Stereotactic ablative body radiotherapy (SABR): an alternative to surgery in stage I-II non-small-cell cancer of the lung?

René-Olivier Mirimanoff

Department of Radiation Oncology, Clinique de La Source, Lausanne CH-1004, Switzerland
Correspondence to: René-Olivier Mirimanoff, MD. Clinique de La Source (CLS), Avenue Vinet 30, CH-1004 Lausanne, Switzerland.
Email: rene-olivier.mirimanoff@chuv.ch.

Author’s introduction: René-Olivier Mirimanoff obtained his MD degree at the University of Geneva, Switzerland. After residencies in medicine and oncology in Philadelphia (USA), Lausanne and Geneva (Switzerland), he did his full residency and fellowship in radiation oncology at Massachusetts General Hospital (MGH, Harvard), in Boston (USA), where he graduated and became US board certified in 1984. After being on the staff in Geneva, he was appointed as Professor and Chairman in Radiation Oncology (RO) at CHUV (University Hospital) in Lausanne in 1988. Under his direction, his Department underwent major changes, with full academic staffing and cutting-edge equipment, to become a significant European Organisation for Reasearch and Treatment of Cancer (EORTC) contributor. Since July 2012, he is Emeritus Professor at the University of Lausanne. His research focused on clinical investigation, his main contributions being on brain, lung and rare tumors. His list of publications comprises over 230 papers, of which 187 in peer-reviewed journals, including New Engl J Med, Lancet, Lancet Oncol, J Clin Oncol, Eur J Cancer, Int J Radiat Oncol Biol Phys, Radiother Oncol, and many others. His is member of a number of international societies, reviewer and board member of several journals. He is founder and the first President of the Scientific Society of Swiss Radiation Oncology (SASRO), founder of the “Rare Cancer Network”, former Chairman of the EORTC Radiation Oncology Group and currently an ESTRO Course Teacher. He is a regular reviewer of research grants in the European Community and Switzerland. Dr. Mirimanoff is Honorary Professor at the Chinese Academy of Medical Sciences in Beijing (2005), Honorary Member of SASRO (2005), medalist of the French SFRO (2006), Emmanuel van der Schueren medalist of the Belgian BVRO (2008), Alumnus of the Year (2010) at MGH, Harvard, and received the ESTRO lifetime achievement award in 2010. He will be recipient of the Golden Hippocrates Award in 2016.
Abstract: For decades, surgery was considered to be the only standard therapy for early-stage non-small-cell lung cancer (NSCLC). However, stereotactic ablative body radiotherapy (SABR) has been used in a growing number of patients and institutions since the early 2000s. Initially, this technique was intended mainly for patients who were deemed to be medically inoperable due to co-morbidities or who refused surgery, but more recently, it has been applied to operable patients as well. Strict criteria for treatment planning, the use of high-technology equipment, and the appropriate selection of dose based on tumor size and location are of paramount importance for a proper application of SABR. Under these conditions, SABR offers high control rates with a moderate risk of severe toxicity, quite comparable to those of modern surgery. This article reviews the basic principles of SABR, its practical aspects, the definition of biologically equivalent doses, the results in terms of tumor control, survival, and toxicity, and an attempt will be made to compare the results of SABR with those of surgery.

Keywords: Stereotactic ablative body radiotherapy (SABR); non-small-cell lung cancer (NSCLC); early-stage; local control; survival

Submitted Apr 21, 2015. Accepted for publication Oct 22, 2015.
doi: 10.3978/j.issn.2304-3865.2015.11.02
View this article at: http://dx.doi.org/10.3978/j.issn.2304-3865.2015.11.02

Introduction

In spite of the remarkable progress in the biological understanding, the pathological and clinical diagnosis and in the various treatments of non-small-cell lung cancer (NSCLC), its overall prognosis remains disappointing, even in early stages. For decades, surgery was considered to be the only standard therapy in early disease, however the 5-year survival rates after a surgical resection ranged 60% to 80% in stage I and only 40% to 50% in stage II (1). Surgery, like other curative treatments for NSCLC, including stereotactic ablative body radiotherapy (SABR), (see below) may be followed by important complications, and can even lead to a decline of quality of life (2), especially in elderly patients (3). Old age and the presence of multiple co-morbidities were, and still are, responsible for the undertreatment of early stage NSCLC in a significant proportion of patients. For example in the Netherlands, up to the late 1990s, 32% of patients aged 75 years and older could not receive any curative local treatment for stage I NSCLC (4), but that was in an era before new treatments such as video-assisted thoracoscopic surgery (VATS), radiofrequency ablation, cryosurgery and SABR, were commonly available. However since the past decade, SABR has been more and more used in a large number of patients in a growing number of institutions. It is generally accepted that this technique represents now an alternative to surgery, under well-defined conditions, and can be administered to elderly patients and to patients with multiple co-morbidities, as reported by recent reviews (5-9). The present review is primarily intended for interested chest physicians, thoracic surgeons, medical oncologists, and radiation oncologists not yet experienced in SABR, and who wish to become more familiar with this technique. In this article, the basic principles of SABR, its practical aspects, the definition of dose, and the results including tumor control and toxicity will be reviewed, and an attempt will be made to compare the results of SABR with those of surgery.

Stereotactic ablative body radiotherapy (SABR)

Rationale and indications

Long before the advent of SABR (also called SBRT for stereotactic body radiotherapy), conventional radiotherapy (RT) was sometimes given to patients with early stage NSCLC who were not candidates for surgery for medical reasons or who refused surgery (10). However, conventional RT techniques had their limitations. Sixty Gy, in conventional daily fractionation of 2 Gy, represents a biologically equivalent dose (BED10) of 72, which is clearly insufficient to control a NSCLC in most situations (see below). So even if some inoperable patients could be cured with conventional RT, the overall results were rather poor, with at least 40% local failures and 5-year overall survival (OS) rates of only 15-20% (10,11). Compared to conventionally fractionated RT, which for NSCLC typically delivers a dose of about 60 Gy in daily fractions...
of 1.8 to 2 Gy in 6 weeks, SABR represents a completely different philosophy, which delivers very high doses of highly conformal RT to relatively small volumes in a few days. This technique derives from the principles applied in intracranial stereotactic radiosurgery (SRS), which has been administered for decades for intracranial neoplasms (12), and thus uses rather similar technical tools and rather similar radiobiological principles. Thus SABR delivers very large doses, referred to by Timmerman et al., as “ablative” doses (13) of RT in a few high-dose fractionation schedules, typically in 3 to 8 fractions (see below). Compared to conventional RT, this implies major radiobiological reconsiderations, referred to by Timmerman et al. in another paper as the “hypofractionated revolution” (14). Thus it is critical for the newcomers in the field to have a good understanding of these radiobiological principles, otherwise major problems and complications will be likely to be met. Besides this, in order to apply SABR, one needs to have at hand the most sophisticated technological tools, high competence in physics, imaging, RT planning and RT delivery, to administer safely this high, compact dose to the target. Each treatment should insure a steep gradient of dose for a maximum avoidance of normal sensitive structures, while hitting the target with the highest precision.

**Target definition and treatment planning**

After having confirmed the indication for SABR, the first technical step for planning SABR includes a careful identification of the target with the best currently available imaging tools, including a high-quality CT with appropriate windowing. Then, a planning 4 D CT is obtained, to define not only the gross tumor volume (GTV) and the clinical target volume (CTV), but also and most importantly the internal target volume (ITV, Figure 1), which represents the space occupied by the tumor during the whole respiratory cycle. Depending on the tumor volume and motion the radiation can be applied either (I) to the whole ITV, particularly in case of limited tumor volume and tumor motion, or (II) by using a “gating” technique, in which the irradiation is applied only during part of the respiratory cycle, or (III) using a “tracking” technology, in which the tumor is “followed” by the beam during the respiratory cycle. During the planning procedure, (like for any high-precision RT), it is essential to determine the best treatment plan by optimal dose-volume histograms (DVH). Normal tissue constraints values, which are defined specifically for large fractions, have to be used, and can be found in related papers (13). Although many different technical approaches have been used, to ensure the best distribution of dose high-technology linacs with intensity modulation RT (IMRT), or volumetric arc therapy (V-MAT) and image-guided RT (IGRT) technology, or other tools like the Tomotherapy or the Cyberknife systems, or even the proton-beam technology, have to be used. However at present, there are no data demonstrating the superiority of any of these treatment techniques over any other ones. Stable and reproducible positioning is essential, using either various frame systems to better immobilize the patients, or frameless systems using markers and image-guided systems.

**Definition of dose in SABR**

SABR implies a large total dose in a few fractions. It should be remembered that what is called a radiobiological dose

![Figure 1 Schematic representation of an internal target volume (ITV) with comparative CT scans of the same patient in expiration and inspiration, and treated at our institution (CLS) (courtesy Mrs Bressan RTT, CLS).](image-url)
has not at all the same meaning as a physical dose. Due to the decreased or absent DNA repair when large individual doses per fraction are given to any tissue (tumor or normal tissue), a dose such as 20 Gy given in one fraction is much more efficient than 20 Gy given in 10 fraction (Figure 2). Similarly, 60 Gy in 3 fractions is much more “tumoricidal” (and hugely more toxic!) than 60 Gy given in 30 fractions. Thus, to establish RT protocols with biologically equivalent doses while using different fractionation schedules, various calculation formulae can be used, like the BED equation, where (3,10):

\[
BED = D \frac{\alpha}{\alpha + \beta} + d \alpha / \beta,
\]

where \(BED\) = biologically equivalent dose; 
\(D\) = total dose in Gy; 
\(d\) = dose per fraction in Gy; 
\(\alpha / \beta\) = coefficient for tumors or normal tissues.

Some examples of SABR schedules and one of a conventional schedule of 60 Gy in 30 fractions, and their corresponding BED10 values are displayed on Table 1. One can see that any dose used for SABR is largely superior to that of the conventional RT scheme. The differences are even greater when taking an \(\alpha/\beta\) value of 3 (BED3) for normal tissues. Specific normal tissue tolerance-dose constraints for 1-5 fractions schedules have to be used and recommendations can be found in the literature (13). This explains why, choosing a protocol for tumors in the vicinity of sensitive normal structures, most investigators today will carefully adapt the total dose and the dose per fraction to decrease the risk of major tissue complications (see below).

### Results of SABR

#### Local control and survival

Following SABR for stage I and II NSCLC, several endpoints should be considered to evaluate its success or failure. Local control should ideally be the most important endpoint in SABR, as it should reflect directly the efficacy of the technique, especially when compared to surgery. However the interpretation of the radiographic response is often difficult. A complete disappearance of the tumor is observed only in a minority of patients, and even in case of permanent local control one can still identify some abnormalities, even months later (Figure 3). In a review, 60% to 100% of patients were expected to have radiographic changes after SABR (15). The changes observed on CT could be scored into five categories: (I) diffuse consolidation, (II) patchy consolidation and ground-glass opacities (GGO), (III) diffuse GGO, (IV) patchy GGO, and (V) no evidence of increased density (15). All these can mean permanent local control. FDG-PET may be helpful but inflammatory response may persist more than 12 months (15,16). Interestingly, late radiological changes may differ depending on the SABR technique that was used. Arc-SABR trended towards more pronounced radiological changes, with a different pattern, compared to changes seen after fixed-beams SABR (16). Overall, after SABR, permanent local

### Table 1 Examples of five different schedules used for SABR compared with one conventional schedule of 60 Gy in 30 fractions, and their corresponding BED10 values

<table>
<thead>
<tr>
<th>Total dose in Gy/number of fractions</th>
<th>BED10</th>
</tr>
</thead>
<tbody>
<tr>
<td>48/4</td>
<td>106</td>
</tr>
<tr>
<td>45/3</td>
<td>113</td>
</tr>
<tr>
<td>60/30</td>
<td>72</td>
</tr>
<tr>
<td>60/8</td>
<td>105</td>
</tr>
<tr>
<td>60/5</td>
<td>132</td>
</tr>
<tr>
<td>60/3</td>
<td>180</td>
</tr>
</tbody>
</table>

SABR, stereotactic ablative body radiotherapy; BED, biologically equivalent dose.
control of the tumor is observed in 81% to 97% of treated patients (Table 2) (17-32). The next other major endpoint is survival: in the same series, survival at 3 years was reported to be between 52% and 64% (Table 2) (17-32). Examples of some typical series show fairly consistent results. Onishi et al. have treated 245 patients in 13 Japanese institutions, with a median BED dose of 108 Gy (57-180 Gy) (17). Local progression occurred in 14.5%, and the 3-year-survival was 56%, with a cause-specific survival of 78%, indicating that a significant proportion of deaths were not cancer-related but were due to other co-morbidities (17). Lagerwaard et al. from VUMC, Amsterdam, have reported on 206 patients treated with 3 schedules of 3×20 Gy, 5×12 Gy and 8×7.5 Gy, depending on T stage and proximity of sensitive structures (19) (see also below). Median survival was 34 months, local failures were observed only in 3% and regional failures in 9% (19). Bradley et al. prospectively registered and analyzed 91 patients from Washington University School of Medicine (WUSM) in St Louis, with 3×18 Gy for peripheral tumors and 5×9 Gy for tumors in close vicinity of critical structures (22). Most patients had

**Table 2** Summary of studies on SABR for early stage non-small cell lung cancer (17-32)

<table>
<thead>
<tr>
<th>Series</th>
<th>Year</th>
<th>Patients</th>
<th>LC</th>
<th>2 y S</th>
<th>3 y S</th>
</tr>
</thead>
<tbody>
<tr>
<td>Onishi (17)</td>
<td>2004</td>
<td>245</td>
<td>85%</td>
<td>–</td>
<td>56%</td>
</tr>
<tr>
<td>Xia (18)</td>
<td>2006</td>
<td>43</td>
<td>81%</td>
<td>78%</td>
<td>–</td>
</tr>
<tr>
<td>Lagerwaard (19)</td>
<td>2007</td>
<td>206</td>
<td>81%</td>
<td>–</td>
<td>64%</td>
</tr>
<tr>
<td>Chen (20)</td>
<td>2008</td>
<td>65</td>
<td>88%</td>
<td>–</td>
<td>57%</td>
</tr>
<tr>
<td>Baumann (21)</td>
<td>2009</td>
<td>57</td>
<td>92%</td>
<td>–</td>
<td>60%</td>
</tr>
<tr>
<td>Bradley (22)</td>
<td>2010</td>
<td>91</td>
<td>86%</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Timmerman (23)</td>
<td>2010</td>
<td>59</td>
<td>97%</td>
<td>–</td>
<td>56%</td>
</tr>
<tr>
<td>Ricardi (24)</td>
<td>2010</td>
<td>57</td>
<td>92%</td>
<td>–</td>
<td>60%</td>
</tr>
<tr>
<td>Matsuo (25)</td>
<td>2011</td>
<td>101</td>
<td>93%</td>
<td>80%</td>
<td>–</td>
</tr>
<tr>
<td>Widder (26)</td>
<td>2011</td>
<td>202</td>
<td>95%</td>
<td>72%</td>
<td>–</td>
</tr>
<tr>
<td>Takeda (27)</td>
<td>2012</td>
<td>173</td>
<td>80%</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Shibamoto (28)</td>
<td>2012</td>
<td>180</td>
<td>87%</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Taremi (29)</td>
<td>2012</td>
<td>108</td>
<td>–</td>
<td>63%</td>
<td>–</td>
</tr>
<tr>
<td>Hamamoto (30)</td>
<td>2012</td>
<td>128</td>
<td>87%</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Crabtree (31)</td>
<td>2014</td>
<td>151</td>
<td>97%</td>
<td>–</td>
<td>52%</td>
</tr>
<tr>
<td>Kestin (32)</td>
<td>2014</td>
<td>483</td>
<td>91%</td>
<td>–</td>
<td>–</td>
</tr>
</tbody>
</table>

LC, local control; 2 y S: 2-year survival; 3 y S: 3-year survival.
either a poor performance status or poor lung functions. Two-year local control was achieved in 86% of patients (22). Crabtree et al. have reviewed 151 SABR patients, whom they compared to 458 surgical patients (31) (see below). BED10 varied between 85.5 and 151.2. For the SABR group, 3-year local control was 89% and OS 52% (31).

**Optimal dose of SBRT**

Since the first reports on SABR, many different schedules of dose have been explored, in order to find the best therapeutic ratio, taking into account the best probability of tumor control with a minimal risk of causing major complications.

One initial dose-escalation phase I study was initiated at Indiana University to assess toxicity and local control rates (33). Forty-seven patients were treated with SABR escalating from a starting dose of 24 Gy in three fractions up to 72 Gy in three fractions. Patients were stratified by T stage and tumor size. The maximum tolerated dose (MTD) was 66 Gy in 3 fractions for tumors larger than 5 cm and was not reached for T1 tumors at 60 Gy in 3 fractions or tumors less than 5 cm at 66 Gy in 3 fractions (33). Chi et al. have extensively reviewed and linked radiobiological modeling and clinical outcome from 9 series of patients (34). Their estimates indicate a clear dose-response relationship: for example with BED10 values of 72, 84, 106 (see corresponding doses and fractions in Table 1), the progression-free survival at 30 months (PFS30) is only 15%, 24%, and 34%, respectively (34). With higher BED10, the PFS30 increases markedly: BED10 values of 113 and 125 correspond to a PFS30 of 95% and 99%, respectively and then of course a plateau is reached (34). Beyond these BED10 values, one may question the necessity to administer higher RT doses, as the toxicity of normal tissues increases even more (see below). Kestin et al. have reviewed 505 T1 and T2 NSCLC in 483 patients treated by SABR in 5 institutions in the USA, Germany, The Netherlands and Canada (32). All were treated with on-line image-guidance RT (32). Five different schedules, with a median prescription BED10 of 132 Gy, were used. A clear dose-response relationship for local control was demonstrated, with an optimal BED10 > to 125 Gy (32). Zhang et al. have performed a meta-analysis on 2,587 patients from 34 observational studies (35). BED was divided into 4 dose groups: low (<83.2), medium (83.2-106), medium to high [106-146] and high (>146). As expected, overall 2- and 3-year survivals were higher in the medium dose and medium to high dose groups compared to the low dose group. However, and interestingly, the former two groups were also superior to the high dose group for OS (35). Not all studies however suggest the concept of the dose-response hypothesis in SABR. Van Baardwijk et al. in their systematic overview of 15 studies on SBRT (=SABR) found no correlation between the freedom from local progression and the EQD2,T, or equivalent dose in 2 Gy fractions (36). In any case, it seems clear from most studies that a BED10 beyond a certain value (around 120-130 or so) may not only be unnecessary, as one could estimate from Chi et al.’s data (34), but may be even detrimental, due to an increased toxicity.

**Patterns of failures**

As seen in Table 2, the rate of local control in most series is above 85%. Provided that a careful initial work up is made, to exclude nodal disease, the regional failure rate should also be low (see below). Thus the largest proportion of failures are represented by distant metastases (37). Bradley et al. in their series of 91 patients from WUSM, have reported that the majority of the failures were distant, with 19 of them being distant metastases, with or without some local (7 cases) and/or regional component (22). Distant failures negatively impacted the OS. In addition, 15 patients developed a second primary lung cancer (22). The largest study to date regarding the pattern of failure was published by Senthi et al. from VUMC Amsterdam (38). They have assessed 676 patients treated by SABR between 2003 and 2011 (38). Eighteen percent had a disease recurrence, with an actuarial 2-year rate of local, regional (nodal) and distant recurrence rate of 4.9%, 7.8%, and 14.7%, respectively, and with corresponding rates at 5 years of 10.5%, 12.7% and 19.9%. New pulmonary lesions, or second primary tumors, developed in 6% of all patients (38).

**Toxicity**

With SABR, like with high-dose, conventional RT, there is a potential risk of major complications, such as radiation pneumonitis, oesophagitis and brachial plexopathy. In addition, new severe and sometimes fatal complications have been reported with SABR, including tracheo-oesophageal fistulae, brocho-pulmonary fistulae, cardiotoxicity and chest wall necrosis (39). The latter complication, along
with rib fractures, may be particularly severe and painful. Timmerman et al. in a seminal paper have reported their experience on 70 patients treated for T1 and T2 (<7 cm) with 60-66 Gy in 3 fractions (BED10: 180-211) (40). Significant grade 3 to 5 toxicity occurred in 14 patients, with 6 toxic deaths. Patients with peripheral lung tumors had a 2-year freedom from severe toxicity of 83% versus only 54% for centrally located tumors (40). They have defined an area referred to as the zone of the proximal bronchial tree, in which very high BED doses should be prohibited (40). In this regard, Lagerwaard et al. have designed a “risk-adapted” protocol taking into account tumor size and location (19). T1 peripheral tumors received 60 Gy in 3 fractions (BED10: 180), T1 with broad contact with the thoracic wall or T2 tumors received 60 Gy in 5 fractions (BED10: 132) and tumors adjacent to the heart, hilus and mediastinum received 60 Gy in 8 fractions (BED10: 105) (19). With this protocol, which reduces considerably the biological dose given to the normal “central” organs, severe toxicity was observed only in 3% of all patients, without compromising the overall local control, which was excellent with only 3% crude local failure rate (19).

Figure 4 shows two examples of central or peripheral lesions treated at our institution (CLS), with doses adapted to their location. Sentihi et al. have reviewed the toxicity of SABR based on 20 publications including 563 central lung tumors (5). They confirm that with the above-mentionned precautions, tumor location did not impact OS or toxicity (5).

**Medically operable or inoperable patients**

At its beginning, SABR was intended for patients who were deemed inoperable because of age or multiple co-morbidities and unacceptable surgical risks. However patients who refused surgery were also candidates for this new procedure. Onishi et al. in their series of 245 patients, reported that 158 were considered to be inoperable and 87 to be operable (17). There was a highly significant difference in survival (P<0.01) between the two categories in favor of the operable patients, the latter having a 3-year survival of 88%. In the group of inoperable patients, the rate of intercurrent deaths (deaths from other causes) was 19.1% versus only 3.4% for operable patients (17). This explains at least in part the large difference in OS between the two groups. Lagerwaard et al. found that in their prospective database of SABR, 177 patients (25% of their cases) were deemed potentially operable, using strict criteria (41). In this group of patients, the 1- and 3-year survival rates were 94.7% and 84.7%, respectively, and the local control rates 98% and 93%, respectively (41). Interestingly, in certain circumstances, a biopsy prior to SABR can be omitted. Verstegen et al. have analyzed a fairly large cohort of patients, with (209 patients) or without (382) a biopsy prior to SABR (42). Local control and OS were exactly the same in the two groups of patients (42). To not biopsy certain patients is based on the fact that only 1-4% of FDG-PET positive lesions undergoing surgery are benign (42), and thus with a careful imaging assessment, the risk without a biopsy of treating a benign lesion instead of a cancerous lesion is low.

At this point the question is raised whether SBRT could be an alternative to surgery, even in the most favorable category of patients with early NSCLC.
**Surgery versus SABR for stage I and II NSCLC**

Evidently to compare the efficiency of SABR vis-à-vis surgery, randomized controlled trials are needed. Unfortunately, three major initiatives in the USA and Europe have failed recently, due to poor accrual (43). One has thus to rely on matched-paired analyses, in which carefully matched patients in each comparative groups are analyzed. Verstegen et al. have matched SABR patients and patients treated by VATS from six hospitals in The Netherlands (44). The cohort consisted of 64 SABR and 64 VATS. Post SABR local control rates were superior at 1 and 3 years (96.8% and 93%, versus 86.9% and 82.6% respectively, P=0.04), but distant recurrence and OS were not different (44). Crabtree et al. from WUSM compared 462 surgical patients to 76 receiving SABR, and found that surgical patients were healthier and had a better tumor control compared with those receiving SABR (31). However when they did a propensity analysis, they found that local recurrence and disease-specific survival were similar in the two groups (31). A meta-analysis of six studies containing 864 matched-paired patients was performed by Zhang et al. (45). Pooled data at 1 and 3 years indicate a better long-term OS with surgery. However the rate of cancer deaths was the same in the two groups of patients, which strongly suggests that in spite of the matching of patients, those undergoing SABR may have been less healthy than the surgical patients. This was indirectly demonstrated by the fact that there was no significant difference in cause-specific survival, disease-free survival or local control between the SABR and the surgical patients (45). Solda et al. have reviewed 45 reports containing 3,771 patients treated with SABR and compared them to 2,038 surgical patients (46). They found that the 2-year survival was 70% after SABR versus 68% after surgery (46). As regards performance status (PS) and comorbidity as independent prognostic factors which may be used for treatment decisions, Louie et al. have constructed univariate and multivariate models to establish recursive partitioning analyses (RPA) classes and a nomogram (47). RPA identified two risk classes based on tumor diameter, age, PS and co-morbidity index, but performed poorly in surgical patients, whereas the nomogram retained a strong performance for surgery and SABR (47).

Finally, the enthusiasm generated by all the “positive” data on SBRT should be tempered by a more critical assessment of this new technique. Brada et al. in a recent editorial (48) have expressed a series of reservations vis-à-vis the “overconfidence” and “self-congratulation” around SABR (48). For example they remind that other newer approaches, like new surgical techniques (VATS), radiofrequency or thermal ablation may provide equivalent tumor control as SABR (48). They also underscore that the local control after SABR may be overestimated, given for example the difficulty to assess this endpoint with the current imaging means (see above), and that long-term toxicity may be underestimated as well. They question whether SABR impacts on the natural history of co-morbid situations. They also emphasize that more studies are needed to better define a series of unsolved or insufficiently solved issues, for example on respiratory and cardiac co-morbidities, on the optimal dose and fractionation and on long-term toxicities (48).

**Conclusions**

SABR is now a well-established technique for the treatment of early stage NSCLC, which requires a high quality of the teams and of the techniques to be used. Besides this high technology, a good understanding of the radiobiological principles is of paramount importance, in order to decrease the risk of severe complications. A dose-adapted scheme has to be used in each institution practicing SABR. Probably a BED10 dose over 120-130 is unnecessary for peripheral lesions, and should be even lower for centrally located tumors, probably not beyond 110. Results show at this point a very good local control, and an acceptable toxicity, provided a proper overall evaluation is made and the appropriate biological effective dose is selected. SABR is now a first choice for medically inoperable patients. For operable patients, at the present time, surgery remains the standard, but SABR can be a good second option for patients who refuse surgery. At the present time though, more studies are needed because a number of problems have not been entirely solved and longer follow-ups are required.

**Acknowledgements**

The author would like to thank Mrs Stefania Bressan who has provided figures 1, 3 and 4 in this article.

**Footnote**

*Conflicts of Interest:* The author has no conflicts of interest to...
declare.

References


