiotherapy (localization different than primary treated lesions). Only 1 patient was re-irradiated using stereotactic radiotherapy (CK) at 9 months after SRS (the same lesion). Three patients received additional surgical treatment after SRS (vertebroplasty in 1, laminectomy in 1, and total excision of paraspinal tumour in 1 case).

#### Overall survival and general condition

Median overall survival was 20.6 months. Median progression-free survival in patients who attended the follow-up was 12.6 months. General condition did not deteriorate in 12/21 (57%). Eight patients were evaluated below the previous value on the Zubrod scale (8/21; 38%). In 1 case an improvement was observed (5%).

### Side effects

No side effects were observed in our study. There were no cases of acute or late toxicities in the investigated population. Early tolerance of the spinal cord was satisfactory, and no case of acute spinal oedema was detected. No case of acute post-irradiation damage of the adjacent structures (laryngitis, oesophagitis or diarrhoea) was noted, either. Similarly, there was no case of remote toxicities. Deterioration in neurological status diagnosed in 4 patients was a result of ongoing metastatic spread or local progression.

# Discussion

According to the meta-analysis by Sohn *et al.*, the number of studies and the quality of available literature on spinal metastases are generally low or very low [15]. There are only two randomized controlled studies on radiosurgical treatment of spinal metastases, and only one investigation has been closed and published to date [11, 16]. The majority of the available literature represents retrospective studies analyzing outcomes of SRS in populations from 20 to 1075 patients [1, 3, 15, 17–24].

# Pain control and neurological improvement

In our evaluation, pain control was satisfactory in all patients who attended scheduled control visits (stable in 62.5% and 50% in the first and last assessment; improvement in 37.5% and 50% at the first and last visit; respectively). The deterioration in neurological status observed in our population was a consequence of progression of the disease.

Outcomes similar to ours have been presented in the literature. Ryu *et al.* described complete pain relief in 46%, partial relief in 18.9%, and stable symptoms in 16.2% at 8 weeks after SRS completion. The overall 1-year pain control rate was 84%. Significant improvement in neurological function was confirmed in 40% of patients at 1 week after SRS [17]. Likewise, Gerszten *et al.* observed long-term satisfactory pain control in 86% of patients in median follow-up of 21 months. Neurological status improved in 85% of cases [24]. Furthermore, Gibbs *et al.* confirmed a decrease in pain and neurological symptoms in 84% of

patients after SRS [15]. Nikolajek *et al.* described improvement in pain control in 75% of patients (all received SRS after previous conventionally fractionated radiotherapy) and progression of neurological symptoms in 12.9% [22]. Gagnon *et al.* described a decrease of pain within 1 week for most patients and total pain reduction in 38% at the end of the month after SRS [27]. Similarly, in the analysis by Wowra *et al.*, most patients experienced significant pain relief in a follow-up period of 1.4 years. Pain recurrence was observed in 3 (6%) patients [28]. Finally, Wang *et al.* reported significant pain improvement and reduction in opioid use in most patients during 6 months after stereotactic body radiation therapy (SBRT) (52.9% did not experience any pain), as well [16].

#### Local control and overall survival

Local control in patients receiving SRS is usually estimated at about 90% [3, 15, 24]. Our outcome with 88.2% of local control at a mean of 11 months after SRS is in line with the available literature. The median overall survival of 20.6 months and median progression-free survival of 12.6 months are similar to the results of other studies, as well.

Degen et al. confirmed 95% local control in the mean follow-up of 350 days [32]. Similarly, Chang et al. reported a 1-year 84% progression-free survival [30]. According to Gill's evaluation, local control in 1 year and in 2 years were 80% and 73%, respectively. The overall survival rate was 80 and 57% in 1 year and in 2 years, respectively [3]. Furthermore, Ryu et al. reported the 1-year survival rate as 74.3% [17]. Nikolajek et al. confirmed 12.9% of local failures. The actuarial rates of freedom from local failure at 6/12/18 months were 93%/88%/85%, respectively. Median overall survival was 16.2 months after SRS [22]. According to Gerszten et al. and Yamada et al., satisfactory control was achieved in 90% of patients [25, 26]. Finally, Wang et al. estimated actuarial tumour progression-free survival rates of 80.5% and 72.4% at 1 year and 2 years after SBRT, respectively. The median overall survival time was 23 months after SBRT, and 1-year and 2-year actuarial survival rates were 71.9% and 48.8%, respectively [16].

#### Doses and fractionation schedules

Dose prescription is determined based on primary tumour histopathology and spinal cord tolerance [15]. Single fraction SRS doses usually range from 8 to 24 Gy. There are numerous hypofractionation schedules, e.g.: 4 Gy × 5 fractions, 6 Gy × 5 fractions, 8 Gy × 3 fractions or 9 Gy × 3 fractions [15]. In our evaluation the mean applied dose was 16.8 Gy (range 8-40 Gy), delivered in 1-4 fractions of mean 8.7 Gy (range 4–15 Gy). This dose range is similar to that reported by other authors commonly applying 10 Gy to 30 Gy in 1 to 5 fractions [3, 14–18, 23–28]. An important issue is also the time needed to deliver a single fraction. Due to their complexity and numerous beams used, CyberKnife plans require long irradiation times [29]. In the current study it ranged from half an hour to one and a half hours per fraction. Such a long delivery time is not suitable for patients in poor general condition. Consequently, this kind of treatment should be offered to patients in good general status and with a long expected time of survival, preferentially to patients with single metastases and a controlled primary, which could give a chance of a cure.

### Complications

Stereotactic radiosurgery is considered as a safe treatment option. The reported complications of spinal radiosurgery include: oesophagitis, dysphagia, diarrhoea, paraesthesia, laryngitis, compression fracture and transient radiculitis [15, 30-35]. Radiation-induced spinal cord injury is an extremely rare complication [15]. Our evaluation did not reveal any early and/or late complications of SRS treatment. Similarly, many authors did not report any complications [3, 15, 17, 27, 28]. Nikolajek et al. documented 1 case of progressive paraparesis. Apart from that, no other toxicities have been observed [22]. Gibbs and colleagues described 3 cases and subsequently 6 cases of spinal cord injuries in their studies [18, 23]. However, Wang et al. [16] described 12 cases of non-neurological grade 3 toxicities (nausea, vomiting, diarrhoea, fatigue, dysphagia, neck pain, diaphoresis, pain associated with severe tongue oedema and trismus, non-cardiac chest pain). No radiation-induced myelopathy occurred. Against the background of the cited papers, the treatment tolerance in our series is excellent; the patients are however still being followed up whenever possible in order not to miss possible late complications.

# Hormonal status

In our study, radiosurgical treatment did not change the PSA level significantly in the early period after SRS. The later elevation of the PSA levels can be attributed to progression in unirradiated sites (all but 1 patient with increasing PSA levels had other metastases) rather than to local failure in the irradiated area. According to available literature, application of CK radiosurgery may cause a decrease of PSA level [36–39]. In contrast to some other studies, in our investigation a large subset of patients was receiving hormone therapy for many weeks or months preceding SRS. Moreover, many patients required modification of hormonal treatment after SRS. Thus, precise and reliable determination of dependence between PSA level fluctuations and CK radiosurgery timing and dose is hampered in our patients.

In conclusion, robotic stereotactic radiosurgery, as a part of multimodality therapy for spinal tumours, is safe and effective. Because of long treatment times, patients in good general status should be selected for this kind of treatment, preferentially without signs of active disease outside the treated area.

The authors declare no conflict of interest.

#### References

1. Harel R, Angelov L. Spine metastases: Current treatments and future directions. Eur J Cancer 2010; 46: 2696-707.

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- Hayat MJ, Howlader N, Reichman ME, Edwards BK. Cancer statistics, trends, and multiple primary cancer analyses from the surveillance, epidemiology, and end results (SEER) program. Oncologist 2007; 12: 20-37.
- 3. Gill B, Oermann E, Ju A, et al. Fiducial-free CyberKnife stereotactic body radiation therapy for single vertebral body metastases: acceptable local control and normal tissue tolerance with five-fraction approach. Front Oncol 2012; 2: 1-6.
- 4. Ries LA, Melbert D, Krapcho M, et al. SEER Cancer Statistics Review 2008; 1975-2005.
- Perrin RG, Laxton AW. Metastatic spine disease: epidemiology, pathophysiology, and evaluation of patients. Neurosurg Clin N Am 2004; 15: 365-73.
- Constans JP, de Divitiis E, Donzelli R, et al. Spinal metastases with neurological manifestations. Review of 600 cases. J Neurosurg 1983; 59: 111-8.
- Camins MB, Jenkins AL, Singhal A, Perrin RG. Tumors of the vertebral axis: benign, primary malignant, and metastatic tumors. In: Winn RH, Youmans JR (eds.). Youmans neurological surgery. 5<sup>th</sup> ed. W.B. Saunders Company, Philadelphia 2004; 4835-68.
- 8. van der Linden YM, Dijkstra SP, Vonk EJ, Marijnen CA, Leer JW; Dutch Bone Metastasis Study Group. Prediction of survival in patients with metastases in the spinal column: results based on a randomized trial of radiotherapy. Cancer 2005; 103: 320-8.
- Bone Pain Trial Working Party.
  Gy single fraction radiotherapy for the treatment of metastatic skeletal pain: randomised comparison with a multifraction schedule over 12 months of patient follow-up. Radiother Oncol 1999;
   111-21.
- Rades D, Stalpers LJ, Veninga T, et al. Evaluation of five radiation schedules and prognostic factors for metastatic spinal cord compression. J Clin Oncol 2005; 23: 3366-75.
- 11. Guckenberger M, Hawkins M, Flentje M, Sweeney RA. Fractionated radiosurgery for painful spinal metastases: DOSIS a phase II trial. BMC Cancer 2012: 12: 530
- 12. Rykaczewski B, Zabek M. A meta-analysis of treatment of vestibular schwannoma using Gamma Knife radiosurgery. Contemp Oncol (Pozn) 2014; 18: 60-6.
- Sokal P, Lebioda A, Harat M, et al. Preliminary results of linac-based radiosurgery in arteriovenous malformations and cerebral tumours in the Oncology Centre in Bydgoszcz. Contemp Oncol (Pozn) 2013; 17: 29-33.
- 14. Levine AM. Stereotactic radiosurgery for the treatment of tumors of the spine. Semin Spine Surg 2009; 21: 129-38.
- 15. Sohn S, Chung CK. The role of stereotactic radiosurgery in metastasis to the spine. J Korean Neurosurg Soc 2012; 51: 1-7.
- Wang XS, Rhines LD, Shiu AS, et al. Stereotactic body radiation therapy for management of spinal metastases in patients without spinal cord compression: a phase 1-2 trial. Lancet Oncol 2012; 13: 395-402.
- 17. Ryu S, Jin R, Jin JY, Chen Q, Rock J, Anderson J, Movsas B. Pain control by image-guided radiosurgery for solitary spinal metastasis. J Pain Symptom Manage 2008; 35: 292-98.
- 18. Gibbs IC, Kamnerdsupaphon P, Ryu MR, Dodd R, Kiernan M, Chang SD, Adler JR Jr. Image-guided robotic radiosurgery for spinal metastases. Radiother Oncol 2007; 82: 185-190.
- 19. Wowra B, Muacevic A, Zausinger S, Tonn JC. Radiosurgery for spinal malignant tumors. Dtsch Arztebl Int 2009; 106: 106-12.
- Choi CY, Adler JR, Gibbs IC, Chang SD, Jackson PS, Minn AY, Lieberson RE, Soltys SG. Stereotactic radiosurgery for treatment of spinal metastases recurring in close proximity to previously irradiated spinal cord. Int J Radiat Oncol Biol Phys 2010; 78: 499-506.
- 21. Gerszten PC, Burton SA. Clinical assessment of stereotactic IGRT: spinal radiosurgery. Med Dosim 2008; 33: 107-16.
- Nikolajek K, Kufeld M, Muacevic A, Wowra B, Niyazi M, Ganswindt U. Spinal radiosurgery – efficacy and safety after prior conventional radiotherapy. Radiat Oncol 2011; 6: 173.
- 23. Gibbs IC, Patil C, Gerszten PC, Adler JR Jr, Burton SA. Delayed radiation-induced myelopathy after spinal radiosurgery. Neurosurgery 2009; 64: A67-72.
- Gerszten PC, Burton SA, Ozhasoglu C, Welch WC. Radiosurgery for spinal metastases: clinical experience in 500 cases from a single institution. Spine (Phila Pa 1976) 2007; 32: 193-9.

- 25. Gerszten PC, Mendel E, Yamada Y. Radiotherapy and radio- surgery for metastatic spine disease: what are the options, indications, and outcomes. Spine (Phila Pa 1976) 2009; 34: S78-92.
- Yamada Y, Bilsky MH, Lovelock DM, et al. High-dose, single-fraction image-guided intensity-modulated radiotherapy for metastatic spinal lesions. Int J Radiat Oncol Biol Phys 2008; 71: 484-90.
- 27. Gagnon GJ, Nasr NM, Liao JJ, Molzahn I, Marsh D, McRae D, Henderson FC Sr. Treatment of spinal tumors using Cyberknife fractionated stereotactic radiosurgery: Pain and quality-of life assessment after treatment in 200 patients. Neurosurgery 2008; 64: 297-306.
- Wowra B, Zausinger S, Drexler C, Kufeld M, Muacevic A, Staehler M, Tonn JC. CyberKnife radiosurgery for malignant spinal tumors: characterization of well-suited patients. Spine (Phila Pa 1976) 2008; 33: 2929-34.
- 29. Blamek S, Grządziel A, Miszczyk L. Robotic radiosurgery versus micro-multileaf collimator: a dosimetric comparison for large or critically located arteriovenous malformations. Radiat Oncol 2013; 8: 205
- 30. Chang EL, Shiu AS, Mendel E, et al. Phase I/II study of stereotactic body radiotherapy for spinal metastasis and its pattern of failure. J Neurosurg Spine 2007; 7: 151-60.
- 31. Benzil DL, Saboori M, Mogilner AY, Rocchio R, Moorthy CR. Safety and efficacy of stereotactic radiosurgery for tumors of the spine. J Neurosurg 2004; 101 Suppl 3: 413-8.
- 32. Degen JW, Gagnon GJ, Voyadzis JM, McRae DA, Lunsden M, Dieterich S, Molzahn I, Henderson FC. CyberKnife stereotactic radiosurgical treatment of spinal tumors for pain control and quality of life. J Neurosurg Spine 2005; 2: 540-9.
- 33. Gagnon GJ, Henderson FC, Gehan EA, Sanford D, Collins BT, Moulds JC, Dritschilo A. Cyberknife radiosurgery for breast cancer spine metastases: a matched-pair analysis. Cancer 2007; 110: 1796-802.
- 34. Haley ML, Gerszten PC, Heron DE, Chang YF, Atteberry DS, Burton SA. Efficacy and cost-effectiveness analysis of external beam and stereotactic body radiation therapy in the treatment of spine metastases: a matched-pair analysis. J Neurosurg Spine 2011; 14: 537-42.
- 35. Hamilton AJ, Lulu BA, Fosmire H, Stea B, Cassady JR. Preliminary clinical experience with linear accelerator-based spinal stereotactic radiosurgery. Neurosurgery 1995; 36: 311-9.
- 36. Ahmed KA, Barney BM, Davis BJ, Park SS, Kwon ED, Olivier KR. Stereotactic body radiation therapy in the treatment of oligometastatic prostate cancer. Front Oncol 2013; 2: 215.
- Muacevic A, Kufeld M, Rist C, Wowra B, Stief C, Staehler M. Safety and feasibility of image-guided robotic radiosurgery for patients with limited bone metastases of prostate cancer. Urol Oncol 2013; 31: 455-60
- 38. Jereczek-Fossa BA, Beltramo G, Fariselli L, et al. Robotic image-guided stereotactic radiotherapy, for isolated recurrent primary, lymph node or metastatic prostate cancer. Int J Radiat Oncol Biol Phys 2012; 82: 889-97.
- 39. Napieralska A, Miszczyk L, Tukiendorf A, Stąpór-Fudzińska M. Wyniki leczenia przerzutów raka stercza do kości po zastosowaniu radiochirurgii CyberKnife. Ortop Traumatol Rehabil [in press].

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