Early stage non-small cell lung cancer treated with pencil beam scanning particle therapy: retrospective analysis of early results on safety and efficacy

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Abstract

Background: To evaluate the safety and efficacy of particle therapy (PT) using pencil beam scanning (PBS) technique for early stage non-small cell lung cancer (NSCLC).

Methods: From 08/2014 to 03/2018, 31 consecutive patients with sum of the longest diameters of primary tumor and hilar lymph node < 5 cm, N0–1, M0 NSCLC treated with PT were retrospectively analyzed. Gating/active breathing control techniques were used to control tumor motion in 20 and 7 patients. PBS-based proton radiotherapy (PRT) or carbon ion radiotherapy (CIRT) plans were designed via Syngo® planning system. PRT, PRT + CIRT boost, and CIRT were used in 6, 6 and 19 patients, respectively. Prescriptions were categorized to 3 levels: 5–7.5 GyE * 8–10Fx, 4–5 GyE * 15–16 Fx and 2.25–3.5 GyE * 20–31 Fx.

Results: Thirty-one patients (20 males and 11 females) with a median age of 71 (50–80) years were enrolled with a median follow-up time of 12.1 (2.9–45.2) months. Fourteen were adenocarcinomas, 7 squamous cell carcinomas, 4 non-specified NSCLC and 6 had no histological diagnosis (4/6 had previous resected lung cancer). The median tumor size was 3.1 (1.1–4.7) cm. No grade 4–5 toxicities were observed. One patient experienced grade 3 (per the Common Terminology Criteria for Adverse Events version 4.03) radiation-induced lung injury (RILI) at 6.7 months from radiation started. Grade 2 acute toxicities included hematological toxicities (5 cases), RILI (2), plural pain (1) and dermatitis (1). Grade 2 late toxicities included RILI (3) and asymptomatic rib fracture (1). Three patients had progressed disease at 4.0–10.6 months after the initiation of PT. One experienced local failure with simultaneous distant failure and died of brain metastasis at 10.8 months; one developed regional and distant failure and died of lung infection at 8.7 months; the other experienced isolated distant failure only and his disease was well controlled after salvage systemic therapy. The estimated rates of progression-free survival, local control, cause-specific survival and overall survival at 1, 2 years were 85.5% and 85.5%, 95.2% and 95.2%, 95.0% and 95.0%, 90.7% and 90.7%, respectively.

Conclusions: PBS-based PT appears safe and effective for early stage NSCLC. Further follow-up and investigation is warranted.

Trial registration: ISRCTN, ISRCTN78973763. Registered 14 August 2018- Retrospectively registered, http://www.isrctn.com/ISRCTN78973763.

Keywords: Particle therapy, Proton, Carbon ion, Radiotherapy, Pencil beam scanning, Early stage non-small cell lung cancer

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Background
Lung cancer is the leading cause of cancer death worldwide. Approximately 80% of lung cancer patients are diagnosed with non-small cell lung cancer (NSCLC). Surgery is currently the standard therapy for early stage NSCLC. However, patients with severe underlying co-morbidities such as chronic obstructive pulmonary disease and cardiac insufficiency are not eligible for surgical procedures. Radiation therapy (RT), including stereotactic body radiotherapy (SBRT), has become the most favorable options for these patients [1].

With the physical dosimetrical advantage of particle beams and additional biological advantage of carbon ion beams compared with conventional radiation, particle therapy (PT) might be able to achieve better sparing of the normal tissue and even better tumor control and has therefore been used for the treatment of NSCLC patients with impaired pulmonary or heart function [2].

There are two major techniques implemented for PT: passive scattering (PS) and pencil-beam scanning (PBS). Theoretically, PBS can provide better dose distribution than PS in early stage NSCLC [3, 4]. However, PBS was relatively slowly applied to lung cancer worldwide because of the possible inaccuracy of dose distribution caused by the interplay effect, complicated tissue density variation of thorax area under PBS, respiratory motions, and range uncertainties of particle beams. Only several patients of early stage NSCLC were mentioned being treated by PBS-proton radiotherapy (PRT) in a previous study [5], no experience has been reported for carbon-ion radiotherapy (CIRT) using PBS technology for early stage NSCLC. After investigating how to deal with the interplay effect and delivery uncertainties for lung cancer, we successfully used PBS technique of PT to treat early stage NSCLC. In this retrospective study, we present our initial experience and clinical results at Shanghai Proton and Heavy Ion Center (SPHIC).

Methods
Patient and pretreatment evaluations
Patients with T1-2a, N0–1, M0 (American Joint Committee on Cancer staging manual version 7th) NSCLC and deemed medically inoperable or declined surgery were included in this study. The sum of the longest diameters of primary tumor and hilar lymph node was required to be less than 5 cm. All patients had pathologically confirmed NSCLC, or clinically diagnosed as NSCLC for the Fluorodeoxyglucose-Positron Emission Tomography/Computed Tomography (FDG-PET/CT) avid lung lesion and confirmed by two senior radiologists independently using computed tomography (CT) images. This study was conducted in accordance with the Declaration of Helsinki, and the protocol was approved by the Institutional Review Board (IRB) of SPHIC (27/10/2017, 171020EXP-01). Written informed consents from all patients for using their data were obtained for this study.

Pretreatment evaluation for all patients included a complete disease history and physical examination, complete blood count, serum electrolytes, renal and liver function tests, electrocardiogram, pulmonary function tests and a mandatory FDG-PET/CT scan for clinical staging.

Particle therapy (PT)

Patient immobilization
The patients were positioned in supine or prone position based on the location of the radiation targets using a vacuum bag. A thermoplastic mask was used to fix the position and restrict the breath motion for patients using free breathing or gating.

Tumor motion management, simulation CT and contouring
All patients were evaluated for tumor motion under fluoroscopy before simulation. If the tumor motion in all the directions was less than 5 mm, the patient would be treated under free breathing (FB); if the motion exceeded 5 mm, a breath control technique, either active breathing control (ABC) or respiratory gating, was required to mitigate the residual motion to < 5 mm during treatment.

The simulation CT scanning started from the angle of mandible to adrenal glands to include tumor lesions, entire lungs, whole neck and all the organs/tissues by which the beams probably pass. CT scanning was performed under the same respiratory mode as that would be used for PT treatment. All CTs for planning were plain CT, and a contrast CT afterwards was scanned for image fusion only when needed.

For patients to be treated under FB or ABC, a plain simulation CT was scanned under the same respiratory mode. The gross tumor volume (GTV) was delineated on the plain CT.

For patients using gating, a 4-dimensional (4D) simulation CT was performed. Ten phases of a whole respiratory cycle were reconstructed. The gating window (respiratory phase time when beam is on, usually around the end of exhale) was selected to restrict the tumor motion to < 5 mm. The GTV was delineated on one phase of them (usually the end of exhale), then propagated to all other phases using deformable registration and confirmed by physicians. The GTVs generated on each phase were combined to form the internal gross tumor volume (iGTV).

The clinical target volume (CTV) was defined as a 0.6–0.8 cm expansion from GTV/iGTV. Range uncertainties and set-up errors were counted when creating the planning target volume (PTV). In most instances, it was CTV plus a 0.5–0.7 cm margin in lateral and a 0.7–1.5 cm margin along beam direction.
Prescriptions
The dose of PT is defined as equivalent dose to Gy of photon (GyE). Three radiation beam sources were used for this group of patients: PRT only, PRT + CIRT boost, or CIRT only. The selection of radiation sources or their combination depended on the clinical trials patients involved. For patients who did not participate in clinical trials, patients were treated according to our IRB approved institutional treatment protocols. From March 2016 ~ July 2017, 3 prospective phase I/II clinical trials approved by hospital IRB were started to enroll patients with stage I NSCLC. According to the distance from the primary lesion to critical organs at risk (OARs), patients were categorized to have peripheral, median, and central lesions. Overall, dose and fractions used included 5–7.5 GyE * 8–10 Fx, 4–5 GyE * 15–16 Fx and 2.25–3.5 GyE * 20–31 Fx.

At least 99% of GTV covered by 99% of the prescription dose, 99% of CTV covered by 95% of the prescription dose, and 90% of PTV covered by 90% of the prescription dose were required. All treatment plans aimed to meet stringent guidelines for protection of OARs, including lungs, spinal cord, esophagus, trachea, proximal bronchi, and heart/big vessels. The normal tissue dose-volume constraints for conventionally fractionated and ablative radiotherapy of the National Comprehensive Cancer Network guideline as well as published results from M.D. Anderson Cancer Center (MDACC) and the University of Florida Proton Therapy Institution were considered to set up the dose constraints for our patients according to fraction size [6–10].

Particle therapy planning
The Siemens Syngo® treatment planning system was used for all patients’ PT planning. PRT (beam energy 30–250 MeV) and CIRT (beam energy 85–430 MeV) plans were designed by using 2 to 4 beams with PBS technique. Single beam optimization (SBO) or intensity modulated particle therapy (IMPT) method was decided to be used in treatment by evaluating delivery uncertainties and doses to critical organs during planning. Without the feature of robustness evaluation in the Syngo® system, several steps were applied to reduce the uncertainties. First, similar to the experience of MDACC [5], the threshold of 5 mm was selected for motion control in our center. Based on the physical experiment on Alderson phantom conducted in our center, the deviation of dose distribution was found to be acceptable (< 5%) when respiratory movement was 4.4 mm or less. Second, iGTV of the plan was contoured to include all tumor trajectory in 10 phases, and the density of the intersection of iGTV and GTV will be overridden to a similar density of the primary lesion before planning. Third, when planning, physicists will avoid the beam angles passing through the heart and great vessels and keep the beam path less inhomogeneous. Forth, for hypo-fractionation plans (fraction dose ≥6 GyE), strategies of repeated SBO which is mimicking the volumetric rescanning [11], and specific beam PTV expansion were typically used to increase the plan robustness. IMPT was only used for lesions with minor motions, or large volumes and irregular shapes. Fifth, large sport size (full width at half maximum, FWHM = 10 mm) and decreased iso-energy slice spacing (Grid resolution = 2 mm) were used to increase target dose heterogeneities [12]. Sixth, 4D plan recalculation and accumulated dose also have been used for hypo-fractionation plans to evaluate dose distribution. Last but not least, before the first treatment and every week during treatment, simulation CT and plan recalculation on the latest CT will be conducted for every patient, to check the dose distribution, and replanning was demanded when recalculation revealed poor coverage of targets or overdosing to the OARs. Two orthogonal X-ray images were taken to verify and registrate patient position according to bone structures before daily irradiation. The set-up error of ≤3 mm before the treatment was allowed. Typical cases of CIRT are shown in Fig. 1.

Follow up and evaluation
All patients were evaluated weekly for treatment-induced toxicities and disease response/progression during their treatment. After the completion of PT, all patients are required to be evaluated according to our institutional follow-up protocol for lung cancer at 3 months after the 1st day of PT, every 3–4 months within the first 2 years, every 6 months between year 3 and 5, and annually thereafter. Treatment-induced side effects were scored using the Common Terminology Criteria for Adverse Events (CTCAE) version 4.03, for events observed after the first dose of irradiation. Toxicities occurred 90 or more days after the completion of PT were defined as late toxicities. The Response Evaluation Criteria in Solid Tumors (RECIST) version 1.1 was used for tumor response evaluation.

Statistical analyses
The overall survival (OS) time was calculated from the date of pathological diagnosis of the primary disease or radiological diagnosis of disease for patients without pathological confirmation until death or the date of the last follow-up. The time to local, regional, or distant failure was calculated from the date of the first fraction of PT until documented first considered treatment failure. Local control (LC) and various survival rates were calculated using the Kaplan-Meier method. All analyses were performed in SPSS statistics version 21 (Armonk, NY; USA).
Results

Patient characteristics
Between August 2014 and March 2018, 31 consecutive and non-selected patients with NSCLC met the inclusion criteria were included in this study. Histological diagnoses were obtained in 25 patients. Biopsy were either declined or contraindicated for the remained 6 patients, including 4 diagnosed as the second primary lung cancer with previous resected primary NSCLC in other locations. All patients had stage I-IIa disease. Eight patients refused surgery, and 23 patients were medically inoperable (including chronic obstructive pulmonary, interstitial lung disease, cardiovascular disease, etc.). Patient characteristics are detailed in Table 1.

Particle therapy and chemotherapy
Particle therapy
The breath control techniques, beam types and doses of PT used are detailed in Table 2. Most patients (27/31) were treated with breath control techniques. Prescriptions of biological effective dose (BED, $\alpha/\beta = 10$) < 100 GyE (using relatively low fraction dose) mostly prescribed to patients with N1 disease. The mean values of doses to thoracic OARs of all patients were listed in Table 3.

Chemotherapy
Chemotherapy was recommended to 16 patients with larger tumor (> 4 cm in diameter) and/or N1 disease; however, only 6 received some form of chemotherapy due to co-morbidities or patients’ refusal. Chemotherapy regimens and schedules were detailed in Table 2.

Survival and disease control
Till the last follow up on July 16th, 2018, the median follow-up time was 12.1 (range 2.9–45.2) months for the entire cohort of the patients. Three patients had progressed disease and 2 out of them died during follow-up. One experienced simultaneous local and distant failure at 7.3 months, then died of brain metastasis at 10.8 months; one suffered simultaneous regional (lymph node out of the radiation field) and distant failure at 4 months, and died of fungal pneumonia at 8.7 months after the completion of CIRT. The third patient developed distant metastasis only at 10.6 months, and the disease was well controlled after salvage chemotherapy and immunotherapy. In addition, one patient diagnosed as T1N1M0 disease had a single bone lesion considered as benign before PT. The bone lesion was proved as metastasis after the completion of PT. He received palliative photon RT to
it and systemic chemotherapy. Both the thoracic and bone lesions remained stable at his last follow-up.

Two patients developed a second primary lung cancer after 25.3 and 31.5 months of particle therapy. The new lesion of one patient occurred from a chronic lung lesion which already existed near to the original primary lesion before PT and was not included in the proton irradiation area; that of the other was close to PT irradiation area but with different pathological type. In addition, two patients were diagnosed with rectal and gastric cancer, respectively, at 1 year and 3 months after PT and received surgery or palliative treatment.

Acute and late toxicities
Grade 2 acute toxicities included hematological toxicities (5 cases), pneumonitis (2), plural pain (1) and dermatitis (1). Grade 2 late toxicities included radiation-induced lung injury (RILI) (3 cases) and asymptomatic rib fracture (1). One patient experienced grade 3 RILI at 6.7 months from the initiation of PT and recovered to grade 1 after hospitalization. No other toxicities of grade ≥3 were observed (Tables 4 and 5).

Discussion
In the current study, by using proper breath control technique and other physical measurements to mitigate the interplay effect and delivery uncertainty, we successfully used PBS technique of PRT or CIRT to treat a group of 31 consecutive and non-selected patients with early stage NSCLC. After a median follow-up time of 12.1 months, we achieved a CSS and LC rates of 95.0% and 95.2% at 2 years with mild acute and late PT-related toxicities in a cohort of patients with stage I and IIa NSCLC (~1/3 of patients with stage IIa disease) and a median age of 71. Furthermore, approximate 80% of the patients had medically inoperable disease and most patients had centrally located lesions (29/31) per RTOG (Radiation Therapy Oncology Group) 0813 protocol [13]. Only 1 patient developed local recurrence due to, at least in part, the relatively low BED_{10} (77.2 GyE) because of poor baseline pulmonary functions. All 3 patients who developed distant metastases had larger tumor size (diameter > 3 cm) with insufficient chemotheraphy. PT-induced toxicities were mild and relieved quickly after proper medical treatment. Only 1 patient who had infectious pneumonia just before the initiation of PT experienced grade 3 RILI and had recovered to grade 1 after hospitalization. PBS-based PT appeared feasible, safe, and effective for early stage NSCLC in a short-term observation.

Radiation therapy is the most important alternative to surgery for unresectable or medically inoperable early stage NSCLC. Radiation therapy with conventional fractionation has generally resulted in inferior LC and OS when compared with surgery. For medically inoperable stage I or II NSCLC, the OS and local failure rates were 24–40% and 25–70% at 2 years, respectively [14]. SBRT achieved better tumor control with more precisely delivered doses to the primary lung lesions with better protection of normal tissues, as compared to conventional RT. Reports [15–17] demonstrated that clinical outcomes after SBRT could be compared to those of surgery with 2.5-year local control rates of 67.9–97%, 73–91.9% and 2.5-year overall survival rates of 48–90%, 43–55.7%. However, for central/ultra-central type lesions, SBRT is of high risk for severe toxicities. Grade 3 toxicities could reach as high as 38% and fatal pulmonary hemorrhage (grade 5) could be even up to 15% [18, 19]. In this study, only 1 patient developed grade 3 RILI among 7/31 patients with ultra-central type lesions.

| Table 1 Characteristics of patients | Number of Patients |
|-------------------------------------|--------------------|
| Age (year)                          | 71 (50–80)         |
| Gender                              |                    |
| Male                                | 20                 |
| Female                              | 11                 |
| ECOG score                          | 0/1                |
| Pathology                           |                    |
| SCC/Adno/ Unclassified/None         | 7/14/4/6           |
| Primary tumor size, median (range)  | 3.1 (1.1–4.7) cm   |
| Primary tumor volume, median (range)| 19.39 (0.97–56.23) cm³ |
| Stage (AJCC staging v7) I/IIa       | 21/10              |
| T1aN0M0                             | 4                  |
| T1bN0M0                             | 7                  |
| T2aN0M0                             | 10                 |
| T1aN1M0                             | 1                  |
| T1bN1M0                             | 4                  |
| T2aN1M0                             | 5                  |
| Tumor location                      |                    |
| Left/right lung                     | 9/22               |
| Upper/middle/lower lobe             | 19/2/10            |
| Peripheral/median/central type-SPHIC institutional definition | 3/15/13 |
| Smoking history (packyears), median (range) | 15 (0–150) |
| Refusal of surgery                  | 8                  |
| Medically inoperable                | 23                 |

Abbreviations: ECOG Eastern Cooperative Oncology Group, SCC Squamous cell carcinoma, Adno Adenocarcinoma, GTV Gross tumor volume, AJCC American Joint Committee on Cancer, SPHIC Shanghai Proton and Heavy Ion Center
Due to their physical characteristics, particle beams provide superior dose localization for deep-situated tumor. The beams penetrate body and abruptly stop to form a Bragg peak where most of the energy was released. Dosimetric studies have shown that PT not only provides comparable or even better coverage of the PTV relative to standard SBRT plans, but also can provide better protection for OARs such as lungs, heart, spinal cord and esophagus \[20–22\]. In addition to the physical dosimetric advantages, carbon-ion is also featured with high linear energy transfer (LET) thus higher relative biological effect (RBE) and lower oxygen enhancement ratio compared to photon. The reported 2-year LC and OS rates of proton therapy reached about 80–97% and 74–97.8%, respectively, which are comparable to SBRT with a generally lower incidence of severe toxicities \[23–25\]. For CIRT, Japanese studies showed a 3-year LC and OS of 82% and 75%, 5-year LC of 90–94.7% and OS of 45–50% without any grade ≥3 pulmonary toxicities \[26–28\].

PBS technology can provide more conformal dose distribution to targets and spare the normal tissues better theoretically, and is considered the future trend of technology development for PT \[3, 4\]. However, all studies published so far on early stage NSCLC’s particle therapy used passive scattering technique except several patient cases \[5\]. The major concerns for using PBS technology in the management of intra-thoracic lesion(s) are inaccuracy of dose distribution caused by inter- and intra-fractional organ motions (interplay effect), significant difference of CT densities among multiple organs and the stopping power uncertainty of particle beam which is more obvious when tumor size or anatomy changes during the treatment \[11\]. In our center, we followed up almost all the requirements in the

| Characteristics of treatments | Number of Patients |
|-------------------------------|-------------------|
| **Breath control techniques** |                   |
| Gating                        | 20                |
| Active breathing control      | 7                 |
| Free Breathing                | 4                 |
| **Prescriptions**             |                   |
| 5–7.5 GyE X 8–10 Fx           | 12                |
| 4–5 GyE X 15–16 Fx            | 3                 |
| 2.25–3.5 GyE X 22–31 Fx       | 16                |
| BED_{10} (GyE), median (range)| 96 (75–119)       |
| **Radiation techniques/sources** |                |
| PRT only                      | 6                 |
| PRT + CIRT boost              | 6                 |
| CIRT only                     | 19*               |
| **Chemotherapy**              |                   |
| 1 cycle (neoadjuvant of pemetrexed+ cisplatin) | 1 |
| 3 cycles (concurrent of Taxol+ cisplatin) | 1 |
| 4 cycles (neoadjuvant of etoposide+ cisplatin) | 1 |
| 4 cycles (neoadjuvant of pemetrexed X 3+ docetaxel X 1) | 1 |
| 5 cycles (neoadjuvant of gemcitabine+ cisplatin X 3 + concurrent of Taxol+ cisplatin X 2) | 1 |
| 6 cycles (neoadjuvant of gemcitabine+ cisplatin) | 1 |

**Table 2** Characteristics of treatments

**Table 3** The mean values of doses to thoracic OARs of all patients (GyE)

|                          | MLD | Lungs V5 (%) | Lungs V20 (%) | MHD | MED | Dmax of Eso | Dmax of SC | Dmax of MBT |
|--------------------------|-----|--------------|---------------|-----|-----|-------------|------------|-------------|
| **Mean**                 | 6.31| 18.42        | 11.13         | 1.88| 4.24| 29.28       | 13.65      | 46.01       |
| **SD**                   | 2.80| 7.45         | 5.77          | 2.56| 5.58| 25.65       | 14.12      | 29.06       |

**Abbreviations:** MLD Mean lung dose, V5/V20 The volume of lung parenchyma that received 5/20 Gy or more, MHD Mean heart dose, MED Mean esophageal dose, Dmax Maximum dose, Eso Esophagus, SC Spinal cord, MBT Main bronchus tree, SD Standard deviation
consensus by Chang et al. [11] except for those we could not accomplish because of the feature of Syngo® system. However, we took several steps to increase our plans’ robustness as mentioned in the ‘methods’ part. This paper was intended to present the promising preliminary results of our study.

PBS technique used for lung cancer was controversial with the concerns of the interplay effect. To the best of our knowledge, this is the first report in early stage NSCLC using PBS technique PRT and CIRT for the entire patient cohort. With the breath control techniques and other physical planning methods used, the 2-year CSS and LC rates reached 95.0% and 95.2% for whole cohort with mild toxicities. Considering the relatively unfavorable disease characteristics in this

| Table 4 Acute toxicities of the entire cohort |
|--------------------------------------------|
| Acute toxicities  | Grade 1 | Grade 2 | Grade 3-5 |
|-------------------|---------|---------|-----------|
| Dermatitis        | 3       | 1       | 0         |
| Esophagitis       | 3       | 0       | 0         |
| Pneumonitis       | 3       | 2       | 0         |
| Plural pain       | 1       | 1       | 0         |
| Cough             | 2       | 0       | 0         |
| Leukocytopenia    | 4       | 3       | 0         |
| Neutropenia       | 4       | 2       | 0         |
| Anemia            | 3       | 0       | 0         |
| Thrombocytopenia  | 1       | 0       | 0         |

| Table 5 Late toxicities of the entire cohort |
|--------------------------------------------|
| Late toxicities  | Grade 1 | Grade 2 | Grade 3-5 |
|-------------------|---------|---------|-----------|
| RILI              | 11      | 3       | 1         |
| Rib fracture      | 0       | 1       | 0         |
| Pleural effusion  | 2       | 0       | 0         |

Abbreviation: RILI Radiation-induced lung injury

Fig. 2 The estimated curves of local control and various survival rates. Panel a shows that of progression-free survival (PFS), panel b local control (LC), panel c cause-specific survival (CSS), and panel d overall survival (OS)
cohort, our results in terms of disease control and OS are very promising compared to previous reports using SBRT, PS-PT or PS-CIRT. Higher BED should be administered under a prospective dose escalation study to reduce local failure. Better results could be expected with optimal systemic therapy such as target therapy and/or immunotherapy.

There were some limitations of this study. First, although we collected the clinical data prospectively, it is still a retrospective study with a small number of patients. Second, the particle resources (proton or carbon ion) and prescriptions used for these 31 patients varied, thus complicated the statistical analyses in a study with a small sample size. Prospective phase I/II clinical trials per different tumor locations have been carrying out for dose escalation of pure carbon ion beam in patients with early stage NSCLC. Further investigation and follow-up of particle therapy using PBS technique are warranted to better define its role in early stage NSCLC.

Conclusions

To the best of our knowledge, this is the first report in early stage NSCLC using proton and/or carbon ion beams with pencil beam scanning technique. Within a short-time follow-up, PBS-based PT achieved safe and effective results for early stage NSCLC. Further follow-up and investigation is warranted.

Abbreviations

4-D: 4-dimensional; ABC: Active breathing control; Adm: Adenocarcinoma; AJCC: American Joint Committee on Cancer; BED: Biological effective dose; BED$_{10}$: Biological effective dose (α/β = 10); CIRT: Carbon ion radiotherapy; CSS: Cause-specific survival; CT: Computed tomography; CTCAE: Common Terminology Criteria for Adverse Events; CTV: Clinical target volume; ECOG: Eastern Cooperative Oncology Group; FB: Free breathing; FDG: Fluorodeoxyglucose; PAWM: Full width at half maximum; GTV: Gross tumor volume; GyE: Equivalent dose to Gy of photon; IGTV: Internal gross tumor volume; IMPT: Intensity modulated particle therapy; IMRT: Intensity modulated radiation therapy; IRB: Institutional Review Board; LC: Local control; LET: Linear energy transfer; MDACC: M.D. Anderson Cancer Center; NSCLC: Non-small cell lung cancer; OARs: Organs at risk; OS: Overall survival; PBS: Pencil beam scanning; PET/CT: Positron Emission Tomography/Computed Tomography; PFS: Progression-free survival; PRT: Proton radiotherapy; PS: Passive scattering; PT: Particle therapy; PTV: Planning target volume; RBE: Relative biological effect; RECIST: Response Evaluation Criteria in Solid Tumors; RILI: Radiation-induced lung injury; RT: Radiation therapy; RTOG: Radiation Therapy Oncology Group; SBO: Single beam optimization; SBRT: Stereotactic body radiotherapy; SCC: Squamous cell carcinoma; SPHIC: Shanghai Proton and Heavy Ion Center

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Availability of data and materials

The datasets generated and/or analyzed during the current study are not publicly available due to hospital policy but are available from the corresponding author on reasonable request.

Authors’ contributions

JC, JL, GI and JM designed the study. JC, NM, JZ and JM collected the data and performed the statistical analyses. JC, JL, CC, MF, GJ and JM acquired funding. JC, NM and JZ drafted the manuscript. JL, CC, MF, GI and JM edited the manuscript. All authors contributed to revising and giving final approval to the manuscript.

Ethics approval and consent to participate

This retrospective study was approved by the institutional review board and written informed consents from all patients were obtained.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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References

1. Mirmanoff R-O. Stereotactic ablative body radiotherapy (SABR): an alternative to surgery in stage I-II non-small-cell cancer of the lung? Chin Clin Oncol. 2015;2(4):42. https://doi.org/10.3978/j.issn.2304-3865.2015.11.02.

2. Chang JY, Jabbour SK, De Ruyscher D, Schild SE, Simone CB, Rengan R, et al. Consensus statement on proton therapy in early-stage and locally advanced non-small-cell lung cancer. Int J Radiat Oncol Biol Phys. 2016;95:505–16.

3. Macdonald OK, Kruse JJ, Miller JM, Garces YI, Brown PD, Miller RC, et al. Prospective phase I/II clinical trial of intensity modulated proton therapy for thoracic malignancies. Int J Radiat Oncol Biol Phys. 2009;75:950–61. https://doi.org/10.1016/j.ijrobp.2009.04.023.

4. Register SP, Zhang X, Mohan R, Chang JY. Proton stereotactic body radiotherapy for clinically challenging cases of centrally and superiorly located stage I non-small-cell lung cancer. Int J Radiat Oncol Biol Phys. 2011;80:1015–22. https://doi.org/10.1016/j.ijrobp.2010.03.012.

5. Chang JY, Li H, Zhu XP, Liao Z, Zhao L, Liu A, et al. Clinical implementation of intensity modulated proton therapy for thoracic malignancies. Int J Radiat Oncol Biol Phys. 2014;90:0808–18. https://doi.org/10.1016/j.ijrobp.2014.07.045.

6. Ettinger DS, et al. Non-small cell lung cancer. In: NCCN clinical practice guidelines in oncology (NCCN guidelines®). 2017. https://www.nccn.org/professionals/physician_gls/default.aspx#nscl. Accessed 15 Oct 2017.
7. Chang JY, Komaki R, Wen HY, De Gracia B, Bluett JB, McAleer MF, et al. Toxicity and patterns of failure of adaptive/ablative proton therapy for early-stage, medically inoperable non-small cell lung cancer. Int J Radiat Oncol Biol Phys. 2011;81:350–7. https://doi.org/10.1016/j.ijrobp.2010.04.049.

8. Chang JY, Li Q-Q, Xu Q-Y, Allen PK, Rebueno N, Gomez DR, et al. Stereotactic ablative radiation therapy for centrally located early stage or isolated parenchymal recurrences of non-small cell lung cancer: how to fly in a “no fly zone”. Int J Radiat Oncol Biol Phys. 2014;88:1120–8. https://doi.org/10.1016/j.ijrobp.2014.01.022.

9. Colaco RJ, Huh S, Nichols RC, Morris CG, D’Agostino H, Flampouri S, et al. Dosimetric rationale and early experience at UFPTI of thoracic proton therapy and chemotheraphy in limited-stage small cell lung cancer. Acta Oncol. 2013;52:506–13. https://doi.org/10.3109/0284186X.2013.769063.

10. Gomez DR, Gillin M, Liao Z, Wei C, Lin SH, Swanick C, et al. Phase 1 study of dose escalation in hypofractionated proton beam therapy for non-small cell lung cancer. Int J Radiat Oncol Biol Phys. 2013;86:665–70. https://doi.org/10.1016/j.ijrobp.2013.03.035.

11. Chang JY, Zhang X, Knopp A, Li H, Mori S, Dong L, et al. Consensus guidelines for implementing pencil-beam scanning proton therapy for thoracic malignancies on behalf of the PTCOG thoracic and lymphoma subcommittee. Int J Radiat Oncol Biol Phys. 2017;99:41–50. https://doi.org/10.1016/j.ijrobp.2017.05.014.

12. Richter D, Graeff C, Jäkel O, Combs SE, Durante M, Bert C. Residual motion mitigation in scanned carbon ion beam therapy of liver tumors using enlarged pencil beam overlap. Radiother Oncol. 2014;113:292–5. https://doi.org/10.1016/j.radonc.2014.11.020.

13. Andrea B, et al. Stereotactic Body Radiation Therapy in Treating Patients with Stage I Non-Small Cell Lung Cancer. https://clinicaltrials.gov/ct2/show/NCT00750269?term=RTDg=0813&rank=1. Accessed 3 Sept 2018.

14. Rowell NP, Williams CJ. Radical radiotherapy for stage VII non-small cell lung cancer in patients not sufficiently fit for or declining surgery (medically inoperable): a systematic review. Thorax. 2001;56:628–38.

15. Bolly G, Filion É, Rakovich G, Kopek N, Tremblay L, Samson B, et al. Stereotactic ablative radiation therapy for the treatment of early-stage non-small-cell lung cancer: CEPO review and recommendations. J Thorac Oncol. 2015;10:872–82. https://doi.org/10.1097/JTO.0000000000000524.

16. Sun B, Brooks E, Komaki R, Liao Z, Jeter MD, McAleer MF, et al. 7-year follow-up after stereotactic ablative radiotherapy for patients with stage I non-small cell lung cancer: results of a phase 2 clinical trial. Cancer. 2017;123:3031–9. https://doi.org/10.1002/cncr.30693.

17. Nagata Y, Kimura T. Stereotactic body radiotherapy (SBRT) for stage I lung cancer. Jpn J Clin Oncol. 2018;48:405–9. https://doi.org/10.1093/jco/hyy034.

18. Corradetti MN, Haas AR, Rengan R. Central-airway necrosis after stereotactic body-radiation therapy. N Engl J Med. 2012;366:2327–9. https://doi.org/10.1056/NEJMoa1203770.

19. Tekabi H, Haasbeek N, Dahele M, De Gracia B, Bluett JB, McAleer MF, et al. Outcomes of Hypofractionated high-dose radiotherapy in poor-risk patients with “Ultracentral” non-small cell lung cancer. J Thorac Oncol. 2016;11:1081–9. https://doi.org/10.1016/j.jto.2016.03.008.

20. Hoppe BS, Huh S, Flampouri S, Nichols RC, Oliver KR, Morris CG, et al. Double-scattered proton-based stereotactic body radiotherapy for stage I lung cancer: a dosimetric comparison with photon-based stereotactic body radiotherapy. Radiat Oncol. 2015;10:425–30. https://doi.org/10.1186/s13014-015-0434-2.

21. Ebara T, Shimada H, Kawamura H, Shinai K, Saito JH, Kawashima M, et al. Dosimetric analysis between carbon ion radiotherapy and stereotactic body radiotherapy in stage I lung cancer. Anticancer Res. 2014;34:5099–104.

22. Shinai K, Kawashima M, Satoh JH, Abe T, Fukata K, Shigeta Y, et al. Clinical outcomes using carbon-ion radiotherapy and dose-volume histogram comparison between carbon-ion radiotherapy and photon therapy for T2b-4N0M0 non-small cell lung cancer-a pilot study. Amendoa R, editor. Public Library of Science. PLoS ONE. 2017;12:e0175589. https://doi.org/10.1371/journal.pone.0175589.

23. Nihei K, Ogino T, Ishikura S, Nishimura H. High-dose proton beam therapy for stage I non-small cell lung cancer. Radiat Oncol Biol. 2006;65:107–11. https://doi.org/10.1016/j.ijrobp.2006.10.031.

24. Hata M, Tokuyue K, Kagei S, Sugahara S, Nakahama H, Fukumitsu N, et al. Hypofractionated high-dose proton beam therapy for stage I non-small cell lung cancer: preliminary results of a phase II clinical study. Radiat Oncol Biol. 2007;68:786–93. https://doi.org/10.1016/j.ijrobp.2006.12.063.

25. Nakayama H, Sugahara S, Tokita M, Satoh H, Tsuboi K, Ishikawa S, et al. Proton beam therapy for patients with medically inoperable stage I non-small-cell lung cancer at the university of Tsukuba. Int J Radiat Oncol Biol Phys. 2010;78:667–71. https://doi.org/10.1016/j.ijrobp.2009.07.1707.

26. Miyamoto T, Baba M, Sugane T, Nakajima M, Yashiro T, Kagei K, et al. Carbon ion radiotherapy for stage I non-small cell lung cancer using a regimen of four fractions during 1 week. J Thorac Oncol. 2007;2:916–26. https://doi.org/10.1097/JTO.0b013e3181560a68.

27. Miyamoto T, Baba M, Yamamoto N, Koto M, Sugawara T, Yashiro T, et al. Curative treatment of stage I non-small-cell lung cancer with carbon ion beams using a hypofractionated regimen. Int J Radiat Oncol Biol Phys. 2007;67:750–8. https://doi.org/10.1016/j.ijrobp.2006.10.006.

28. Iwata H, Murakami M, Demizu Y, Miyawaki D, Terashima K, Niwa Y, et al. High-dose proton therapy and carbon-ion therapy for stage I non-small cell lung cancer. Cancer. Wiley Subscription Services, Inc, A Wiley Company. 2010;116:2476–85. https://doi.org/10.1002/cncr.24998.