

Stereotactic body radiation therapy via helical tomotherapy to replace brachytherapy for brachytherapy-unsuitable cervical cancer patients – a preliminary result

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Aim: To review the experience and to evaluate the results of stereotactic body radiation therapy (SBRT) via helical tomotherapy (HT), for the treatment of brachytherapy-unsuitable cervical cancer.

Methods: Between September 1, 2008 to January 31, 2012, nine cervical cancer patients unsuitable for brachytherapy were enrolled. All of the patients received definitive whole pelvic radiotherapy with or without chemotherapy, followed by SBRT via HT.

Results: The actuarial locoregional control rate at 3 years was 78%. The mean biological equivalent dose in 2-Gy fractions of the tumor, rectum, bladder, and intestines was 76.0 ± 7.3 , 73.8 ± 13.2 , 70.5 ± 10.0 , and 43.1 ± 7.1 , respectively. Only two had residual tumors after treatment, and the others were tumor-free. Two patients experienced grade 3 acute toxicity: one had diarrhea; and another experienced thrombocytopenia. There were no grade 3 or 4 subacute toxicities. Three patients suffered from manageable rectal bleeding in months 11, 14, and 25, respectively. One stage IVA patient experienced fistula formation in month 3.

Conclusion: SBRT via HT provides the possibility for treatment of locally advanced cervical cancer in patients who are unsuitable for brachytherapy. Long-term follow up and enrollment of more such patients to receive SBRT via the HT technique are warranted.

Keywords: biological equivalent dose, complication, image guidance, intensity modulated radiation therapy, rectal bleeding

Background

For patients with contraindications to intracavity brachytherapy (BT), further conformal external beam radiotherapy (EBRT), in the form of a cervical boost, may provide benefits of increasing the dose to the central pelvis.¹ Nevertheless, a previous study has shown that EBRT with conventional techniques used throughout the treatment course for cervical cancer is associated with poor outcomes and a high incidence of side effects.²

The application of intensity-modulated radiotherapy (IMRT) to gynecologic malignancies has been reported to provide excellent planning target volume (PTV) coverage and is associated with fewer sequelae (by normal tissue sparing) than conventional radiotherapy (RT). Additionally, using these characteristics of IMRT as a final boost in gynecologic malignancies with contraindications for BT has also shown encouraging results.³

Helical tomotherapy (HT), an image-guided IMRT, can deliver highly conformal dose distributions and provides a more impressive critical organ-sparing ability for cervical cancer than does IMRT.⁴ With the uniqueness of HT, the applications of

stereotactic body RT (SBRT), administered via a HT system in place of BT, could be an effective and well-tolerated treatment for cervical cancer.⁵

Here, we report on patients with locally advanced cervical cancer that were treated with HT-guided SBRT rather than BT because clinical judgment indicated contraindications to the use of that type of conventional treatment modality.

We aimed to assess the tolerance, clinical outcomes, and toxicities.

Methods

Patient characteristics

During the period between September 1, 2008 to January 31, 2012, nine patients undergoing whole pelvic radiotherapy (WPRT) for locally advanced cervical cancer contraindicated for BT were retrospectively enrolled, with the approval of the Institutional Review Board. Staging investigations included complete history and physical examination, fiberoptic endoscopic evaluation, complete blood count, liver and renal function tests, chest X-ray, and magnetic resonance imaging (MRI) scans or computed tomography (CT) scans of the pelvic region. The disease was staged according to the International Federation of Gynecology and Obstetrics (FIGO) criteria.⁶

Radiotherapy

Radiotherapy was administered as WPRT followed by SBRT via HT. The total dose of SBRT delivered to patients was 27-16 Gy/5-9 fractions. Weekly cisplatin, beginning on the first day of radiation, was administered during external radiation. A dose of 40 mg/m² cisplatin (maximum dose, 70 mg) was used and administered to patients via a peripheral vein, if patients received concurrent chemoradiation therapy (CCRT).

Delineation of target volumes

All patients underwent a CT planning scan (Somatom[®] Plus 4 CT scanner; Siemens AG, Munich, Germany), from the diaphragm to 5 cm below the ischial tuberosities. CT with 3 mm slice thickness was taken for treatment planning. Target objects and normal structures were contoured on a Pinnacle³ treatment planning system (Philips Medical System, Fitchburg, WI, USA). Delineation and constraints were based on Radiation Therapy Oncology Group (RTOG) 0418 protocol, the International Commission on Radiation Units and Measurements (ICRU) Report 50 and Report 62 recommendations, and our hospital guidelines.⁴ Briefly, the gross tumor volume was defined as all known gross disease determined from CT, clinical information, and MRI. The gross tumor

volume plus a 7 mm expansion was defined as the primary tumor clinical target volume (CTV), excluding the bowel, bladder, and rectum, if they were not clinically involved. The internal target volume was defined as the volume of the vagina and paravaginal soft tissues that was in both the empty and full bladder CT scans. The PTV provided a 7 mm margin around the nodal CTV and internal target volume with three-dimension (3D) expansion. The treatment plan was carried out on the full bladder scan. Identification of the nodal CTV usually began with the identification of the iliac vessels down to the level of S3. The average margin was 7 mm. Bone, iliopsoas muscle, and the intraperitoneal small bowel were excluded from the nodal CTV. Approximately 1.5 cm of tissue anterior to the S1-3 sacral segments was usually added to the CTV in order to include the presacral lymph nodes and uterosacral ligaments. The CTV of the nodes ended 7 mm from L4/L5 interspace, to account for the PTV. The PTV for nodes was stopped at the L4/L5 interspace. The lateral margin of the vaginal PTV extended to the obturator muscle, and at least 3 cm of the vagina needed to be treated. The 90% isodose surface covered between 95% and 98% of the PTV 50.4; volumes of overdose exceeding 115% < 5% of the PTV 50.4 volume could be considered acceptable.

Normal structures were contoured using the full-bladder CT scan. Dose-volume constraints for normal tissues were as follows: small bowel (2 cm above the most superior vessel contour) <30% to receive \geq 40 Gy; rectum < 60% to receive \geq 30 Gy; bladder < 35% to receive \geq 45 Gy; femoral head \leq 15% to receive \geq 30 Gy; pelvic bone marrow, V10 < 95% and V20 < 76%.

SBRT via HT substitute for BT

The CTV_{boost} was defined as the area of residual tumor and gross disease determined from primary CT or MRI which will be boosted by SBRT. The PTV_{boost} provided a 5 mm margin around the CTV_{boost} with 3D expansion. The treatment plans were carried out with a full bladder scan and with, or without, rectal balloon (30-40 cc) insertion. All patients received megavoltage CT (MVCT) scanning every time before SBRT treatment. Patients were treated every day or every other day after WPRT was completed.

Toxicity

Interruptions in RT were at times necessitated by uncontrolled diarrhea or other acute complications. If RT was held, then chemotherapy was also held. Chemotherapy stopped at the completion of RT. Radiation was only stopped in cases of grade 4 hematologic or nonhematologic toxicity and until

toxicity was resolved to at least grade 3; however, cisplatin was withheld in any case involving grade 3 toxicity, until the toxicity regressed to any grade of <3, and in patients with grade 3 toxicity that persisted > 2 weeks, chemotherapy was no longer administered.

Determination of organs at risk (OARs) dose and complications

The mean and maximum doses for the bladder, intestine, and rectum were recorded and summed with the previous plan for evaluation. The resulting dose was calculated into the biological equivalent dose (BED) in 2-Gy fractions (EQD2) using a linear-quadratic model, assuming α/β ratio = 10 for tumor and $\alpha/\beta = 3$ for OARs. Doses were normalized using this formula and were denoted by Gy3 (critical normal organs) or Gy10 (tumor). The BED for the bladder, intestine, and rectum was determined by adding the components of EBRT and SBRT. The equation used in the calculation for the total mean and max BED is as follows:

$$\begin{aligned} \text{BED}_{\text{total}} &= \text{BED}_{\text{WPRT}} + \text{BED}_{\text{boost}} \\ &= Nd (1 + d/[\alpha/\beta]) + Sb (1 + b/[\alpha/\beta]) \quad (1) \end{aligned}$$

where Nd is the tumor or OARs dose of EBRT in Gray, d is the fraction dose of EBRT in Gray, Sb is the tumor or OARs dose of SBRT in Gray, and b is the fraction dose of SBRT in Gray.⁷

Follow up

Upon treatment completion, patients were evaluated every 3 months for the first year, every 4 months during the second year, every 6 months during the third year, and annually thereafter. At each visit, a physical and pelvic examination, blood count clinical chemistry, and chest x-ray were performed. A Papanicolaou (PAP) test, CT or MRI scan, ultrasound, and other imaging studies were conducted, when appropriate. Suspected cases of persistent or recurrent disease were confirmed by biopsy, whenever possible. Acute and late (occurring >90 days after beginning RT) toxicities were defined and graded according to the Common Terminology Criteria for Adverse Events (CTCAE) v 3.0.⁸

Statistical methods

Descriptive statistics (mean, median, proportions) were calculated to characterize the patient, disease, and treatment features as well as toxicities after treatment. The overall survival (OS), disease-free survival, locoregional control, and metastases-free survival rates were estimated using the Kaplan–Meier product-limit method. All analyses

were performed using the Statistical Package for the Social Sciences ([SPSS] v 12.0; IBM Corporation, Armonk, NY, USA).

Results

Patient characteristics

Nine women were included. They had a median age of 68 years (range, 46–93 years) and belonged to FIGO Stage IIB to IVA (Table 1). The medium tumor volume was 41.6 (6.6–200.7) cm³. During WPRT, six received CCRT and three had RT alone. The medium length of cycles of chemotherapy was 6 weeks. All of the patients were treated with WPRT, with or without chemotherapy, followed by image-guided SBRT. A total dose of 70.4–78 Gy was given to seven patients (78%). Two patients received 65.4–66.4 Gy. The mean EQD2 of the tumor, rectum, bladder, and intestines were 76.0 ± 7.3, 73.8 ± 13.2, 70.5 ± 10.0, and 43.1 ± 7.1, respectively (Table 2). The most frequent reason for being unable to perform intracavitary treatment was inability to cannulate the cervical os (44%) and the second most common was medical unsuitability or contraindication (33%). One patient was unfit for intracavitary treatment because of the risk of anaphylactic shock in anesthesia. The other patient showed unusual anatomic configurations for the uterus due to myoma, and frequent contact bleeding.

Treatment outcome

The median survival was 13 months (range, 4–40 months). The actuarial 3-year OS, disease-free survival, locoregional control, and metastases-free survival rates were 46.9%, 25.9%, 77.8%, and 28.6%, respectively (Figure 1). Two stage IVA patients without concurrent chemotherapy had residual tumors after radiotherapy was completed. The others (7/9) had no locoregional recurrence upon confirmation of follow-up images and PAP. Four of nine (44%) patients experienced distant metastasis: Patient 1 had bone metastasis; Patient 5 had lung metastasis; Patient 7 had bone and lung metastasis; and Patient 9 experienced liver metastasis. Fifty-six percent of patients (5/9) were surviving at the time of this report.

Acute and subacute toxicity

Acute toxicity and late toxicity are detailed in Table 3. One patient presented with grade 3 diarrhea and another had grade 3 thrombocytopenia during treatment. There were no grade 3 toxicities for anemia, leucopenia, nausea, vomiting, genitourinary effects, or body weight loss. There were no grade 3 or 4 subacute toxicities for hematologic, genitourinary, or gastrointestinal effects. However, 3 patients

Table 1 Patients' characteristics and received techniques

Patient	Age	T	N	M	Stage	Tumor size (cm)	CCRT or RT alone	Cycles of C/T (wks)	Whole pelvic technique	Whole pelvic doses (cGy)/fractions	Cervical boost with SBRT Doses (cGy)/fractions	Total doses (cGy)	Total doses days	Reason for SBRT
1	78	2b	0	0	IIB	3.7 × 1.6 × 3.2	RT	–	IMRT	5040/28	2000/5	7040	65	To uncannulate the cervical os
2	68	2b	0	0	IIB	3.0 × 3.2 × 4.0	CCRT	6	HT	5040/28	2700/6	7400	81	Anaphylactic shock in anesthesia
3	53	2b	0	0	IIB	3.4 × 2.4 × 3.0	CCRT	4	HT	5040/28	1600/8	6640	82	To uncannulate the cervical os
4	49	2b	0	0	IIB	2.4 × 2.1 × 2.6	CCRT	6	IMRT	5040/28	1500/5	6540	81	To uncannulate the cervical os
5	77	3b	1	0	IIIB	6.4 × 5.6 × 11.2	CCRT	6	HT	5040/28	2700/6	7740	63	CVA with poor medical condition
6	46	3b	0	0	IIIB	5.1 × 5.4 × 8.9	CCRT	6	IMRT	5400/27	2400/6	7800	45	Myoma with contact bleeding
7	68	3a	1	0	IIIB	4.0 × 4.0 × 5.2	CCRT	5	HT	5000/25	2700/6	7700	79	To uncannulate the cervical os
8	93	4a	1	0	IVA	5.6 × 5.4 × 8.9	RT	–	HT	5040/28	2000/5	7040	75	Old age
9	56	4a	1	0	IVA	6.0 × 3.6 × 5.9	RT	–	HT	5040/28	2700/9	7740	86	Bed rest with poor medical condition

Abbreviations: T, primary tumor; N, regional lymph nodes; M, distant metastasis; CCRT, concurrent chemoradiation therapy; RT, radiotherapy; C/T, chemotherapy; SBRT, stereotactic body radiation therapy; HT, helical tomotherapy; IMRT, intensity-modulated radiotherapy; CVA, cerebral vascular accident.

Table 2 The equivalent dose in 2-Gy fractions (EQD2), using a linear-quadratic model and assuming α/β ratio = 10 for tumor and α/β = 3 for organs at risk (OARs)

Patient	Tumor EQD2	Rectum			Bladder			Intestines						
		Sum dose (Gy)	Mean doses	Maximal doses	Sum dose (Gy)	Mean doses	Maximal doses	Sum dose (Gy)	Mean doses	Maximal doses				
1	72.9	64.7	64.7	76.3	96.1	114.9	53.6	76.6	78.3	115.2	33.9	68.7	48.2	102.3
2	82.7	45.4	45.4	81.9	69.0	130.8	43.5	83.0	64.1	132.6	25.8	74.0	39.5	115.0
3	65.6	41.0	41.0	72.5	55.3	97.7	41.3	72.6	55.7	97.8	29.2	61.4	39.3	82.8
4	64.6	47.5	47.5	70.8	65.6	99.8	45.2	70.1	63.0	98.8	34.2	63.7	47.6	89.3
5	82.2	44.3	44.3	83.1	67.1	132.8	46.0	83.1	70.8	133.3	32.8	72.7	51.4	115.0
6	82.0	45.5	45.5	81.3	70.1	124.1	49.5	80.1	72.0	121.8	22.2	51.8	29.7	69.1
7	82.7	53.6	53.6	82.2	83.1	131.5	42.1	80.8	66.8	129.2	27.8	70.9	43.8	111.4
8	72.9	47.7	47.7	75.7	67.9	110.7	50.6	75.7	73.3	110.8	26.1	57.9	38.1	84.8
9	78.8	64.5	64.5	83.8	90.8	121.4	63.6	84.3	90.3	122.1	34.4	80.7	50.2	116.8
Median	78.8	47.5	47.5	81.3	69.0	121.4	46.0	80.1	70.8	121.8	29.2	68.7	43.8	102.3
Mean ± SD	76.0 ± 7.3	50.5 ± 8.7	50.5 ± 8.7	78.6 ± 4.9	73.8 ± 13.2	118.2 ± 13.3	48.4 ± 7.0	78.5 ± 5.0	70.5 ± 10.0	117.9 ± 13.4	29.6 ± 4.5	66.9 ± 9.0	43.1 ± 7.1	98.5 ± 17.5

Abbreviation: SD, standard deviation.

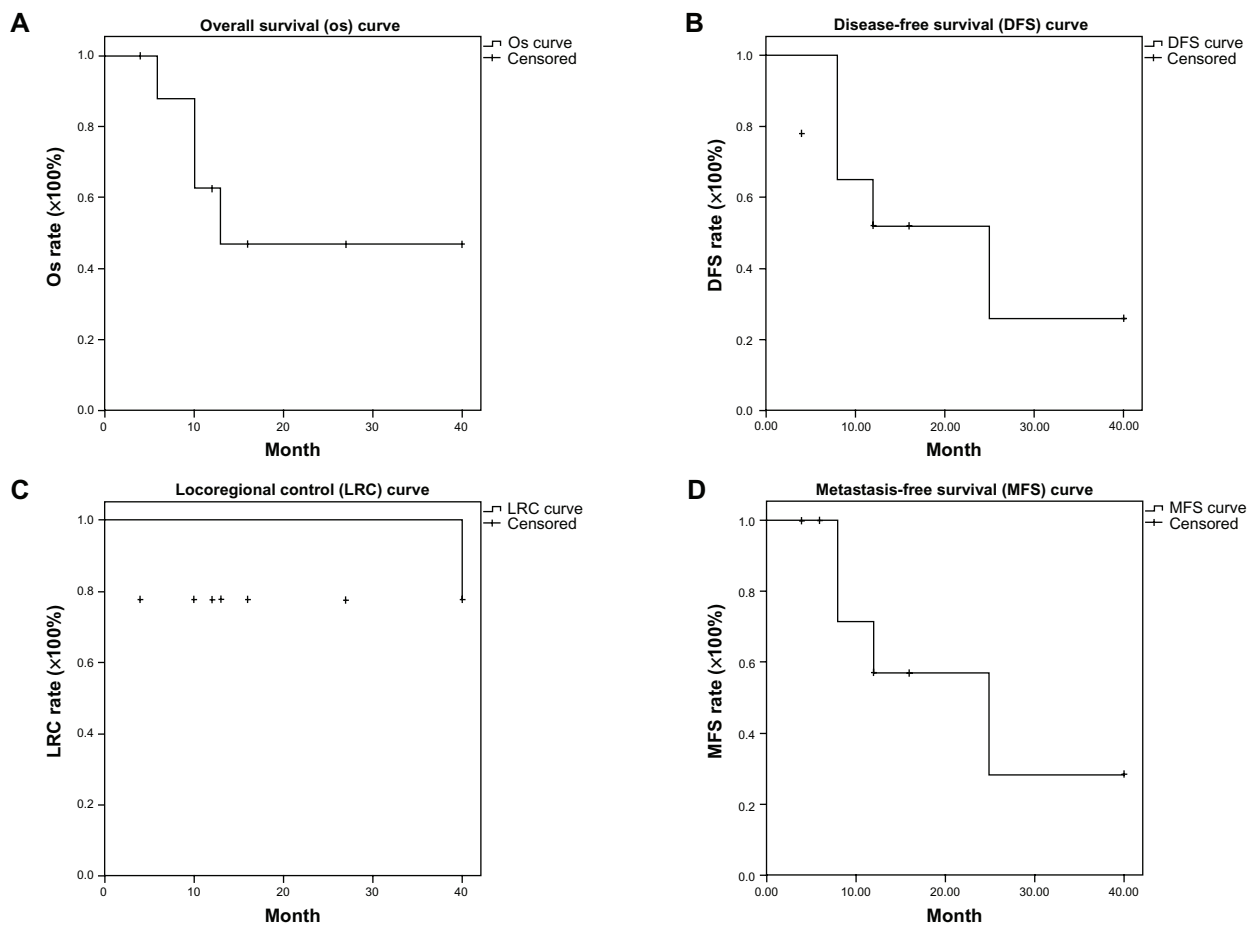


Figure 1 Locally advanced cervical cancer patients received whole pelvic radiotherapy concurrent with or without chemotherapy, followed by stereotactic body radiation therapy via helical tomotherapy. **(A)** Overall survival curve; **(B)** disease-free survival curve; **(C)** locoregional control curve; and **(D)** metastases-free survival curve.

suffered from manageable rectal bleeding in months 11, 14, and 25, respectively. One stage IVA patient experienced fistula formation in month 3.

Discussion

There are a variety of situations in which BT cannot be carried out, such as difficult cannulation of the cervical os, anesthetic comorbidity,¹ or unusual anatomic configuration.⁵ In these situations, EBRT could be an alternative to BT in cervical cancer patients.^{1,3,5}

Where EBRT has been used to replace BT, local recurrence has been documented in 6.6%–65% of cases, with most occurring within 3 years. Ferreira et al⁹ reported that the locoregional failure rates for use of EBRT as replacement of BT was 65%, which was inferior to EBRT followed by BT (49%). In a 1991 report,¹⁰ EBRT plus BT had a lower local failure rate (41%) than EBRT alone (67%), for stage IIIB cancer of the uterine cervix. However, other evidence showed that results of EBRT alone were comparable to the results of the combination of BT and external irradiation. In the report

by Akine et al,¹¹ the local control rate for stage IIIB patients treated with EBRT was 19%. Montana et al¹² obtained better relapse-free survival for the combination therapy groups at 2 years (61% vs 36%), but this difference was not sustained beyond 5 years. Mollà et al³ also confirmed the concept of using IMRT to deliver a final boost, and this approach might well be considered an acceptable alternative to BT. In our previous case report of pathology findings,⁵ replacement of BT with image-guided SBRT via HT also resulted in a disease-free state, without local failure. Barraclough et al¹ reported that the 3-year, cancer-specific OS rate was 49% for patients who were treated with EBRT and that the local recurrence rate was 41%. Ulmer et al¹³ used external irradiation alone in 119 patients with stage III tumors and obtained a 5-year survival rate of 30.3%. In the current study, the actuarial 3-year OS and local recurrence rates were 47%, and 22%, respectively, and with the exception of the two patients with stage IV who had residual tumors after treatment, seven of the nine (78%) patients whose treatment was delivered by SBRT had no local failure. The current results suggest the

Table 3 Acute and late toxicity for patients with locally advanced cervical cancer who received whole pelvic radiotherapy concurrent with or without chemotherapy, followed by stereotactic body radiation therapy via helical tomotherapy

Toxicity*	Patients, n (%)						
	Acute toxicity						
	Nausea/ vomiting	Diarrhea	Genitourinary effects	Body weight loss	Anemia	Leukopenia	Thrombocytopenia
Gr 1	9 (100)	7 (78)	8 (89)	10 (100)	8 (89)	8 (89)	6 (67)
Gr 2	0	1 (11)	1 (11)	0	1 (11)	1 (11)	2 (22)
Gr 3	0	1 (11)	0	0	0	0	1 (11)
Gr 4	0	0	0	0	0	0	0
Gr 5	0	0	0	0	0	0	0
Patient	Late toxicity						
	Rectal bleeding		Genitourinary effects		Fistula formation		
1	Grade 2 rectal bleeding Occurred in month 25		No occurrence until present (26 months)		No occurrence until present (26 months)		
2	No occurrence until present (16 months)		No occurrence until present (16 months)		No occurrence until present (16 months)		
3	No occurrence until present (12 months)		No occurrence until present (12 months)		No occurrence until present (12 months)		
4	No occurrence until present (4 months)		No occurrence until present (4 months)		No occurrence until present (4 months)		
5	Grade 2 rectal bleeding Occurred in month 11		No occurrence prior to expiration (13 months)		No occurrence prior to expiration (13 months)		
6	Grade 2 rectal bleeding Occurred in month 14		No occurrence until present (40 months)		No occurrence until present (40 months)		
7	No occurrence prior to expiration (10 months)		No occurrence prior to expiration (10 months)		No occurrence prior to expiration (10 months)		
8	No occurrence prior to expiration (6 months)		No occurrence prior to expiration (6 months)		No occurrence prior to expiration (6 months)		
9	No occurrence prior to expiration (10 months)		No occurrence prior to expiration (10 months)		Occurred in month 3		

Notes: *The grade of toxicity followed the Common Terminology Criteria for Adverse Events (CTCAE) v 3.0.⁸

possibility of replacing BT with image-guided SBRT via HT for those patients who are unsuitable for BT treatment.

Favorable outcomes, with pelvic control rates of 69%–76%, were presented^{14–16} with the BED 65–100.9 Gy10 (EQD2 = 54.2–84.1 Gy) at point A, with a combination of EBRT and high-dose rate BT for an advanced stage tumor of the uterine cervix. Ito et al¹⁴ suggested that a 76% pelvic control rate could be achieved with the BED 65–90.4 Gy10 (EQD2 = 54.2–75.3 Gy) at point A. The University of Wisconsin also shared their experiences with the BED 85–100.9 Gy10 (EQD2 = 70.8–84.1 Gy) at point A, showing that 71% of 3-year pelvic control rates could be achieved.¹⁵ Similarly, Toita et al¹⁶ also confirmed that BED 70–80 Gy10 (EQD2 = 58.3–66.7 Gy) at point A could provide an impressive 3-year pelvic control rate (76%) for advanced stage diseases. In the current study, the range of BED and the mean doses of BED for tumors were 77.5–99.2 Gy10 (EQD2 = 64.6–82.7 Gy) and 91.2 ± 8.8 Gy10 (EQD2 = 76.0 ± 7.3 Gy), respectively. The 3-year locoregional control was 78%, which suggested the BED at

77.5–99.2 Gy10 (EQD2 = 64.6–82.7 Gy) might be adequate to control advanced cervical cancer in patients who cannot receive BT and in whom BT is replaced with image-guided SBRT. However, this should be evaluated with caution, since the number of cases is still limited and not all the patients in the current study were receiving CCRT.

Several investigators have analyzed probability of late complications as a function of total BED at the ICRU 38 reference points. Ranging from 97–169 Gy3 (EQD2 = 80.8–140.8 Gy), with median or mean doses for the BED of the rectum, the late complication rate was 11%–52%.^{7,16–18} Ogino et al¹⁷ demonstrated that the incidence of rectal complication was correlated with BED and suggested that rectal complications could decrease to less than 10% when BED does not exceed 146 Gy3 (EQD2 = 121.7 Gy). The data from Clark et al¹⁸ showed that the rectum BED, without the development of complications, for the CCRT and RT alone groups was 162 Gy3 (EQD2 = 135 Gy) and 125 Gy3 (EQD2 = 104.2 Gy), respectively. Similarly, Toita et al¹⁶ also suggested that the cumulative BED at the rectal reference point should be kept

below 100–120 Gy₃ (EQD2 = 83.3–100 Gy) to prevent rectal complication. Cheng et al⁷ found that patients with a total maximal proximal rectal BED more than 110 Gy (EQD2 = 91.7 Gy) presented with a significantly increased frequency of Grade 2 or greater rectal complications.

Except for BED of the rectum, the total doses to the rectum have also contributed to complications. Perez et al¹⁹ and Pourquier et al²⁰ reported that with doses below 75–80 Gy delivered in limited volumes, the incidence of grade 2 and 3 complications was less than 5%; however, with higher doses, the incidence of complications increased to 10%–15%. Cheng et al⁷ recommended a proximal rectal dose < 62 Gy of a direct dose sum from WPRT and BT, to avoid an increased frequency of grade 2 or greater rectal complications. In the current study, the minimal values of mean and maximal doses of rectum for these rectal bleeding patients were 80.5 Gy₃ and 137 Gy₃ for BED (EQD2 = 55.3 and 97.7 Gy, respectively) and 44.3 Gy and 81.3 Gy for sum doses, respectively (Table 2). Three of the nine patients experienced manageable rectal bleeding. Toxicities are of concern and may pose a limitation for this technique. Obviously, an analysis based only on mean and maximal BED or sum doses at the rectum might be insufficient to draw conclusions; nevertheless, this should be considered and stimulate further improvement in the techniques of SBRT to decrease rectal complications, so that this can be an alternative to BT.

Calculated total BED at the bladder reference point had no significant correlation with the incidence of bladder complications.¹⁶ In high-dose radiation following a course of conventionally fractionated radiation (50.4 Gy) without toxicity, the maximal limitation for bladder and intestines recommended by RTOG 9708 were 25 and 10 Gy, respectively.²¹ In the current study, neither complication of the bladder, nor of the intestines, was noted. The maximal mean dose of the bladder and intestines for sum dose and BED Gy₃ were 63.6 and 34.4 Gy, 108 and 61.7 Gy₃ (EQD2 = 90.3 and 51.4), respectively. The maximal mean dose of the bladder and intestines after WPRT were 16.6 and 13.0 Gy, respectively. For the bladder and intestines, an analysis of the sum dose and maximal mean dose after conventionally fractionated radiation might be a way to evaluate the treatment plan, to avoid complications when replacing BT with SBRT via HT.

There are some limitations in our current study. First, because of the number of cases and retrospective study design, no statistical conclusions can be drawn. Second, the follow-up period was short, so long-term results and close monitoring are further required. Third, it is not easy to confirm that the sum of maximal doses for bladder, intestine, and

rectum are coming from the same points, although we tried to use the concepts of BED to provide a means for judging the correlation between doses and toxicities. Fourth, not all the patients had implanted fiducial markers and so the radiotherapy margin could not be reduced effectively, even with image-guided technique.

Conclusion

The present study effectively used image-guided SBRT via HT to provide impressive results for cervical carcinoma patients with contraindications to BT. The proposed technique may be considered an acceptable alternative to BT, by further improving on techniques and its consideration of both mean and maximal BED, and sum doses at the rectum, to avoid rectal complications. Long-term follow up is needed to confirm these preliminary findings.

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Disclosure

The authors report no conflicts of interest in this work.

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