Long-term follow-up after first-line bronchoscopic therapy in patients with bronchial carcinoids

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ABSTRACT

Background Carcinoid of the lung is considered to be a low-grade malignancy. A subgroup presents as an endobronchial tumour. Surgical resection is considered the standard approach because of its metastatic potential and the possibility of an iceberg phenomenon for the endobronchial subgroup. Advances in noninvasive and minimally invasive technologies seem to justify a more lung parenchyma-sparing approach.

Methods In patients presenting with bronchial carcinoids, initial bronchoscopic treatment (IBT) is first attempted for complete tumour eradication and sufficient tissue sampling for the proper differentiation of typical (TC) versus atypical (AC) histological type. Furthermore in cases with postobstruction problems the desobstruction is aimed at improving the patient’s condition and by that alleviate surgery if that is needed. High resolution CT is performed 6 weeks post IBT to determine local tumour growth. Surgical resection follows in case of extraluminal disease, residual carcinoid inaccessible for IBT, or late recurrences not salvageable by repeat IBT.

Results Minimum follow-up was 5 years from start of treatment for 112 patients (65 women, 47 men), with a median age of 47 years (range 16–77 years). Eighty-three patients (74%) had TC, and 29 (26%) AC. IBT only was ultimately curative in 42% of the cases (47/112); 42 TC, 5 AC. Disease-specific mortality including surgical mortality has been 2% (3/112) in patients with extraluminal carcinoids (3 AC).

Conclusions IBT, if with unsuccessful rescue surgery, is justifiable with excellent long-term outcome. IBT made surgery unnecessary in 42% of the cases. Iceberg phenomenon and metastatic potential in this group of patients with bronchial carcinoids are clinically insignificant.

INTRODUCTION

Carcinoids represent only a small portion of all pulmonary tumours, that is, 2–5%. They belong to the spectrum of neuroendocrine tumours consisting of tumourlets, typical carcinoids (TCs), atypical carcinoids (ACs), large cell neuroendocrine carcinomas and small cell lung cancers in which TCs represent the relatively benign and small cell lung carcinomas the most malignant entities of this spectrum.1–2 Within this group of carcinoids the initial presentation can be in the parenchyma or mainly in the airway, described as bronchial carcinoids.

TC shows relatively low malignant behaviour with excellent 5-year and 10-year survival rates of 87–100% and 82–87%, respectively. In case of positive lymph nodes in TC, 5-year survival rates are still up to 92–100%. ACs have a higher malignant potential with significantly lower 5-year and 10-year survival rates of 56–75% and 35–56%, respectively. In case of positive lymph nodes 5-year and 10-year survival rates for ACs dropped to 25–78% and 25–59%, respectively.3–6

Distinction between TC and AC is based on the classification criteria by Travis et al,7 which are based on a retrospective analysis of surgically resected specimens. Historically, surgical resection is considered the treatment of choice for both entities mentioned above. Lung parenchyma preserving surgical techniques, such as surgical bronchotomy and sleeve resections are clinically acceptable.4,5,8 Current advancements in diagnostic and therapeutic modalities and further understanding of tumour biology enable and encourage many to explore less or minimally invasive diagnostic and therapeutic techniques for optimal preservation of healthy lung tissue.7,9,10 We have previously reported that initial bronchoscopic treatment (IBT) seems to be a justifiable approach for bronchial carcinoids instead of immediate surgical exploration and resection.11–13 Moreover, recent literature showed favourable results for bronchoscopic treatment (BT) as well, in carefully selected patients.14–16 In this study we report the long-term outcome of our IBT strategy in patients with centrally located bronchial carcinoids.

PATIENTS AND METHODS

We have outlined this approach in great detail in the aforementioned report.13 Figure 1 is a summary of our original IBT approach.

Our current approach is shown in figure 2 and the current approach, in brief, is as follows:


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All patients referred to our institution for suspicion of harbouring a bronchial carcinoid undergo rigid and flexible bronchoscopy in one session for diagnostic purposes and complete resection of the intraluminal tumour growth. Six weeks post IBT for allowing postobstructive changes to resolve, a high resolution CT (HRCT) scan with intravenous contrast and repeat fibre-optic bronchoscopy is performed to assess IBT result. In case of residual extraluminal disease or AC histology or late recurrence not amenable for IBT, surgical resection is performed. In the later phase of this study period (since May 2000), AC successfully eradicated with IBT was subjected to close surveillance and not immediately followed by surgical resection. Successful IBT is defined if there is no sign of residual disease based on detailed bronchial wall images on thin slice HRCT with intravenous contrast and videobronchoscopy with negative biopsies. Close surveillance is by repeat HRCT and fibre-optic bronchoscopy 6-monthly for the first 2 years and annually up to 5 years; thereafter patients have been referred back to their pulmonologists with advice for yearly check-up. For intraluminal carcinoid recurrence during follow-up, BT is again attempted first to obtain tumour eradication prior to the decision for surgical resection.

We reviewed the prospectively gathered data of every referred patient to our centre—since the beginning of our protocol—from 1991 till July 2009. This final analysis includes only those patients with at least 5 years follow-up after their first IBT. Every patient has been given detailed explanations of the pros and cons of our strategy. SPSS V20 was used for data collection purposes.

RESULTS
From 1991 until July 2009, 112 patients have been accrued. For patients alive in July 2014 the median follow-up has been 112 months (range: 60–277 months). Four patients died within the first 5 years after inclusion. Patients’ characteristics are shown in table 1, and IBT results in figure 3 and table 2.

A description of what is shown in figure 3 is further elucidated as follows.

In 37/112 (33%) patients (33 TC, 4 AC) there was no sign of residual tumour (complete remission (CR)) after the first IBT session. In 45 patients (40%: 29 TC, 16 AC) extraluminal growth was immediately apparent and surgical resection was performed. Of the remaining 30 patients (21 TC, 9 AC) with residual disease, 14 (11 TC, 3 AC) had residual intraluminal disease completely eradicated by repeating IBTs (range one to four sessions), one intraluminal residual carcinoid (TC) was inaccessible for IBT: this patient was unfit for surgical resection. The remaining 15 patients (21 TC, 9 AC) extraluminal disease ineligible for BT. Thirteen of them (13/15; 87%) underwent surgical resection (7 TC, 6 AC). Three patients (3 TC) did not undergo surgery: one aforementioned patient had intraluminal residual disease inaccessible for IBT and unfit for surgery, and two patients had extraluminal residual disease and had refused surgery. All three are still alive and asymptomatic after IBT (follow-up being 140 months, 168 months and 205 months, respectively).

Fifty-one patients with CR after IBT (44 TC, 7 AC) have been closely monitored (table 3). Six (4 TC, 2 AC) had local recurrences (12%). In the two intraluminal recurrences (both TC) after 10 months and 63 months, repeat BT again resulted in CR. The remaining four...
(2 TC, 2 AC) developed extraluminal recurrences at 47 months, 104 months, 116 months and 198 months and underwent rescue surgery. It is important to point out that the extent of surgical resection would not have differed if patients had been operated at their first presentation prior to IBT despite the long interval period up to 198 months. For the final analysis, these four patients were grouped in the surgically resected cohort.

Sixty-two patients underwent surgical resection. We performed 5 pneumonectomies, 15 bilobectomies, one sleeve bilobectomy, 40 lobectomies (of which 15 were sleeve lobectomies) and one segmentectomy. For patients undergoing resections pTNM stages according to the 7th edition from the International Association for the Study of Lung Cancer of the TNM classification of malignant tumors were as follows: pT1aN0 (30 cases), pT1aN2 (2 cases), pT1bN0 (9 cases), pT2aN0 (7 cases), pT2bN0 (1 case); one case showed no residual disease. From 12 cases (all N0) the exact T-status could not be retrieved.

One patient who required emergency pneumonectomy for uncontrollable bleeding during IBT, recovered fully with specimen showing extraluminal TC N0. Remarkably, 139 months thereafter she presented with a solitary chest wall metastasis of a TC which was subsequently radically resected, and a dermal metastasis of the scalp developed (187 months postpneumonectomy) and was radically resected. She is now again free of disease for 8 months.

Metastatic disease developed in four patients (1 TC, 3 AC), two of them died. All had been treated with rescue surgery. One patient developed tracheal metastases of AC, repeatedly treated with IBT, and is alive. Overall distant metastases were seen in 1.2% (1/83) of TC and 10% (3/29) of AC.

Twelve patients have died: eight TC, four AC (table 4). Two patients died because of metastatic extraluminal AC despite surgical treatment and one patient (AC) because of postoperative complications: that is, hypoxic encephalopathy after cardiac tamponade post lobectomy. The remaining nine patients died of unrelated causes; eight had only IBT and one rescue surgery. Disease-specific mortality including surgical mortality has thus been 3/112 (2.6%). Overall disease-specific survival is 97%. Within 5 years after IBT, four patients died of disease-unrelated and treatment-unrelated causes, a 5-year survival rate of 97% (108/112). Disease-specific 5-year survival is 100% with our combined resection strategy. There were 60 patients with a follow-up of >10 years: 12 patients have died within this period; 2 because of the disease, 10 because of unrelated causes while 3 patients refusing surgery are still alive. Overall 10-year

**Figure 3** Results of current initial bronchoscopic treatment (IBT) strategy in 112 patients with bronchial carcinoids. AC, atypical carcinoid; BT, bronchoscopic treatment; TC, typical carcinoid.
survival is therefore 48/60 (80%); disease-specific 10-year survival rate is still 97% (58/60).

We compared tumour classification based on bronchoscopic versus surgical specimens in every surgically treated patient. In eight patients with extraluminal disease a shift from typical towards atypical histology classification was documented without any influence on treatment management and outcome.

DISCUSSION
The standard treatment for TC and AC is (surgical) resection. Carcinoid tumours harbour a malignant potential (AC being more malignant than TCs) and a ‘tip of the iceberg’ growth has been postulated in which the visible intraluminal part might just be a tiny fraction of tumour growing from deeper inside the tracheobronchial wall.

So, subjecting patients to non-surgical resection by less-invasive options, for example BT, was deemed unacceptable as this is not an approach through which resection margins could be checked for residual tumour; nor is sampling of lymph nodes possible. Surgical parenchyma-sparing techniques such as bronchotomy are, nevertheless, acknowledged to be acceptable from the oncological viewpoint; this is in contrast to surgery for non-small cell lung cancer.

After achieving a long-lasting CR in a small group of patients with endobronchial TC after endobronchial resection, we started a prospective study to diligently address the abovementioned issues.

Current long-term data clearly show the relatively low and slow malignant potential of centrally located bronchial carcinoids with only two deaths and one patient still alive despite extraluminal atypical metastatic disease after radical surgery.

One could suggest that delaying surgery by the described approach might induce rapid regrowth and lead to unresectability, more extensive surgery or fast distant spread; this however, is not supported by the observations in this study. There were six patients with local recurrences after BT and in the four patients in whom surgical delay might have been caused, the extent of surgical resection and long-term outcome have not been compromised. Four patients with extraluminal disease on diagnostic HRCT performed 6 weeks after BT and radically resected with N0 status still developed metastases later on. For the three patients not achieving CR after BT, but not operated upon, survival with disease has been long.

Iceberg phenomenon can be clinically observed by looking at the detailed images of current HRCT in defining tumour growth pattern, in addition to bronchoscopic examinations in the vast majority of our cases. One would argue that too many repeat HRCTs and bronchoscopies are burdensome, causing long-term emotional distress and increased costs. A more detailed analysis about cost-effectiveness of BT strategy is underway. Our unpublished data on the Hospital Anxiety and Distress Scale, showed no difference in scores for both group of patients. One may carefully state that the ‘so-called’ burden of repeat investigations and uncertainty does not seem to outweigh the 42% success rate of BT as a complementary initial approach to surgical resection. At least to our knowledge, there has not been any publication to address this issue properly.

As a result of continuous reflection on optimising our IBT strategy diagnostically and therapeutically during this longitudinal study, back referral to the pulmonologists after successful IBT is now much earlier and follow-up has been scheduled less frequently, that is, HRCT repeat scan occurs at an interval of 2 months.

Table 2: Clinical outcome of patients with bronchial carcinoid after initial bronchoscopic treatment (IBT) with or without completion surgery

<table>
<thead>
<tr>
<th>Histology</th>
<th>N</th>
<th>Treatment</th>
<th>Outcome</th>
<th>Alive</th>
<th>Dead</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Typical (83)</td>
<td></td>
<td>IBT (38)</td>
<td>CR (33)</td>
<td>120</td>
<td>168</td>
<td>(73–241)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Residue (3)</td>
<td>10 and 63</td>
<td>98 and 29</td>
<td>(140–205)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Recurrence (2)</td>
<td>47 and 104</td>
<td>57 and 111</td>
<td>(60–267)</td>
</tr>
<tr>
<td>Atypical (29)</td>
<td>37</td>
<td>Surgery (37)</td>
<td>CR (35)</td>
<td>129</td>
<td>192</td>
<td>IBT—Recurrence Surgery—July 2014</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Recurrence (2)</td>
<td>116 and 198</td>
<td>113 and 79</td>
<td>IBT—Recurrence Surgery—July 2014</td>
</tr>
</tbody>
</table>

No residual tumour detected macroscopically (videobronchoscopy, high resolution CT) and microscopically (biopsy and brush specimens).

CR, complete remission; BT, bronchoscopic treatment.

Table 3: Follow-up (FU) of patients still alive in months from initial bronchoscopic treatment (IBT) until July 2014

<table>
<thead>
<tr>
<th>Carcinoid</th>
<th>Treatment (n)</th>
<th>Outcome (n)</th>
<th>Median FU (mo)</th>
<th>Range FU (mo)</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Typical (75)</td>
<td>IBT (38)</td>
<td>CR (33)</td>
<td>120</td>
<td>(73–241)</td>
<td>IBT—Recurrence</td>
</tr>
<tr>
<td></td>
<td>Surgery (37)</td>
<td>CR (35)</td>
<td>129</td>
<td>(60–267)</td>
<td>Surgery—July 2014</td>
</tr>
<tr>
<td>Atypical (25)</td>
<td>IBT (5)</td>
<td>CR (5)</td>
<td>109</td>
<td>(83–170)</td>
<td>IBT—Recurrence</td>
</tr>
<tr>
<td></td>
<td>Surgery (20)</td>
<td>CR (18)</td>
<td>88.5</td>
<td>(60–159)</td>
<td>Surgery—July 2014</td>
</tr>
</tbody>
</table>

No residual tumour detected macroscopically (videobronchoscopy, high resolution CT) and microscopically (biopsy and brush specimens).

CR, complete remission; BT, bronchoscopic treatment.
bronchoscopic therapy suffices. By doing so, a large percentage of patients presenting with bronchial carcinoids (42%) can be treated curatively as well as safely in the most parenchyma-sparing way, without compromising outcome.

In conclusion, our long-term follow-up shows that IBT strategy in patients with bronchial carcinoids, complemented with surgical resection in case of extraluminal disease, resulted in excellent long-term outcome. Further studies of bronchial carcinoids’ molecular heterogeneity seem warranted. IBT strategy showed to be a justifiable alternative for surgical resection in 42% of the cases, and needs to be implemented in the treatment of patients presenting with bronchial carcinoids.

Contributors All four authors contributed substantially to the conception and design of the work, acquisition, analysis and interpretation of data for this study. Every author revised this article critically for important intellectual content and gave final approval of the version to be published. Furthermore every author agrees to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Competing interests None.

Ethics approval Institutional review board.

Provenance and peer review Not commissioned; internally peer reviewed.

REFERENCES