PET data obtained at a gantry angle of 90°. The measurement time for each gantry angle is expected to be only a few minutes; therefore, the effects on the acquired parallel-plane PET data of metabolic changes in the body or source activity are negligible.

Additionally, this high spatial resolution will likewise be useful in hypoxic region imaging using FMISO because the hypoxic region distribution in the tumor is complex (Nehmeh *et al* 2008).

5. Conclusion

We performed a basic study to determine the accuracy of image registration using a PET-based molecular image guided method. Planar images were reconstructed from parallel-plane PET data to obtain the PET-based digitally reconstructed planar image (PDRI) used in the registration. In-plane PDRIs had higher resolution and therefore usable for image registration. Phantom experiments using in-plane PDRIs showed that there is no significant difference between radiographic and PDRI registrations. Our results suggest that m-IGRT image registration using PET-based reconstructed planar images along the in-plane direction is feasible for clinical use. Furthermore, the system will provide additional information for image registration when bony structures cannot be recognized with radiographic registration methods.

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DISEASES OF THE ESOPHAGUS

Original article

Submucosal tumor appearance is a useful endoscopic predictor of early primary-site recurrence after definitive chemoradiotherapy for esophageal squamous cell carcinoma

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SUMMARY. **Chemoradiotherapy (CRT) for esophageal cancer is disadvantageous because of a high locoregional failure rate. Detecting early small recurrent cancers at the primary site is necessary for potential salvage treatment. However, most endoscopists are inexperienced and therefore, a role for surveillance endoscopy after complete remission (CR) has not been established. We retrospectively evaluated serial surveillance endoscopic images from patients eventually proved to have primary-site recurrence in order to identify useful endoscopic features for early diagnosis. From January 2000 to December 2004, 303 patients with esophageal squamous cell carcinoma underwent definitive CRT, and 133 of them achieved CR. The surveillance endoscopic images stored at intervals of 1–3 months for the 16 patients with recurrence only at the primary tumor site and the 61 patients with no recurrence were collected for reexamination. Among 133 patients who achieved CR, 16 (12%) developed only local recurrence at the primary site. Thirteen of the 16 primary-site recurrent tumors (81%) appeared as submucosal tumors (SMT), with the remaining appearing as erosions or mild strictures. Of biopsy-proven recurrences, 81% were preceded by newly developed lesions such as SMT, erosions, or mild strictures detected by earlier surveillance endoscopies. For all 77 patients achieving CR with no metastasis, 86% of the evolving SMT with negative biopsies were eventually confirmed as cancer at later endoscopies. Thirteen of the 21 evolving lesions were subsequently confirmed as recurrent cancer. Early primary-site recurrence of esophageal cancer after a complete response to CRT is detectable with frequent endoscopic surveillance. SMT appearance is a useful endoscopic sign of early recurrence, as well as a predictor of subsequent diagnosis of recurrence.**

KEY WORDS: **chemoradiotherapy, esophageal cancer, recurrence, surveillance.**

INTRODUCTION

Definitive chemoradiotherapy (CRT) is widely accepted as a standard treatment option in the management of locally advanced esophageal cancer because of its high response rate and significant survival benefit.[1,2] A major drawback to this nonsurgical approach is locoregional treatment failure. At least 40% of patients undergoing CRT experienced local failure, some of whom did not develop distant metastases.[1,3–5]

These primary-site recurrence patients are traditionally managed with salvage esophagectomy for a chance of long-term survival, particularly in those with an earlier pathological stage (T1N0 and T2N0).[6,7] However, high perisurgical mortality and morbidity rates are major concerns.[7,8] Recently developed nonsurgical techniques, such as salvage endoscopic mucosal resection and photodynamic therapy,

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have the advantages of greater safety and fewer treatment-related sequelae, while conferring promising survival benefits for local failures after definitive CRT.[9,10] Technically, endoscopic mucosal resection and photodynamic therapy are feasible only when the volume of the locally recurrent tumor is small enough to be amenable to these endoscopy-based procedures. Therefore, the application of these newer treatments depends crucially on the ability to identify early recurrent tumors by endoscopy.

A strategy of frequent surveillance endoscopy initiated early after remission of the cancer should theoretically improve the chances of detecting primary-site recurrent tumors in their early stages. This requires the prompt recognition of minute tumors arising from the former neoplastic bed, instead of from the uninvolved normal esophageal mucosa. However, the complete regression of cancer cells results in residual fibrosis, radiation-induced tissue injury, and the distortion of normal microstructures,[11,12] which may render relapsing neoplastic growth morphologically different from typical primary tumors. Apparently, most endoscopists are inexperienced in hunting for these difficult lesions. To our knowledge, no study of the skills in endoscopic detection of such lesions has been published. Not surprisingly, a follow-up endoscopy after the completion of CRT is considered 'optional' in the National Comprehensive Cancer Network clinical practice guidelines for esophageal cancer.[13] We believe that a reliable endoscopic diagnostic technique is necessary to support a strategy of intense endoscopic follow-ups.

As a cancer referral and research hospital, our institute is unique in its implementation of a vigorous endoscopic follow-up program after primary treatment for all patients with esophageal cancer. Therefore, it is possible to analyze the filed imaging data of endoscopic monitoring on the post-CRT mucosa. In the present study, we aimed to identify useful endoscopic findings through reviewing the image data pool to predict recurrent esophageal cancers limited to the primary site after complete remission (CR) is achieved by CRT.

MATERIALS AND METHODS

Patient population

Between January 2000 and December 2004, 303 patients with esophageal squamous cell carcinoma underwent definitive CRT at the National Cancer Center Hospital East, Kashiwa, Japan. The CRT consisted of 50.4–60 Gy irradiation, together with two cycles of continuous infusion with 5-fluorouracil (5FU) and cisplatin. Up to four courses of CRT were added for those patients who showed a good initial response to treatment.[9]

Table 1 Clinical data of 133 patients achieving complete remission with definitive chemoradiotherapy

Characteristic		Number of patients	%
Sex			
Male		110	82.7
Female		23	17.3
Age (years)			
Mean	62		
Range	39–76		
T stage			
T1		30	22.6
T2		21	15.8
T3		70	52.6
T4		12	9.0
N stage			
N0		46	34.6
N1		87	65.4
M stage			
M0		123	92.5
M1		10	7.5
Clinical stage			
I		16	12.0
II		45	33.8
III		62	46.6
IV		10	7.5
Macroscopic classification			
Type 0		30	22.6
Type 1		19	14.3
Type 2		60	45.1
Type 3		24	18.0

Response to treatment was assessed at the completion of CRT. CR was defined when all the following criteria were met: (i) the disappearance of the tumor lesion or ulcer at the primary site, with negative biopsies; (ii) no esophageal stricture or any condition that prevented a thorough endoscopic examination of the whole esophagus; (iii) no remaining measurable disease or distant metastasis on computer tomography and chest roentgenography; and (5) these criteria were met for at least 4 weeks.

Of the 303 patients, 133 (43.9%) were defined as being in CR at the completion of CRT. Of these 133 patients, 110 were men, with a median age of 62 years. Pretreatment staging of their esophageal cancers was determined with the tumor-node-metastasis classification of the International Union Against Cancer.[14] Seventy (52.6%) patients had T3 tumors; most patients had N1 (65.4%) or M0 (92.5%) disease. Forty-five (33.8%) and 62 (46.6%) patients were classified as clinical stages II and III, respectively (Table 1).

Study design

After achieving CR, initial follow-up endoscopy to confirm CR was scheduled within at most 1–2 months for each patient, accompanied with other necessary studies for the assessment of metastases. After the confirmation of CR, follow-up endoscopy was scheduled every 2–3 months for the first year and every 4–6

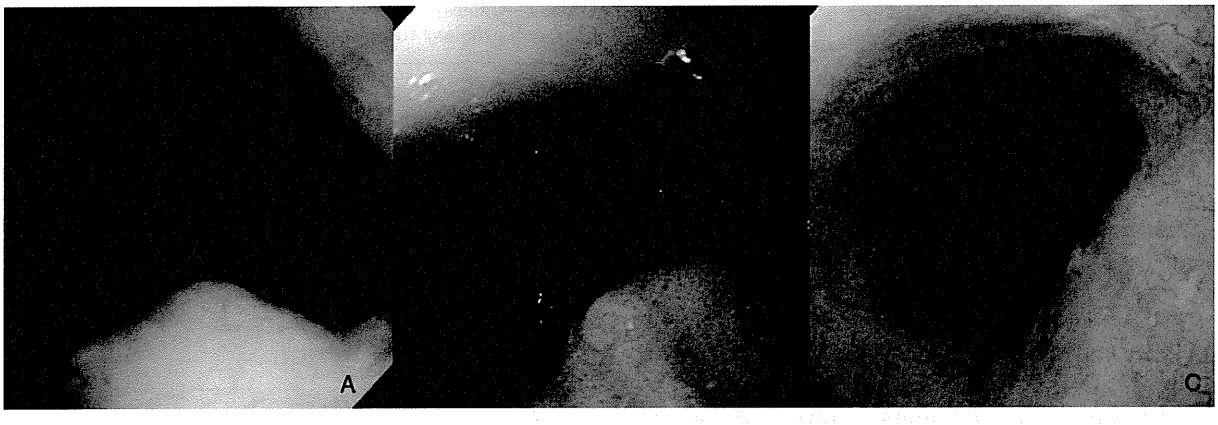

Fig. 1 Initially growing recurrent esophageal cancer at the primary tumor site after complete remission was achieved with chemoradiotherapy may be detected by endoscopy, with features of a submucosal tumor (A), a submucosal tumor with superficial ulcer (B), or a flat erosion (C).

months for 2 years thereafter. Lugol staining and multiple biopsies at the primary site were routinely required.[15] The diagnosis of local recurrence was determined by a positive biopsy.

Of the 133 CR patients, 61 had no recurrence, 56 developed lymph node or distant metastases, and the remaining 16 developed local recurrence at the primary tumor site with no evidence of metastasis. We excluded the 56 patients with lymph node or distant metastases from this study because for them, evaluation of the primary site was not important and only those patients eligible for salvage treatment on local tumors were of interest. Therefore, the endoscopic images of the remaining 77 patients were retrospectively enrolled. This population comprised patients with esophageal squamous cell carcinoma who achieved CR after the initial CRT and developed no metastasis during follow-up, regardless of local recurrence. All of the filed endoscopic images stored after achieving CR, both conventional endoscopy and Lugol-stained chromoendoscopy, were retrospectively collected for reexamination. The stored endoscopic images were evaluated by consensus among three endoscopists experienced in upper gastrointestinal cancer diagnosis (K. T., M. M., K. M.).

RESULTS

Upon the diagnosis of primary-site recurrence for the 16 patients, 13 (81%) had endoscopic findings resembling submucosal tumors (SMT), typically a focal bulge mostly covered by normal-appearing mucosa (Fig. 1A).[16] Eleven of the 13 tumors contained central eroded areas recognized as ulcers or erosions (Fig. 1B and 1C). The remaining three tumors were detected as flat erosions without features of SMT (Table 2).

Images of surveillance endoscopies performed at intervals between CR and the diagnosis of recurrence in the 16 patients were sequentially examined. Newly developed gross lesions at the primary site with negative biopsies were interpreted as recurrent lesions. Evolving lesions were discovered in 13 (81%) patients, including six (38% of the 16 patients) SMT, five (31%) erosions, and two (12%) mild luminal strictures (Table 3).

For all 77 patients achieving CR and free of metastasis, lesions newly developed between CR and the most recent endoscopic surveillance were considered evolving lesions. Therefore, an evolving lesion may be eventually proven to be a recurrence or remain biopsy-negative at the most recent endoscopy. Six of the seven (86%) evolving SMT were subsequently confirmed as recurrent cancer by follow-up

Table 2 Endoscopic findings at primary-site with biopsy-proven recurrence

Endoscopic finding	Number of patients	%
SMT	13	81
SMT with erosion or ulceration	11	
SMT without erosion or ulceration	2	
Erosion	3	19
Total	16	100

SMT, submucosal tumor.

Table 3 Endoscopic findings of newly developed lesion for primary-site recurrent tumors

Preceding newly developed lesions with negative biopsies	Findings at diagnosis of recurrence	Number of patients
SMT	SMT	6
Erosion	SMT	4
Erosion	Erosion	1
Mild stricture	SMT	2
No newly developed lesion	SMT	1
No newly developed lesion	Erosion	2
Total		16

SMT, submucosal tumor.

Table 4 Primary-site biopsy results of the latest surveillance endoscopy for patients who achieved complete remission and remained free of metastasis

Evolving lesion found at preceding endoscopies	Numer of patients (%)	Biopsy result of the latest endoscopy	Number of patients (%)
SMT	7 (9)	Recurrence	6 (86)
		Negative	1 (14)
Erosion	8 (10)	Recurrence	5 (63)
		Negative	3 (37)
Mild stricture	6 (8)	Recurrence	2 (33)
		Negative	4 (67)
No evolving lesion	56 (73)	Recurrence	3 (5)
		Negative	53 (95)
Total	77 (100)		

SMT, submucosal tumor.

endoscopic biopsies. Similarly, five of eight (63%) evolving erosions and two of six (33%) evolving mild strictures were finally confirmed as recurrence. Fifty-six patients were never found to have evolving lesions throughout the follow-up, including three (5%) who were confirmed as recurrence upon the first appearance of an endoscopic lesion. In total, eight of the 21 (38%) patients who developed evolving lesions remained biopsy-negative at their most recent endoscopic follow-up (Table 4).

DISCUSSION

We discovered that the most frequent (81%) endoscopic indicator of primary-site recurrence at its earliest possible stage for a histological diagnosis is SMT. Eighty-one percent of biopsy-proved recurrences were preceded by newly developed lesions such as SMT, erosions, or mild strictures detectable with surveillance endoscopies. Most (86%) evolving SMT with negative biopsies were eventually confirmed as cancer at later endoscopies, but the proportions were lower for other evolving lesions such as erosions (63%) and strictures (33%). This is the first study to describe the morphological changes of early recurring tumors by serial endoscopic observations at short intervals. Our findings will be helpful for improving the skills to detect potentially treatable primary-site recurrence after definitive CRT for esophageal squamous cell carcinoma.

For the endoscopic diagnosis of primary esophageal cancer, several features have been previously described to detect early stage squamous cell carcinoma: localized mucosal erosions in contrast to normal surrounding mucosa; circumscribed mucosal protuberances with irregular configurations; focal areas of mucosal coarsening and congestion; and, rarely, white mucosal plaques.[16] However, these features are not reliable when applied to early recurrent tumors arising from the mucosal bed of a former primary cancer that regressed after CRT. The original esophageal layering and vascular structures have been disrupted by the primary tumor. Furthermore, the expansion and arrangement of recurring neoplastic cells are disrupted by tissue reactions to previous chemotherapy and radiotherapy, as well as by subsequent repair processes. Tumor necrosis, foam cell formation, vascular granulation, inflammatory exudation, and fibrosis are frequent histological sequelae of CRT.[17,18] The minute foci of the initial neoplastic growth may arise from scattered residual cancer cells in deeper tissues, rather than from the superficial mucosal layer, as does the primary cancer.[11] These factors have largely precluded endoscopic ultrasound as a feasible tool in the assessment of residual or recurrent esophageal cancers.[19,20] For the same reason, the endoscopic diagnostic features for recurrent tumors are likely to be different from those for primary tumors.

We speculate that most of the SMT lesions discovered in our study were formed by expanding tumor cells in the submucosal layers, but barely reached the luminal surface because of their depth and constraining fibrosis. Although the overlying mucosa appeared normal, they manifest their first sign by bulging outward. Malignant cells can be captured by biopsy forceps only when they reach the surface in sufficient numbers, or more efficiently, destroy the surface to make an erosion. This might explain why all of the six newly developed SMT yielded negative results at their first biopsies but eventually proved to be recurrences (Table 3).

Several previous studies have aimed to improve the detection of local recurrence by measures other than endoscopy. In addition to pretreatment staging, F-18-fluorodeoxyglucose–positron emission tomography (FDG–PET) is highly sensitive (up to 96%) in detecting recurrent esophageal cancer, but with somewhat lower specificity (68–82%).[21–23] However, its utility in detecting locoregional recurrence is limited by its low specificity (57–75%) for postesophagectomy patients. Postsurgical inflammation and anatomical changes are largely responsible for the false positivity. Detecting small residual or early recurrent cancers is even more challenging because low tumor volume could greatly reduce the sensitivity of FDG–PET. Moreover, such lesions are not distinguishable from post-CRT inflammation or regional lymph-node metastasis.[24,25]

The results of our study disagree with the conventional belief that endoscopy is of limited utility in the management of esophageal cancer after CRT.[13,26] We believe that routine endoscopy, particularly focused on the primary tumor site, is advisable for all patients with esophageal squamous cell carcinoma after the completion of CRT. We also suggest regular endoscopic surveillance at least every three months for those who have achieved CR. The occurrence of

SMT-like lesions after CR is an alarming sign that deserves intensive investigation and follow-up if a modality of salvage treatment is available. Any evolving lesion at the primary site with negative biopsy should be followed closely.

Our retrospective study design has introduced a knowledge bias because the evaluating endoscopists were not totally blinded to the outcomes. Therefore, a randomized controlled trial comparing the clinical outcomes is necessary to establish the role of surveillance endoscopy after definitive CRT for esophageal squamous cell carcinoma.

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CLINICAL INVESTIGATION

PROTON BEAM THERAPY AS A NONSURGICAL APPROACH TO MUCOSAL MELANOMA OF THE HEAD AND NECK: A PILOT STUDY

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Purpose: The aim of this pilot study was to assess the clinical benefit of proton beam therapy for mucosal melanoma of the head and neck.

Methods and Materials: Patients with mucosal melanoma of the head and neck with histologically confirmed malignant melanoma and N0 and M0 disease were enrolled. Proton therapy was delivered three times per week with a planned total dose of 60 Gy equivalents (GyE) in 15 fractions.

Results: Fourteen consecutive patients were enrolled from January 2004 through February 2008. Patient characteristics were as follows: median age 73 years old (range, 56 to 79 years); male/female ratio, 7/7; and T stage 1/2/3/4, 3/2/0/9. All patients were able to receive the full dose of proton therapy. The most common acute toxicities were mucositis (grade 3, 21%) and mild dermatitis (grade 3, 0%). As for late toxicity, 2 patients had a unilateral decrease in visual acuity, although blindness did not occur. No treatment-related deaths occurred throughout the study. Initial local control rate was 85.7%, and, with a median follow-up period of 36.7 months, median progression-free survival was 25.1 months, and 3-year overall survival rates were 58.0%. The most frequent site of first failure was cervical lymph nodes (6 patients), followed by local failure in 1 patient and lung metastases in 1 patient. On follow-up, 5 patients died of disease, 4 died due to cachexia caused by distant metastases, and 1 patient by carotid artery perforation cause by lymph nodes metastases.

Conclusions: Proton beam radiotherapy showed promising local control benefits and would benefit from ongoing clinical study. © **2010 Elsevier Inc.**

Proton beam therapy, Mucosal melanoma, Head and neck.

INTRODUCTION

Although rare worldwide, mucosal melanoma of the head and neck is relatively common in Japan (1). Most reports to date have described small series of patients over long time periods but have not led to any consensus in the approach to treatment. A surgical approach incorporating postoperative radiotherapy has been recognized as a community standard, and the 5-year survival rate of head and neck mucosal melanoma varies from 20% to 45% (2–5). This surgical approach is often complicated by serious cosmetic and functional deformity, and, particularly for nasal and sinonasal mucosal melanoma, satisfactory surgical clearance is often markedly difficult to obtain.

Several reports have described the use of radiotherapy alone for mucosal melanoma of the head and neck, with 5-year survival rates slightly less than those of the surgical approach (6–8). Regarding radiotherapy, The review by Trotti *et al.* (9) of four reports of radiotherapy for mucosal melanoma showed 3-year local control rates of 36% to 61%. In Japan Wada *et al.* (10) recently reported a series of 66 cases of mucosal melanoma of the head and neck, 21 of whom were treated with radiotherapy as the main modality. The rate of complete response in these 21 cases was 29%, and the 3-year disease-specific survival rate was 33%. Since X-ray irradiation has a limitation of dose distribution for tumor areas in proximity to organs at risk, like optic nerve and brain stem, it is often difficult to give enough dosage to planned target volume.

Proton beam therapy (PBT) is characterized by rapid fall-off at the distal end of the Bragg peak and a sharp lateral penumbra, depending on the energy, depth, and delivery (11).

Because of its physical characteristics, PBT provides better dose distribution than X-ray irradiation. PBT is deemed a feasible and effective treatment modality that provides curative high-dose irradiation to the tumor volume without increasing normal tissue toxicity. However, the use of PBT

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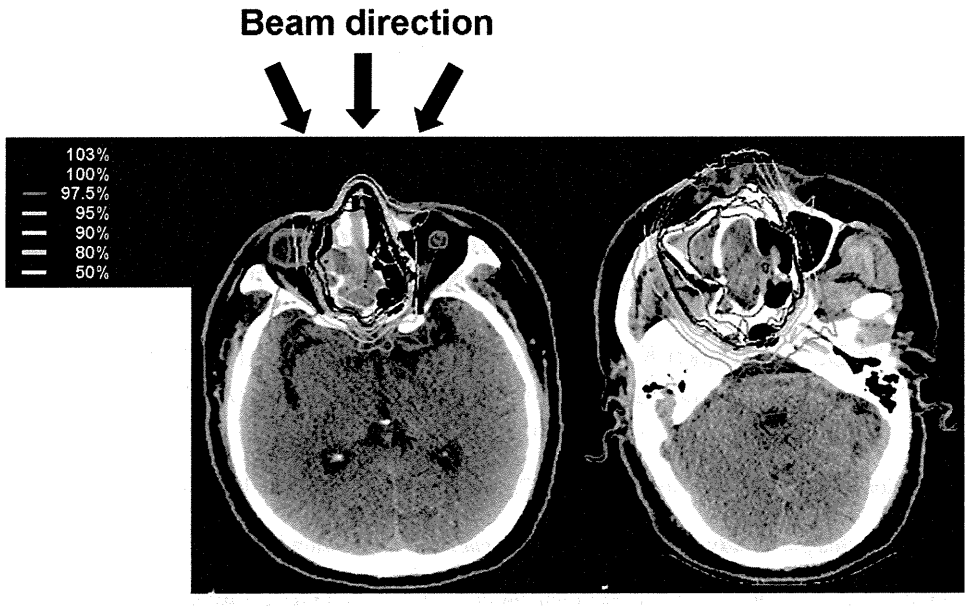

Fig. 1. Target volume and beam arrangement. GTV was defined as the gross tumor lesion determined with pretreatment CT/MRI and PET. CTV was defined as the region of the gross tumor lesion and adjacent sinuses. PTV was basically set as CTV plus 3-mm margin, with acceptance of fine-tuning to the PTV in consideration of organs at risk. Irradiation dose and volume for organs at risk were usually minimized by using a noncoplanar three-field technique.

for mucosal melanoma of the head and neck has not been reported. Here, we conducted a pilot study to examine the utility of hypofractionated PBT as a newly developed treatment modality for mucosal melanoma of the head and neck.

METHODS AND MATERIALS

Patients

Entry criteria for this retrospective study were (*1*) pathologically proven mucosal melanoma of the head and neck; (*2*) clinical TNM status of N0M0; (*3*) Eastern Cooperative Oncology Group (ECOG) performance status of 2 or less; (*4*) adequate organ function; and (*5*) no active concomitant malignancy. This treatment was approved by the institutional review board of the National Cancer Center Hospital, and written informed consent to treatment was obtained from all patients before the initiation of treatment.

Pretreatment clinical evaluation was performed using magnetic resonance imaging (MRI); cervical, chest, and abdominal computed tomography (CT); and/or positron emission tomography-CT (PET-CT). Radiological evaluations for staging were jointly reviewed by radiologists, surgeons, and oncologists at our institution. In the present study, all diseases were staged with the International Union Against Cancer criteria for carcinoma of the nasal cavity or paranasal sinus (12).

Table 1. Patient characteristics

Characteristic	Parameter	No. of patients (n = 14)
Age	Median (range)	73 (56-79)
Gender	Male/female	7/7
Performance Status	0 to 1/2	14/0
Primary site	Nasal cavity	11
	Paranasal sinus	3
TNM stage	T1N0M0	3
	T2N0M0	2
	T3N0M0	0
	T4N0M0	9

Treatment

PBT was delivered three times per week for a planned total dose of 60 Gy equivalents (GyE) in 15 fractions using a 150- to 190-MeV proton beam. The biologically equivalent dose (BED) using a linear-quadratic model is defined as BED = nd [1+ d/1/(α/β)], where n is the fractionation number, d is the daily dose, and the α/β ratio was 2.5 ($Gy_{2.5}$) for malignant melanomas (6). When n = 15 and d = 4 were substituted, BED was 156 $Gy_{2.5}$.

Treatment planning was performed with a three-dimensional CT planning system. In this system, the proton beam was generated with a Cyclotron C235 with an energy of 235 MeV at the exit. Relative biologic effectiveness was defined as 1.1, based on our preclinical

Table 2. Adverse events

	No. of patients with toxicity grade shown*				
Toxicity	1	2	3	4	% 3-4
Dermatitis	7	5	0	0	0
Mucositis	9	2	3	0	21
Infection	0	0	0	0	0
Hearing loss	1	0	0	0	0
Neuropathy					
CN-II	0	0	2	0	12
CN-V	0	0	0	0	0
Keratitis	0	2	0	0	0
Memory impairment	0	0	0	0	0

Treatment-related death: 0%.

* Using Common Terminology Criteria for Adverse Events vesion.3.0.

experiments (13). PBT at our institution is passive irradiation with dual-ring double-scatter methods. Dose distribution was optimized using the spread-out Bragg peak method and obtained using a broad-beam algorithm.

Gross tumor volume (GTV) was determined with pretreatment CT, MRI, and/or PET-CT. The clinical target volume (CTV) was defined as the GTV plus a 5-mm margin and sinuses adjacent to GTV. In cases with brain invasion, the area of T_2-weighted prolongation on MRI was also included in the CTV. The planning target volume (PTV) was basically defined as the CTV plus a 3-mm margin but could be finely adjusted where necessary in consideration of organs at risk. The beam energy and spread-out Bragg peak were fine-tuned such that the PTV encompassed a 90% isodose volume of the prescribed dosage. Irradiation dose and volume for organs at risk was usually minimized using a noncoplanar three-field technique (Fig. 1).

Dose constraints for organs at risk at 4 GyE per fraction were (*1*) surface of brainstem, 45 GyE; (*2*) center of brainstem, 33 GyE; (*3*) optic nerves of the healthy side/chiasm, 42 GyE; and (*4*) optic lens, 13 GyE.

To evaluate the risk of radiation-induced complications in normal tissue, dose–volume histograms were calculated for all patients. Patients were immobilized with custom-made immobilization devices that provided high reproducibility at every treatment fraction. Patient setup was verified before the delivery of each fraction, using a digital radiography subtraction system.

Evaluation of toxicity and efficacy

Toxicities were graded using the Common Terminology Criteria for Adverse Events (CTCAE) version 3.0. Weekly follow-up was continued until acute toxicity was easily manageable, and posttreatment MRI was performed at 6 to 10 weeks after the end of PBT to rule out treatment-induced empyema and brain necrosis. To confirm local control, MRI was performed every 3 to 6 months after the end of treatment, and distant metastases were assessed by CT/PET-CT. The achievement of initial local control was confirmed when all of the following criteria were fulfilled: (*1*) patients were alive at 1 year after the initiation of treatment; (*2*) no progressive disease was detected at the primary site for 1 year; and (*3*) no recurrence was detected at the primary site for 1 year.

Statistical analysis

Overall survival time was calculated from the start of treatment to the date of death or last confirmed date of survival. Survival time was censored at the last confirmed date of survival if the patient was alive. Progression-free survival (PFS) time was defined from the day of initiation of treatment to the first day of confirmation of progressive disease at any site or any cause of death. Overall survival time, PFS time, and local control period were estimated using the Kaplan-Meier product-limits method.

RESULTS

Patient characteristics

Fourteen consecutive patients with mucosal melanoma of the head and neck were treated with PBT at the National Cancer Center East from March 2004 through February 2007. All patients agreed to participate in the present study. Patient characteristics are listed in Table 1. Median age was 72 years (range, 56 to 79 years). Most patients had a good performance status, and over half the patients had T4 disease.

Toxicity

Major adverse reactions to PBT are listed in Table 2. The most common acute toxicities were mucositis (grade 3, 21%) and mild dermatitis (grade 3, 0%). All patients were able to receive the full dose of PBT (60 GyE) given with a median duration of 36 days (range, 33–42 days). Blindness did not occur, although 2 patients had a unilateral decrease in visual acuity. No treatment-related deaths occurred throughout the study.

Efficacy

Initial local control rate was 85.7% (12/14 patients, 95% confidence interval [CI], 57.2%–98.2%). One patient had recurrent disease, and 1 patient died within 1 year after the initiation of treatment.

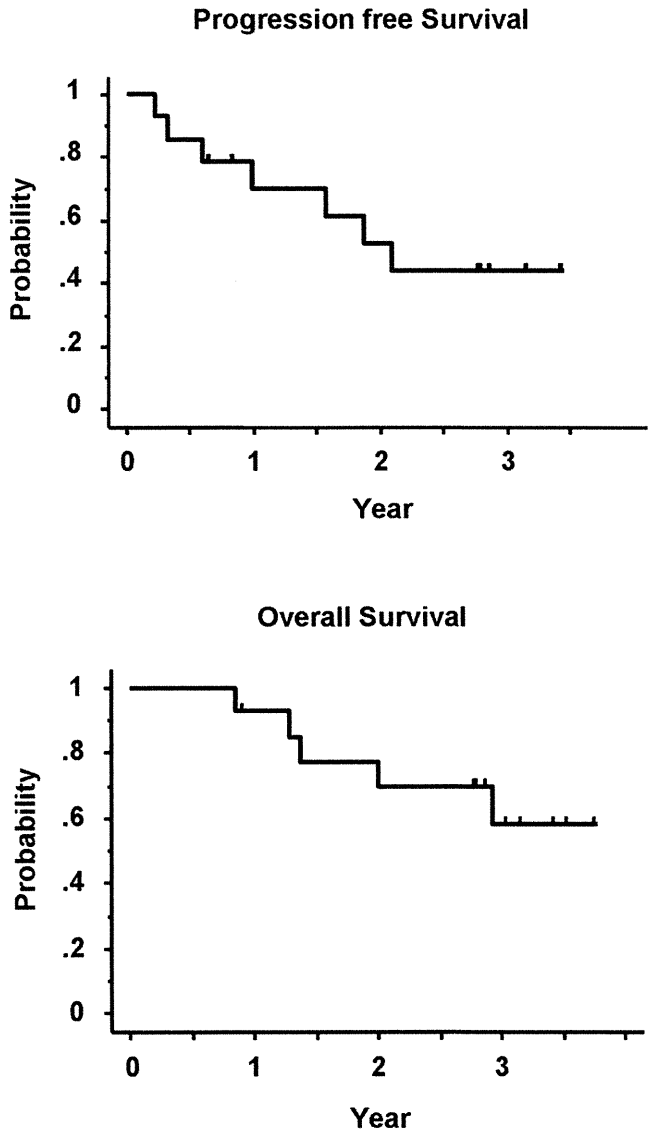

Fig. 2. Progression-free survival (PFS) and overall survival (OS). PFS and OS rates were estimated using the Kaplan-Meier product-limits method. Median PFS was 25.1 months, and 2-year PFS rates were 43.7%. Median survival time was not reached, and 3-year overall survival rate was 58.0% with a follow-up period of 36.7 months.

Table 3. Failure pattern in detail

Case	Time to failure	Failure site	Second-line treatment	Status (time)	Cause of death
1	2.7 M	LN II	Observation	Alive (35.2 M)	
2	22.5 M	LN Ib	Salvage Surgery	Death (35.1 M)	DOD/LM
3	3.8 M	LN Ib, II	Observation	Death (15.4 M)	DOD/DM
6	30.1 M	LN Ib, II	Salvage Surgery	Alive (37.0 M)	
8	11.9M	LN II	Radiation	Death (18.6 M)	DOD/DM
9	7.1 M	LN Ib, II	Salvage Surgery	Death (23.9 M)	DOD/DM
10	8.1 M	Lung	Observation	Death (10.1 M)	DOD/DM
11	18.6 M	Primary site	Observation	Alive (42.7 M)	

Abbreviations: M = months; LN = lymph node; DOD = died of disease; LM = lymph node metastases; DM = distant metastases.

Median PFS was 25.1 months, and 2-year PFS rate was 43.7%. Median survival time with a follow-up period of 36.7 months was not reached, and 3-year overall survival rate was 58.0% (Fig. 2).

Failure pattern and second-line treatment

Six of 14 patients were alive at the end of follow-up with no evidence of disease, while the remaining 8 patients had evidence of disease progression. The most frequent site of first failure was a cervical lymph node outside of the PTV (6/8 patients), followed by local failure in 1 patient (1/8), and lung metastases in one patient (1/8). Failure pattern details are shown in Table 3. With regard to lymph node metastases, 4 patients (4/6) experienced progress within 1 year, and all failure sites were lymph node level Ib or II.

Cause of death

On follow-up, 5 patients died of disease, 4 patients due to cachexia caused by distant metastases and 1 patient by carotid artery perforation cause by lymph nodes metastases.

DISCUSSION

In this study, hypofractionated PBT showed good local control for mucosal melanoma of the head and neck and acceptable toxicity. Prognosis of mucosal melanoma of the head and neck remains poor. In their review of more than 1,000 patients, Mandolis *et al.* (14) reported 5- and 10-year survival rates of 17% and 5%, respectively. Overgaard *et al.* (6) reported a significant relationship between dose per fraction and response, with complete response rates of 59% when fractions of more than 4 Gy were used, compared to 24% with fractions lower than or equal to 4 Gy, while a univariate analysis by Wada *et al.* (9) revealed that a high dose per fraction (3Gy) and high biologically equivalent total dose were associated with better local control and survival.

From these findings, our treatment schedule was planned with consideration for two premises: hypofractionation and high BED. Carbon ion radiotherapy is a promising nonsurgical modality for mucosal melanoma of the head and neck. Yanagi *et al.* (15) reported that with a median follow-up period of 49.2 months, 3-year survival rates were 46.1% in mucosal melanoma patients treated with carbon ion radiotherapy.

The 3-year overall survival rate was 58.0% in the present study. In comparison with the surgical approach or carbon ion therapy, the efficacy of PBT seemed not to be inferior, although recruiting number of patients was small. With regard to late toxicity, decreased visual acuity occurred in 2 patients. Generally, it is often inevitable that the PTV in stage T4 disease with paranasal and/or intracranial invasion includes the unilateral or bilateral optic nerves. In these patients, the better

Table 4. Published cases of late toxicity

Author (study)	Year	Location	Modality	No.of patients	% Treatment outcome	Late toxicity (severe morbidity)
Owens *et al.* (3)	2003	Sinonasal	S	20	5YSR 45%	Not mentioned
			S + RT	24	5YSR 29%	
Temam *et al.* (4)	2005	Sinonasal +α	S/S + RT	30/39	5YSR 20%	Not mentioned
Krengli Owens *et al.* (5)	2006	Head and neck	S/S + RT/others	17/42/15	3YSR 31%	>Grade 3 11% Stenosis of the nasocrimal duct Dry-eye syndrome Optic nerve toxicity Bone necrosis
Wada Owens *et al.* (10)	2004	Sinonasal +α	RT/S+RT	21/10	3YSR 33%	Grade 4 6% soft tissue necrosis; fatal bleeding
Gilligan and Slevin (7)	1991	(Para)-nasal	RT	28	5YSR 17.9%	None
Yanagi *et al.* (15)	2009	Head and neck	Carbon	72	3YSR 46.1%	Grade 2 skin, mucosa*
Present study	2010	Paranasal	Proton	14	3YSR 58.0%	Gade 3 12% unilateral visual acuity

Abbreviations: 5YSR = 5 year survival rate; S = surgery.

* Visual loss after carbon ion radiotheraphy was not mentioned.

dose distribution characteristics of PBT over X-ray should minimize the risk of treatment-related bilateral visual impairment or treatment-related blindness.

Hasegawa *et al.* (16) showed that a certain degree of visual impairment had occurred in 28% of patients whose optic nerves were included in the irradiated volume in carbon ion radiotherapy. There is no report about a direct comparison between PBT and carbon ion radiotherapy.

Previous reports about various approaches to mucosal melanoma are summarized in Table 4.

Cervical lymph nodes were the most frequent site of first failure, and most patients who died finally had distant metastases. Several authors have suggested that aggressive local treatment should be initiated at the presentation of localized melanomas, on the basis that the achievement of local tumor control may increase in survival rate (6, 17). However, it remains controversial whether cervical lymph nodes should be included in the treatment field. We think that what we can do at present is to institute close follow-up after PBT and to detect signs of recurrence or regrowth as early as possible.

CONCLUSIONS

In conclusion, PBT for mucosal melanoma showed promising local control benefit and enough feasibility. To confirm the efficacy and safety, a phase II study of hypofractionated PBT for mucosal melanoma of the head and neck (UMIN-000001505) using the same treatment schedule as the present study is now ongoing in Japan.

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CLINICAL INVESTIGATION

MULTI-INSTITUTIONAL PHASE II STUDY OF PROTON BEAM THERAPY FOR ORGAN-CONFINED PROSTATE CANCER FOCUSING ON THE INCIDENCE OF LATE RECTAL TOXICITIES

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Purpose: Proton beam therapy (PBT) is theoretically an excellent modality for external beam radiotherapy, providing an ideal dose distribution. However, it is not clear whether PBT for prostate cancer can clinically control toxicities. The purpose of the present study was to estimate prospectively the incidence of late rectal toxicities after PBT for organ-confined prostate cancer.

Methods and Materials: The major eligibility criteria included clinical Stage T1-T2N0M0; initial prostate-specific antigen level of ≤20 ng/mL and Gleason score ≤7; no hormonal therapy or hormonal therapy within 12 months before registration; and written informed consent. The primary endpoint was the incidence of late Grade 2 or greater rectal toxicity at 2 years. Three institutions in Japan participated in the present study after institutional review board approval from each. PBT was delivered to a total dose of 74 GyE in 37 fractions. The patients were prospectively followed up to collect the data on toxicities using the National Cancer Institute-Common Toxicity Criteria, version 2.0.

Results: Between 2004 and 2007, 151 patients were enrolled in the present study. Of the 151 patients, 75, 49, 9, 17, and 1 had Stage T1c, T2a, T2b, T2c, and T3a, respectively. The Gleason score was 4, 5, 6, and 7 in 5, 15, 80 and 51 patients, respectively. The initial prostate-specific antigen level was <10 or 10–20 ng/mL in 102 and 49 patients, respectively, and 42 patients had received hormonal therapy and 109 had not. The median follow-up period was 43.4 months. Acute Grade 2 rectal and bladder toxicity temporarily developed in 0.7% and 12%, respectively. Of the 147 patients who had been followed up for >2 years, the incidence of late Grade 2 or greater rectal and bladder toxicity was 2.0% (95% confidence interval, 0–4.3%) and 4.1% (95% confidence interval, 0.9–7.3%) at 2 years, respectively.

Conclusion: The results of the present prospective study have revealed a valuable piece of evidence that PBT for localized prostate cancer can achieve a low incidence of late Grade 2 or greater rectal toxicities. © 2010 Elsevier Inc.

Proton beam therapy, prostate cancer, radiotherapy, clinical trial, rectal toxicity.

INTRODUCTION

The number of patients with organ-confined prostate cancer has been increasing annually because of the widespread screening using prostate-specific antigen (PSA) measurement and aging society. However, organ-confined prostate cancer can now be cured by radical local treatment, including prostatectomy, external beam radiotherapy (EBRT), or brachytherapy, with or without systemic hormonal therapy.

A total dose of >70 Gy using a standard fractionation schedule is considered to be necessary for EBRT to control the disease (1–3); however, the frequency of normal tissue complications increases when the total dose is >70 Gy for conventional EBRT (4, 5). To deliver higher doses to the prostate without increasing the dose to normal tissues, high-technology EBRT, such as three-dimensional conformal radiotherapy (3D-CRT), intensity-modulated radiotherapy (IMRT), and particle therapy have been developed and

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used for more than one decade. Although high-technology techniques of EBRT using X-rays such as 3D-CRT and IMRT have been prospectively evaluated (6–9), the amount of prospective information on the use of proton beam therapy (PBT) alone is limited.

The proton beams used in the modality of particle therapy have distinct physical advantages over conventional photon beams. Proton beams have a low entrance dose, a maximal dose at any prescribed depth called the "Bragg peak," and no exit dose. The "Bragg peak" can be spread out and shaped to conform to the depth and volume of an irregular target. PBT can thus create an inherently three-dimensional conformal dose distribution without exposure of the surrounding normal tissues to excessive doses as compared with the case in conformal photon treatment.

PBT for prostate cancer allows a good dose distribution to be obtained using simple bilateral opposed fields, without exposure of the rectum and bladder to excessive doses. Thus, PBT is theoretically an excellent therapeutic modality owing to its efficacy with reduced toxicity to normal tissues. However, because of the lack of prospective data, it is not yet clear whether PBT can control the incidence of toxicity in the clinical setting. With this background, we started a multi-institutional Phase II trial for patients with organ-confined prostate cancer, focusing on the incidence of late Grade 2 or greater rectal toxicity.

METHODS AND MATERIALS

Patients

The eligibility criteria for inclusion of patients in the present study were (*1*) histologically proven prostate adenocarcinoma, (*2*) clinical Stage II disease (2002 TNM classification, 6th edition, cT1-T2N0M0), (*3*) an initial PSA level of ≤20 ng/mL and Gleason score (GS) of ≤7, (*4*) no history of hormonal therapy or hormonal therapy within 12 months before registration, (*5*) performance status (Eastern Cooperative Oncology Group) 0–2, (*6*) preserved organ function, (*7*) no other active malignancy, and (*8*) written informed consent. Patients with a GS of <7 and PSA level of <10 ng/mL and those with a GS of 7 and/or a PSA level of >10 ng/mL were defined as the low- and intermediate-risk groups, respectively. Patients with Stage cT3-T4, GS 8–10, and/or PSA >20 were not eligible for the present study.

Study endpoints

The primary endpoint was the incidence of late Grade 2 or greater rectal toxicity at 2 years. The secondary endpoints included other toxicities (both acute and late), biochemical relapse-free survival, overall survival, and disease-specific survival. Biochemical failure in the present study was defined as a PSA value of nadir plus 2.0 ng/mL (10, 11), the initiation of any hormonal therapy, or death from any cause.

Study design and statistical analysis

A multi-institutional Phase II study was planned for prospective collection of the toxicity data. The sample size was calculated by the interval estimation method to maintain the accuracy of estimation, using 95% confidence intervals (CIs) of the primary endpoint. The expectation value of the primary endpoint has been defined as <10% according to previous reports of EBRT (12–16). The study sample size was calculated as 150 patients, such that the upper limit of the 95% CI of the primary endpoint was <16% when the actual incidence was <10%. The planned accrual period was 2 years.

Participating institutions

Five institutions were equipped to provide PBT at the beginning of the present study in Japan; three of them participated in the present study, and the institutional review board at each of the three institutions approved the present study (National Cancer Center Hospital East, Kashiwa; Shizuoka Cancer Center, Shizuoka; and Hyogo Ion Beam Medical Center, Hyogo).

Treatment planning

The patients were placed in the supine position and fixed with a vacuum cushion or a thermoplastic cast. The patients were instructed to maintain regular bowel movement; patients with constipation were prescribed laxatives such as magnesium oxide, sennoside, and/or picosulfate sodium to control bowel movement. The bladder filling was controlled by water drinking after urination, and all PBT sessions were performed with a full bladder.

The clinical target volume was defined as the prostate alone for low-risk patients and as the prostate plus the proximal seminal vesicles for intermediate-risk patients, at least encompassing all-known diseases identified by the planning computed tomography scan and other clinical information. The planning target volume consisted of the clinical target volume with optimal margins to account for the uncertainties from the patient setup or internal organ motion, which were estimated at each institution (Table 1). The rectum, from the sigmoid flexure to the anal verge, and the entire bladder as solid organs were delineated as the critical normal structures.

Proton beam therapy

The PBT was delivered at a total dose of 74 GyE in 37 fractions (2 GyE/d). In the low-risk patients, the prostate alone received 74 GyE; in the intermediate-risk patients, a booster dose of 24 GyE was added to the prostate alone after the initial 50 GyE was delivered to the prostate and proximal seminal vesicles. As listed in Table 1, the dose prescription was determined by each institutional method. The dose constraints for the normal tissues were as follows, on the basis of the results from our previous analysis (17) (V_X indicates the percentage of volume receiving more than x GyE): rectum, V_{50} <35%, V_{60} <25%, and V_{70} <15% in the low-risk patients; V_{50} <40%, V_{60} <30%, and V_{70} <20% in the intermediate-risk patients; bladder, V_{65} <50%, V_{70} <35%; femoral head, maximal dose <50 GyE.

Table 1. Details of treatment planning in each institution

Institution	PTV margin (mm)	Dose prescription	Bolus/collimator
NCCHE	7	90% dose to PTV	Individual bolus/ collimator
SCC	5	95% dose to PTV	Individual bolus/ collimator
HIBMC	8–10	To isocenter	No bolus/multileaf collimator

Abbreviations: NCCHE = national cancer center hospital east; SCC = shizuoka cancer center; HIBMC = hyogo ion beam medical center; PTV = planning target volume.

Bilateral opposed fields were used for proton dose delivery. The range modulation by bar-ridge filters was used to generate a spread-out Bragg peak. Proton beams with optimal energy in the range of 190–235 MeV were selected, and individual boluses and collimators were manipulated to conform to the target volume. Daily verification of patient positioning was performed in all patients using orthogonal radiography according to the bony structures. The relative biologic effectiveness of the proton beam was estimated to be 1.1 compared with that of the photon X-rays (GyE = proton Gy × 1.1), in animal experiment conducted at each institution.

Assessments

The registered patients were prospectively followed up to collect data on the toxicities and PSA values at 1 month and once every 3 months after PBT completion for the first 2 years and once every 6 months thereafter.

Late toxicities were defined as those observed >90 days after the start of PBT, but the Radiation Therapy Oncology Group (RTOG)/European Organization for Research and Treatment of Cancer late radiation morbidity scoring scheme was not used in the present study to assess the late toxicities. Instead, so that the observed symptoms could be individually assessed and scored objectively, the National Cancer Institute Common Toxicity Criteria, version 2.0, was used to assess the acute and late toxicities. The symptoms of rectal and/or bladder toxicities assessed in the present study included proctitis, rectal bleeding, rectal pain, hematuria, urinary frequency/urgency, urinary retention, and dysuria (painful urination). Toxicity grading was determined using the severity of each symptom assessed objectively. The Common Terminology Criteria for Adverse Events, version 3.0, was not available when preparing the present study and was not used.

The cumulative incidence of the toxicities and the survival rates were analyzed using the Kaplan-Meier method.

RESULTS

Patients

Between March 2004 and March 2007, 151 patients were enrolled at the 3 institutions for the present study. The patient characteristics are listed in Table 2. Of the 151 patients, 77 and 74 were low- and intermediate-risk, respectively. All the patients enrolled in the present study received the planned PBT up to a dose of 74 GyE in 37 fractions. The median follow-up period was 43.4 months (range, 3–62). Of the 151 patients, 3 were lost to follow-up within the first 2 years, and 1 patient died of other causes on Day 165, without biochemical failure. These 4 patients were excluded from the analysis of late toxicity. All the patients enrolled in the present study were included in the assessment of the acute toxicities and efficacy.

Acute toxicities

The acute rectal and bladder toxicities observed within 90 days of the initiation of PBT are listed in Table 3. All the acute toxicities observed were transient and resolved spontaneously after completion of PBT. No Grade 3 or greater acute toxicities were observed. The rectal toxicities observed included anal pain at defecation, soft stool, anal discomfort, and rectal bleeding. The bladder toxicities were urinary frequency, dysuria, narrow stream, and urinary retention.

Table 2. Patient and tumor characteristics

Characteristic	Value
All patients (*n*)	151
Age (y)	
Median	67
Range	51–82
cT Stage (*n*)	
T1c	75
T2a	49
T2b	9
T2c	17
T3a	1
Gleason score (*n*)	
4	5
5	15
6	80
7	51
iPSA (ng/mL)	
10	102
10–20	49
Hormonal therapy (*n*)	
Yes	42
No	109
Risk group (*n*)	
Low risk	77
Intermediate risk	74

Abbreviation: iPSA = initial prostate-specific antigen.

Late toxicities

The late toxicities at the final follow-up of the 147 patients who had been followed up for >2 years are listed in Table 4. No Grade 3 or greater late rectal toxicities were observed. The late rectal toxicities observed included rectal bleeding, urgency of defecation, and anal pain. The bladder toxicities were transient gross hematuria and urinary retention.

The Kaplan-Meier curves of late rectal and bladder toxicities are shown in Fig. 1. The incidence of late Grade 2 or greater rectal toxicity was 2.0% (95% CI, 0–4.3%) at 2 years (primary endpoint of the present study) and 4.1% (95% CI, 0.4–7.7%) at the final follow-up. The corresponding data for bladder toxicity were 4.1% (95% CI, 0.9–7.3%) at 2 years and 7.8% (95% CI, 2.9–12.8%) at the final follow-up.

Efficacy

The median follow-up period was 43.4 months in the present study. We evaluated the biochemical relapse-free survival

Table. 3. Acute toxicities

Toxicity	Patients (*n*)
Total	151 (100)
Rectum	
Grade 0	135 (89)
Grade 1	15 (10)
Grade 2	1 (0.7)
Bladder	
Grade 0	46 (30)
Grade 1	87 (58)
Grade 2	18 (12)

Data in parentheses are percentages.

Table 4. Late toxicities

Toxicity	Patients (*n*)
Total	147* (100)
Rectum	
Grade 0	115 (78)
Grade 1	27 (18)
Grade 2	5 (3)
Bladder	
Grade 0	128 (87)
Grade 1	9 (6)
Grade 2	8 (5)
Grade 3	2 (1)

Data in parentheses are percentages.
* Number of patients followed up for >2 years.

using the failure definition of nadir plus 2.0 ng/mL; the Kaplan-Meier curve is shown in Fig. 2. Two patients died of other causes on Day 165 and Day 1,202, respectively, without biochemical failure. No patients died of prostate cancer. The biochemical relapse-free survival rate was 94% at 3 years (95% CI, 90–98%).

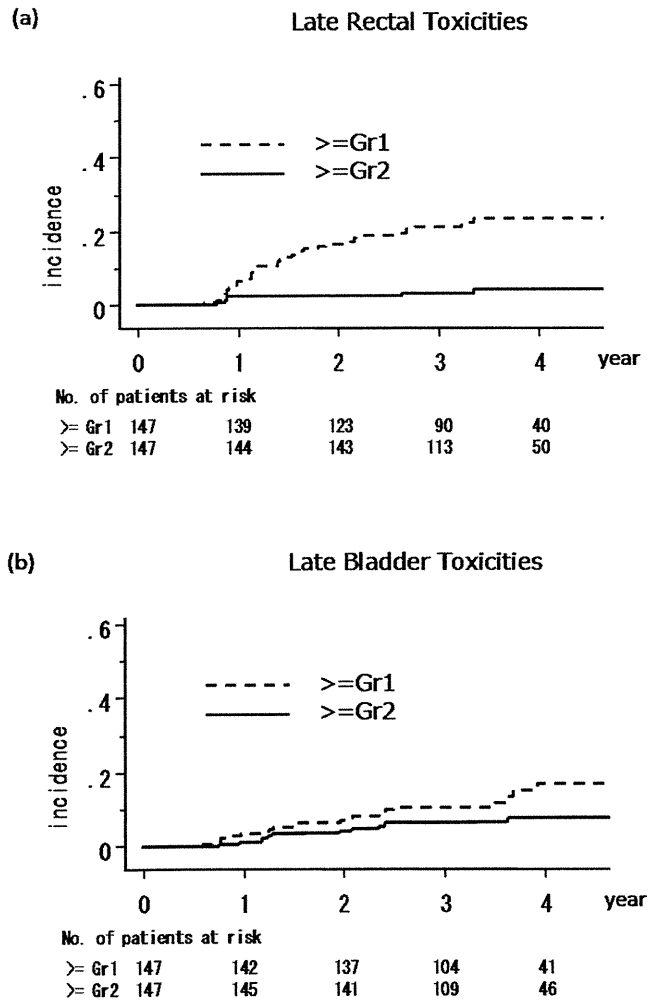

Fig. 1. Kaplan-Meier curves of late (a) rectal and (b) bladder toxicity. Gr = grade.

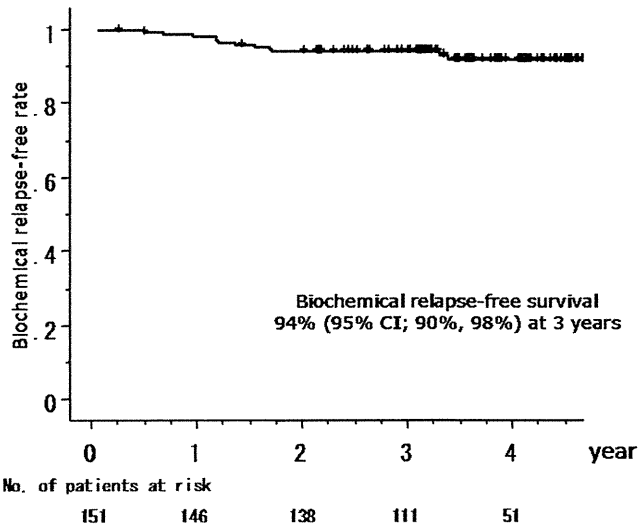

Fig. 2. Biochemical relapse-free survival using definition of nadir plus 2.0 ng/mL. CI = confidence interval.

DISCUSSION

The results of the present prospective study have provided valuable evidence to show a low incidence of late Grade 2 or greater rectal toxicities after PBT for organ-confined prostate cancer. Although PBT is theoretically an excellent modality for EBRT, providing an ideal dose distribution, few well-designed prospective clinical trials are available to corroborate its superiority. The present prospective study was conducted with the aim of scientifically clarifying whether PBT can control the incidence of late rectal toxicity.

The RTOG conducted a Phase I-II dose-escalation study to determine the maximal tolerated dose of 3D-CRT for prostate cancer (RTOG 94-06). The toxicity results of RTOG 94-06 showed a significantly lower incidence of Grade 3 or greater late toxicity but a significantly greater incidence of Grade 2 or less late toxicity than expected from the results of previous RTOG trials (13–15, 18). Although Grade 2 toxicity is generally defined as moderate in severity and is often underestimated, the patients' quality of life can suffer even from such moderate toxicity. Therefore, it is becoming increasingly important to devise sophisticated techniques for high-dose EBRT, such that even the frequency of moderate Grade 2 or less toxicity can be reduced in patients with prostate cancer.

Of the late toxicities that characteristically occur in patients receiving high-dose EBRT for prostate cancer are rectal toxicities, which in most cases, are represented by rectal bleeding, occurring within 2 years of treatment completion. For this reason, the primary endpoint of the present study was defined as the incidence of late Grade 2 or greater rectal toxicity at 2 years after treatment completion.

Since the 1990s, when the existence of a dose–response relationship was suggested in prostate cancer patients undergoing EBRT, dose escalation has been eagerly pursued using high-technology EBRT. In a randomized Phase III trial of 3D-CRT at the M.D. Anderson Cancer Center, the toxicity results revealed an incidence of late Grade 2 or greater

gastrointestinal (GI) toxicity in a high-dose arm (78 Gy in 39 fractions) and standard-dose arm (70 Gy in 35 fractions) of 26% and 12%, respectively (16). Michalski *et al.* conducted a multi-institutional dose-escalation Phase I-II study of 3D-CRT (RTOG 9406; Level 1, 2, and 3, 68.4, 73.8, and 79.2 Gy at 1.8 Gy/fraction; Level 4 and 5, 74 and 78 Gy in 2 Gy/fraction, respectively) and reported the late toxicity profiles at each dose level. Late Grade 2 or greater GI toxicity occurred in 9–13%, 7–9%, 11–14%, 10–16%, and 25–26%, respectively, at dose levels 1–5 (6).

Intensity-modulated RT (IMRT) is a modality of high-technology EBRT. The Memorial Sloan-Kettering Cancer Center has been conducting a single-institutional dose-escalation trial of 3D-CRT and IMRT. Zelefsky *et al.* (7–9) reported that Grade 2 or greater late GI toxicity occurred at an incidence of 16% in patients who had undergone 3D-CRT to a total dose of 75.6–81 Gy (1.8 Gy/fraction). The corresponding incidence was only 2%, even in patients who had undergone IMRT to a total dose of 81–86.4 Gy (7–9). According to a retrospective analysis from the Fox Chase Cancer Center, Grade 2 or greater late GI toxicity occurred at an incidence of 2.4% in patients who had undergone IMRT to a total dose of 74–78 Gy (2 Gy/fraction) (19). However, in some reports, no reduction in the incidence of late toxicity could be achieved despite using IMRT. De Meerleer *et al.* (20) reported that 18% of patients experienced Grade 2 or greater late GI toxicity after receiving a total dose of 74–76 Gy (2 Gy/fractions). Vora *et al.* (21) also reported an incidence of late Grade 2 or greater toxicity of 24% in patients who had undergone IMRT to a total dose of 75.6 Gy.

The PBT facility was installed at Loma Linda University Medical Center in 1990, and the morbidity results for the prostate cancer patients treated to a total dose of 74–75 GyE (1.8–2.0 GyE/fraction) were reported. Late Grade 2 GI toxicity had developed in 21% of the patients at 3 years after treatment completion (22).

The toxicity results in previous reports are summarized in Table 5. The incidence of late Grade 2 or greater rectal toxicity in patients who had undergone 3D-CRT was 9–16% (6, 8). The Memorial Sloan-Kettering Cancer Center and Fox Chase Cancer Center reported a very low incidence of late Grade 2 or greater rectal toxicity (2–2.4%) in patients who had undergone IMRT (7, 9, 19). In contrast, some other centers have reported a high incidence of late rectal toxicity (>15%) even in patients who had undergone IMRT (20, 21). Our results have shown that the incidence of late Grade 2 or greater rectal toxicity was 2.0% at 2 years and 4.1% at the final follow-up; the upper limit of the 95% CIs of these values was 4.3% and 7.7%, respectively. Our results cannot be directly compared with the toxicity data from previous reports, because these studies used different grading scales and also included retrospective and/or single institution-specific data. However, the incidence of late rectal toxicity associated with PBT in the present study was lower than those from the 3D-CRT series and, at least, was not greater than the historical data on the incidence of late Grade 2 or greater GI toxicity after 3D-CRT and IMRT (12–16). The result for the primary endpoint in the present study was 2.0% (95% CI, 0–4.3%) at 2 years, providing at least one piece of scientific evidence of PBT in patients with prostate cancer.

Table 5. Overview of late gastrointestinal toxicity in EBRT for localized prostate cancer

Institution/study	Patients (*n*)	Dose (Gy)	Technique	Grading scale	Grade 1	Grade 2	Grade 3	Follow-up (y)
MDACC (16)	150	70	3D-CRT	RTOG/LENT	36%	11%	1%	6
	151	78			28%	19%	7%	6
					23%, Grade 2/3			2
RTOG 9406 (6)	112	68.4	3D-CRT	RTOG	9-13%, Grade 2/3			9–12
	300	73.8			7-9%, Grade 2/3			7–10
	167	79.2			11-14%, Grade 2/3			9
	256	74			10-16%, Grade 2/3			7–8
	220	78			25-26%, Grade 2/3			6
MSKCC (7–9)	695	75.6–81	3D-CRT	Modified RTOG/CTCAE,	16%, Grade 2/3			5
	561	81–86.4	IMRT	version 3.0	2%, Grade 2/3			7
FCCC (19)	216	74–78	IMRT	Modified RTOG/LENT	2.4%, Grade 2/3			3.5
Ghent University (20)	133	74–76	IMRT	Modified RTOG	47%	17%	1%	3
Mayo Clinic Arizona (21)	145	75.6	IMRT	Modified RTOG	20%	23%	1%	4
LLUMC (22)	643	74	PBT alone	RTOG	—	21%	—	3
		75	X+PBT					
Present study	151	74	PBT alone	NCI-CTC, version 2.0	14%	2.0%, Grade 2/3		2

Abbreviations: EBRT = external beam radiotherapy; MDACC = M.D. Anderson Cancer Center; 3D-CRT, three-dimensional conformal radiotherapy; RTOG = radiation therapy oncology group; LENT = late effects normal tissue task force; MSKCC = memorial sloan-kettering cancer center; IMRT = intensity-modulated radiotherapy; CTCAE = common terminology criteria for adverse events; FCCC = fox chase cancer center; LLUMC = loma linda university medical center; X = photon radiotherapy; PBT = proton beam therapy; NCI-CTC = national cancer institute common toxicity criteria.

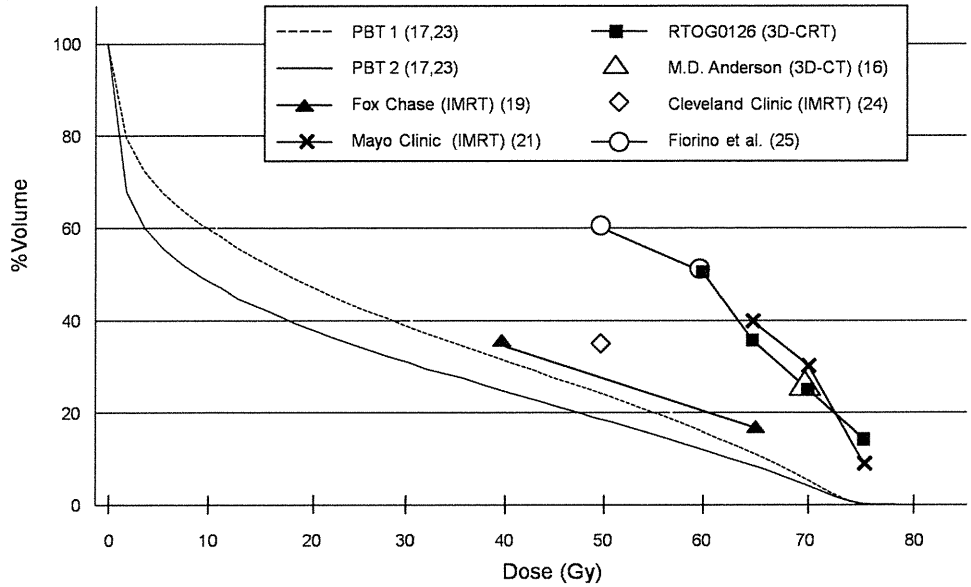

Fig. 3. Average dose–volume histograms (DVHs) for rectum in photon beam therapy (PBT) compared with other dose constraints used in three-dimensional conformal radiotherapy (3D-CRT) or intensity-modulated radiotherapy (IMRT). PBT1 = PBT for intermediate-risk patients; PBT2 = PBT for low-risk patients; RTOG = Radiation Therapy Oncology Group. Data in parentheses indicate reference report.

The previously reported average dose–volume histograms for the rectum in PBT performed at National Cancer Center Hospital East are shown in Fig. 3 (17, 23) and were compared with the other dose constraints used in 3D-CRT or IMRT (16, 19, 21, 24, 25). PBT with simple bilateral opposed fields can thus reduce the dose to the rectum through high to intermediate dose levels, thereby achieving the ideal dose–volume histogram for the rectum. Although bilateral opposed fields can present another issue regarding the dose to the femoral head, the maximal dose to the femoral head in PBT with simple bilateral opposed fields is <35 GyE (45% of the prescribed dose), and V_{30} of the femoral head is <40%.

The total dose of PBT used in the present study was 74 GyE, administered using a conventional fractionation schema (2 GyE/fractions); however, the possible benefit of additional dose escalation and hypofractionation of EBRT for prostate cancer has been suggested, and the efficacy of such a strategy is now under investigation in some randomized trials (16, 26–28). As shown in Fig. 3, the excellent dose–volume histograms for the rectum in PBT might allow the implementation of these strategies; however, additional prospective data are required to ascertain whether PBT administered using these investigational approaches can yield a low frequency of late rectal toxicity.

A more objective grading scale might be necessary to allow comparison of the morbidity data, because different grading scales have been used in previous reports. Thus, the health-related quality of life might be a more rigid and comparable indicator for assessing the toxicities.

Late genitourinary toxicity often occurs after a longer follow-up period (29, 30) and is another issue that needs to be addressed with the use of high-dose EBRT for prostate cancer. As shown in Fig. 1, the incidence curve of late bladder toxicity seems to have been increasing over the years, and that of late rectal toxicity reached a plateau after a few years. Longer follow-up is needed for a more precise assessment of both late genitourinary and GI toxicity.

Quality assurance procedures for clinical assessments have an important role in enhancing confidence in the results of multi-institutional clinical trials. At the beginning of the present study, the specifications for PBT facilities differed among the participating institutions, and the method of delivery of the proton beams to the target organs was defined at the discretion of each institution (Table 1). Although such institutional differences in the method of dose delivery can affect the incidence of toxicity, no significant difference was found in the incidence of toxicity among the three institutions in the present study. Owing to the increasing number of PBT facilities, implementation of quality assurance procedures for PBT in multi-institutional trials, from both the clinical and the physics aspect, is gaining importance.

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Apparent absence of a proton beam dose rate effect and possible differences in RBE between Bragg peak and plateau

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Purpose: Respiration-gated irradiation for a moving target requires a longer time to deliver single fraction in proton radiotherapy (PRT). Ultrahigh dose rate (UDR) proton beam, which is 10–100 times higher than that is used in current clinical practice, has been investigated to deliver daily dose in single breath hold duration. The purpose of this study is to investigate the survival curve and relative biological effectiveness (RBE) of such an ultrahigh dose rate proton beam and their linear energy transfer (LET) dependence.

Methods: HSG cells were irradiated by a spatially and temporally uniform proton beam at two different dose rates: 8 Gy/min (CDR, clinical dose rate) and 325 Gy/min (UDR, ultrahigh dose rate) at the Bragg peak and 1.75 (CDR) and 114 Gy/min (UDR) at the plateau. To study LET dependence, the cells were positioned at the Bragg peak, where the absorbed dose-averaged LET was 3.19 keV/μm, and at the plateau, where it was 0.56 keV/μm. After the cell exposure and colony assay, the measured data were fitted by the linear quadratic (LQ) model and the survival curves and RBE at 10% survival were compared.

Results: No significant difference was observed in the survival curves between the two proton dose rates. The ratio of the RBE for CDR/UDR was 0.98 ± 0.04 at the Bragg peak and 0.96 ± 0.06 at the plateau. On the other hand, Bragg peak/plateau RBE ratio was 1.15 ± 0.05 for UDR and 1.18 ± 0.07 for CDR.

Conclusions: Present RBE can be consistently used in treatment planning of PRT using ultrahigh dose rate radiation. Because a significant increase in RBE toward the Bragg peak was observed for both UDR and CDR, further evaluation of RBE enhancement toward the Bragg peak and beyond is required. © *2010 American Association of Physicists in Medicine.* [DOI: 10.1118/1.3490086]

Key words: proton therapy, relative biological effectiveness, dose rate

I. INTRODUCTION

Respiration-gated irradiation system in proton radiotherapy for a moving target, such as lung or liver cancer, requires a long time to deliver fractional dose.[1,2] Also, if the dose escalation is to be realized in the near future, as in the case for carbon therapy,[3,4] the treatment time would have to be prolonged. The prolongation of fractional and overall treatment time possibly increase the risk of treatment error and psychological distress of the patient. Therefore, a short-time high dose rate irradiation protocol is strongly desired.

At the National Cancer Center Hospital East in Japan, an irradiation system for high dose rate therapy (the "one-shot radiation system") has been developed. This system consists of proton image guided radiation therapy system, which enables us to monitor the tumor position during irradiation, and an AVF cyclotron, which has a maximum beam current of 300 nA. A spot scanning system[5] is also under construction, which would allow us to achieve a better DVH for healthy tissues and reduce side effects. For the latter system, high dose rate radiation is achieved by combining the strong intensity beam from the cyclotron and the high-speed scanning technique. If we compare it to the clinical dose rate used to treat liver tumors, which is around 5–8 Gy/min in our institute, the dose rate that we are planning to use in one-shot radiation therapy is more than tenfold higher. The achievable dose rate is even greater in the spot scanning system.