Survey of the Patterns of Using Stereotactic Ablative Radiotherapy for Early-Stage Non-small Cell Lung Cancer in Korea

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*Purpose*
Stereotactic ablative radiotherapy (SABR) is an effective emerging technique for early-stage non-small cell lung cancer (NSCLC). We investigated the current practice of SABR for early-stage NSCLC in Korea.

*Materials and Methods*
We conducted a nationwide survey of SABR for NSCLC by sending e-mails to all board-certified members of the Korean Society for Radiation Oncology. The survey included 23 questions focusing on the technical aspects of SABR and 18 questions seeking the participants’ opinions on specific clinical scenarios in the use of SABR for early-stage NSCLC. Overall, 79 radiation oncologists at 61/85 specialist hospitals in Korea (71.8%) responded to the survey.

*Results*
SABR was used at 33 institutions (54%) to treat NSCLC. Regarding technical aspects, the most common planning methods were the rotational intensity-modulated technique (59%) and the static intensity-modulated technique (49%). Respiratory motion was managed by gating (54%) or abdominal compression (51%), and 86% of the planning scans were obtained using 4-dimensional computed tomography. In the clinical scenarios, the most commonly chosen fractionation schedule for peripherally located T1 NSCLC was 60 Gy in four fractions. For centrally located tumors and T2 NSCLC, the oncologists tended to avoid SABR for radiotherapy, and extended the fractionation schedule.

*Conclusion*
The results of our survey indicated that SABR is increasingly being used to treat NSCLC in Korea. However, there were wide variations in the technical protocols and fractionation schedules of SABR for early-stage NSCLC among institutions. Standardization of SABR is necessary before implementing nationwide, multicenter, randomized studies.

*Key words*
Non-small cell lung carcinoma, Stereotactic body radiotherapy, Clinical practice pattern, Surveys and questionnaires
Introduction

Surgical resection has long been the mainstay treatment option for early-stage non-small cell lung cancer (NSCLC). In recent decades, the introduction of stereotactic ablative radiotherapy (SABR) has provided an opportunity to cure NSCLC in patients with high surgical risks. SABR showed comparable local control with minimal toxicity compared with surgical resection in a retrospective study [1,2]. However, owing to its short history of development, there are still some controversies regarding the best strategies for SABR, including the optimal indication, dose, fractionation, planning method, and management of respiratory motion. In many countries, several studies have investigated the differences in the patterns of using SABR for NSCLC [1-3]. These differences in practice are potential hurdles to implementing multicenter trials. The purpose of the current study was to understand the current practices of using SABR for early-stage NSCLC in Korea. The lung cancer subcommittee of the Korean Society for Radiation Oncology intends to use these results to develop new protocols and standards for multicenter trials.

Materials and Methods

1. Survey design

This three-part survey was developed by the lung cancer subcommittee of the Korean Society for Radiation Oncology. SABR was defined as hypofractionated radiotherapy (1-8 fractions). The full questionnaire is provided in Supplementary. The first part of the survey recorded the responder and the institution characteristics, including the location; demographic characteristics of radiation oncologists; lung specialists, and medical physicists; how long SABR has been used to treat NSCLC; and the annual number of patients who undergo SABR for NSCLC. The second part of the survey focused on technical aspects of SABR for NSCLC, including the equipment used, planning systems, immobilization and motion-management methods, simulation techniques, image guidance, and quality assurance. The third part of the survey included questions about the clinical decisions and the choice of treatment modality for specific clinical scenarios. Some of the questions allowed multiple responses to reflect diverse opinions.

2. Responses and statistics

The survey was e-mailed to all board-certified members of the Korean Society for Radiation Oncology. The e-mail was sent four times between June 2014 and January 2015 to collect as many responses as possible. The survey results were collected and analyzed on March 2015. A total of 79 radiation oncologists responded, from 61 of the 85 institutions (71.8%) in South Korea that have a radiation oncology department. Statistical analyses were performed using SPSS software (release 18.0.0, SPSS Inc., Chicago, IL). The Χ² test and Fisher exact test were used to identify differences among groups. A two-sided p-value of ≤ 0.05 was considered statistically significant.

Results

1. Characteristics of the institutions and radiation oncologists

Of the 61 institutions that responded to survey, 33 used SABR for NSCLC and were evaluated in this study. There were 28 academic institutions (85%) and five non-academic institutions (15%). Nineteen institutions (58%) were located in Seoul and its metropolitan areas. There were one or two board-certified radiation oncologists at each of 13 institutions (40%), three at each of 12 institutions (36%), and ≥ 3 at each of eight institutions (24%). Most institutions had one radiation oncologist who had specialized in lung cancer (64%) together with one medical physicist (61%). Eleven institutions (33%) had used SABR to treat NSCLC for < 2 years and 10 (30%) had used SABR for ≥ 5 years. The time that SABR was first used to treat NSCLC is shown in Fig. 1. Overall, 37 radiation oncologists with experience of using SABR for lung cancer completed the survey and provided evaluable responses. Twenty-two of the respondents (59%) worked in Seoul and its metropolitan areas. The characteristics of the radiation oncologists are shown in Table 1.

2. Technical aspects and follow-up policy

Multiple responses were allowed for questions on the technical aspects of SABR. Rotational intensity-modulation radiotherapy (IMRT) (59%) was the most common planning method followed by static IMRT (49%). Immobilization methods included the wing board and vacuum lock system, which were used by 54% and 51% of respondents, respectively. For motion management, 54% of the respondents used abdominal compression to control target movement and 51%
used respiratory gating. The technical aspects of SABR are summarized in Table 2. With a fractionated schedule, 49% of respondents treat all patients on consecutive days, 38% used a more prolonged schedule, and 13% used both schedules. Most of the respondents (86%) obtained the first follow-up images at 4-7 weeks after SABR, and the most common imaging modalities were computed tomography (CT) (78%), followed by positron emission tomography (PET) (11%), and both CT and PET (11%).

3. Clinical decisions

The surveyed oncologists were asked whether they think it is necessary to obtain pathological confirmation before performing SABR in cases with a high risk of procedure complications or technical difficulties, if clinical T1 or T2 lung cancer is most likely. The oncologists frequently (46%) recommended that patients should undergo pathologic confirmation by biopsy before performing SABR. The other options included that SABR could be performed only after CT or PET studies had been completed (38%), and that SABR could be performed after a close imaging follow-up (14%). The
Fig. 2. (A) A 75-year-old male with peripherally located stage I, cT1 (1.6 cm), N0 non-small cell lung cancer (NSCLC) (arrow). (B) A 75-year-old male with medically inoperable stage I, cT2 (4.5 cm), N0 NSCLC (arrow).

Fig. 3. Treatment options chosen for each case scenario. SABR, stereotactic ablative radiotherapy.

respondents were also asked how they would handle suspicious mediastinal or hilar lymph node enlargements diagnosed as reactive nodes by CT or PET. Fifty-four percent of the responders recommended pathologic confirmation before SABR, 41% recommended close observation with imaging modalities after SABR, 3% responded that they would not perform SABR in this case, and one clinician would perform SABR in a T1 case and recommend pathologic confirmation in a T2 case. We also asked the respondents to describe how pulmonary function test (PFT) results influence their decision to perform SABR. Most of the respondents (62%) thought that the patient’s clinical condition is more important than the PFT results, while 14% of the respondents stated that they would set limitations on PFT results which would not consider SABR.
4. Case management

Case 1P was a 75-year-old male with stage IcT1 (1.6 cm), N0 NSCLC located peripherally to the mediastinum and 2.2 cm from the chest wall (Fig. 2A). We assumed that he was diagnosed with adenocarcinoma, had good performance status (Zubrod 1), but his pulmonary function was too poor to perform lobectomy. Nearly all of the respondents (94%) recommended SABR and the most frequently selected fractionation schedule was 60 Gy in four fractions (46%) followed by 48 Gy / four fractions (19%) and 60 Gy / three fractions (16%).

Case 1C was identical to case 1P, except the tumor was located < 2 cm from the mediastinum. For case 1C, SABR was recommended by 43% of the respondents and hypofractionated conventional radiotherapy was recommended by 41%. The most frequently selected fractionation schedule was 60 Gy in 10 fractions, chosen by 14% of respondents; however, there was little consensus with regard to the fractionation schedule. Other schedules used included 60 Gy / 15 fractions (11%) and 50 Gy / five fractions (8%).

Case 2P was a 75-year-old male with stage IcT2 (4.5 cm), N0 NSCLC with a performance status of Zubrod 1, but his pulmonary function was poor and he was scheduled to undergo lobectomy (Fig. 2B). Most of the respondents recommended radiotherapy alone (76%). Concurrent chemoradiotherapy and sublobar resection were recommended by 14% and 5%, respectively. SABR was selected as the treatment of choice by 59% of respondents and hypofractionated radiotherapy by 19% of respondents. The most frequently selected fractionation schedule was 60 Gy in four fractions (32%).

Case 2C was identical to 2P, except that it was central NSCLC (< 2 cm from the mediastinum). Radiotherapy alone, concurrent chemoradiotherapy, and sublobar resection were recommended by 76%, 16%, and 8% of respondents, respectively. Only 19% of respondents considered SABR as the treatment of choice; most recommended hypofractionated conventional radiotherapy (44%). The treatment options selected for each case are shown in Fig. 3.

Discussion

In this nationwide survey, we assessed the current practices of using SABR to treat early-stage NSCLC in Korea. SABR for NSCLC was rapidly incorporated into clinical practice in Korea (Fig. 1). Most of the institutions started SABR for NSCLC within the last 5 years. The majority of the radiation oncologists (72%) with experience of SABR used this technique in < 10 NSCLC patients per year. This low volume suggests that we should conduct multicenter trials rather than single-center trials in future studies of SABR for NSCLC.

The technical aspects of SABR varied widely among the institutions and could represent potential hurdles to multicenter trials. In particular, there were marked variations in the planning techniques and methods used to manage respiratory motion. Accordingly, it will be necessary to standardize these factors before commencing multicenter trials. Most institutions used modern imaging modalities with radiotherapy, including four-dimensional CT simulation, IMRT, and volumetric image guidance. These rapid adaptations of modern techniques make it feasible to perform high-quality trials.

Although SABR has been rapidly integrated into clinical practice, there are some issues to address. One of the issues is mandatory pathologic examination before treatment in high-risk patients. In our survey, 46% of the respondents sought pathologic confirmation before performing SABR. Another issue is the value of mediastinal staging. More than half of the respondents stated that pathologic confirmation was necessary before SABR in cases with reactive mediastinal nodes. Of course, extensive diagnostic staging procedures will be required in multicenter trials.

Currently, various fractionation schedules are used for SABR, including 20 Gy×3, 18 Gy×3, 16 Gy×3, 15 Gy×4, 12 Gy×4, and 12 Gy×5 fractions [4]. To compare the efficacy or toxicity of two different dose fractionation schemes, we conventionally use the biologically effective dose (BED), which is calculated from a linear quadratic model. It has also been adopted in studies comparing SABR fractionation schemes. It has been widely debated whether we should apply a LQ model to SABR, but multiple studies have revealed that radiotherapy doses exceeding a BED of 100 Gy or 105 Gy increase the local control rate and survival outcome, which suggests that the LQ model is suitable for comparing conventional dose schemes [5-9]. However, there is no consensus regarding the ideal fractionation schedule for SABR of NSCLC. In the present survey, even in cases with peripheral cT1 NSCLC, the most popular fractionation regimen (60 Gy in four fractions) was used by less than half of the respondents. This might be due to the national health insurance program of South Korea, which covers SABR only when SABR is given in four fractions. In T2 or central cases, the fractionation schedule varied considerably. Accordingly, it will be necessary to standardize the dose and fraction schedule for SABR before starting future trials.

We investigated the opinions of radiation oncologists regarding their use of SABR in cases with centrally located and/or large (cT2) cancers. Results of a phase II trial performed at the University of Indiana revealed a trend toward increased grade 3-5 toxicity in central tumors (27.3% vs.
10.4%, p=0.088) [10], raising concern of increased toxicity of SABR for centrally located NSCLC. Accordingly, fewer respondents recommended SABR for central NSCLC than for peripheral NSCLC in the current survey (Fig. 3). In terms of tumor size, T2 NSCLC was associated with a lower local control rate than T1 NSCLC following SABR in prior studies [11,12]. The part of the survey focusing on preferred treatments for T2 NSCLC revealed that concurrent chemotherapy was more common in T1 cases than in T2 cases. Many radiation oncologists hesitated to use SABR for centrally located T2 NSCLC. Only 19% of the respondents stated that they would perform SABR in cases with centrally located T2 NSCLC. Because of the varying influences of location and size of the cancer on clinical practices, clinical trials should include stratified protocols to account for these factors. Our study has several limitations. First, not all of the board-certified radiation oncologists in Korea responded to the survey, which might introduce selection bias. In addition, because multiple responses were allowed for some of the technical questions, we may overestimate the rate of infrequently used techniques. Finally, although we sought to integrate most of the approaches used in actual practice, some practices were not reported because the list of responses was incomplete.

**Conclusion**

In conclusion, this nationwide survey confirmed that the use of SABR for treating early-stage NSCLC has increased dramatically in Korea in recent years. The technical aspects and clinical decisions for SABR for treating NSCLC varied markedly. The treatment patterns of centrally located T2 NSCLC varied much more than those of peripherally located T1 NSCLC. These results highlight the need to standardize various aspects of the protocol when planning nationwide, multicenter, clinical trials.

**Electronic Supplementary Material**

Supplementary materials are available at Cancer Research and Treatment website (http://www.e-crt.org).

**Conflicts of Interest**

Conflict of interest relevant to this article was not reported.

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