Patients with locally advanced lung cancer and very limited pulmonary function (forced expiratory volume in 1 second [FEV1] \( \leq 1 \) L) have dismal prognosis and undergo palliative treatment or best supportive care. We describe two cases of locally advanced node-positive non-small cell lung cancer (NSCLC) patients with very limited lung function treated with induction chemotherapy and moderate hypofractionated image-guided radiotherapy (Hypo-IGRT). Hypo-IGRT was delivered to a total dose of 45 Gy to the primary tumor and involved lymph nodes. Planning was based on positron emission tomography-computed tomography (PET/CT) and four-dimensional computed tomography (4D-CT). Internal target volume (ITV) was defined as the overlap of gross tumor volume delineated on 10 phases of 4D-CT. ITV to planning target volume margin was 5 mm in all directions. Both patients showed good clinical and radiological response. No relevant toxicity was documented. Hypo-IGRT is a feasible treatment option in locally advanced node-positive NSCLC patients with very limited lung function (FEV1 \( \leq 1 \) L).

Keywords: Hypofractionated radiotherapy, Image-guided radiotherapy, NSCLC, Thoracic radiotherapy

Introduction

Locally advanced node-positive lung cancer patients often present with limited lung function due to chronic pulmonary comorbidities. Because conventional thoracic irradiation and chemoradiotherapy (CRT) are associated with significant pulmonary toxicity, these high-risk patients undergo palliative treatment or best supportive care alone [1].

Stereotactic body radiotherapy (SBRT) is an established treatment option in early-stage inoperable non-small cell lung cancer (NSCLC) [2]. SBRT was also shown to be feasible and safe in patients with limited lung function. Furthermore, several small trials have demonstrated that accelerated hypofractionated thoracic irradiation can be safely delivered in locally advanced lung cancer [3,4]. Nevertheless, these small trials included patients with adequate pulmonary function exclusively. There is no data on the feasibility of multimodality treatment in locally advanced node-positive patients with reduced lung function.

Two high-risk patients with very limited lung function (forced expiratory volume in 1 second [FEV1] \( \leq 1 \) L) and locally advanced node-positive disease were treated with induction chemotherapy and Hypo-IGRT.
chemotherapy followed by hypofractionated image-guided radiotherapy (Hypo-IGRT). In the following report, we describe these cases with follow-up of 9 months.

**Case Report**

1. Patients

A 72-year-old woman (current non-smoker) with the Eastern Cooperative Oncology Group (ECOG) performance status 2 presented with stage IIIA NSCLC. She had chronic obstructive pulmonary disease (COPD) Global Initiative for Chronic Obstructive Lung Disease (GOLD) III at baseline. Tumor-related symptoms included worsening of dyspnea. The histology was an adenocarcinoma of the lung, epidermal growth factor receptor (EGFR) wild-type without anaplastic lymphoma kinase (ALK)-translocation. 18F-FDG/PET-CT (positron emission tomography with 2-deoxy-2-[fluorine-18]-fluoro-D-glucose integrated with computed tomography) in treatment position showed the primary tumor with pleural contact in the right upper lobe and positive lymph nodes in the ipsilateral hilum and mediastinum. The pulmonary function tests revealed a vital capacity (VC) of 2.02 L, FEV1 of 1.0 L (60% predicted) and an estimated DLCO (diffusion capacity for carbon monoxide) of 35% of the norm. Induction chemotherapy constituting cisplatin 80 mg/m² IV (intravenous) day1 and pemetrexed 500 mg/m² IV day1 (every 3 weeks) was initially administered, leading to stable disease. Two and a half months after the last cycle of chemotherapy, Hypo-IGRT was delivered.

A 62-year-old man (current non-smoker) with ECOG performance status 2 presented with stage IIIA NSCLC. He had COPD GOLD IV at baseline and was on supplemental oxygen. Tumor-related symptoms included pain in the parasternal region and worsening of dyspnea. The histology revealed an adenocarcinoma of the lung, EGFR wild-type without ALK-translocation. FDG/PET-CT scan in treatment position showed the primary tumor with pleural contact in the left upper lobe and negative lymph nodes in the ipsilateral mediastinum. The pulmonary function tests revealed a VC of 2.66 L, FEV1 of 0.84 L (56% predicted) and DLCO was not measurable. Induction chemotherapy was also administered analog the above-mentioned case, leading to stable disease. Two months after the last cycle of chemotherapy, Hypo-IGRT was delivered.

2. Hypo-IGRT

Hypo-IGRT was performed following chemotherapy. Patients underwent PET-CT and four-dimensional computed tomography (4D-CT) (10 sequential scans in all respiratory phases) in the treatment position (vacuum pillow and alpha-cradle). 4D-CT was matched with PET-CT for better definition of the target volume. The range of tumor motion was 1–4 mm in every dimensional space. Gross tumor volume (GTV) included primary tumor and involved lymph nodes (short-axis ≥1 cm and/or PET-positive) post-induction chemotherapy. Internal target volume (ITV) was defined through the overlap of GTVs delineated on 10 phases of 4D-CT. Clinical target volume was not generated in both cases, due to much reduced pulmonary reserve. Isotropic margin of 5 mm was added to ITV to generate the planning target volume (PTV). PTV was 255.5 and 314.28 cm³ in patient #1 and #2, respectively. Dose was calculated in 3 mm grid with Collapse-Cone algorithm (Fig. 1).

Total dose to PTV was 45 Gy in 15 fractions, 5 times a week and was prescribed per ICRU with corrections for lung inhomoogeneity. Pulmonary dose-volume parameters were calculated with total lung volume minus PTV. Mean lung dose (MLD), averages of the ipsilateral and contralateral lung were 10.6 Gy, 14.45 Gy, and 4.98 Gy in patient #1 and 9.4 Gy, 13.99 Gy, and 5.17 Gy in patient #2, respectively. In the first patient, V5 and V10 of both lungs were 52% and 32%, respectively. In the second patient, V5 and V10 were 48% and 29%. V20 (both lungs) was 19.1% and 15% in patients #1 and #2, respectively. Radiation was delivered by linear accelerator (Elekta Synergy, Stockholm, Sweden) using 15 MeV 8–9 coplanar fixed photon beams. Image guidance was performed with kilovoltage-cone-beam CT prior to each treatment. Toxicity monitoring was performed twice weekly to record any side effects.

3. Treatment tolerability

The first patient presented with grade 2 esophagitis (Common Terminology Criteria for Adverse Events v4.0) during last 4 days of treatment. Hypo-IGRT was not interrupted and the symptoms were resolved 5 days following irradiation. Lung function after treatment was at 1 month VC/FEV1/DLCO 1.87 L/0.88 L (53.6% predicted)/27.4%; at 3 months 1.73 L/0.98 L (59.8% predicted)/no data; and at 6 months 1.65 L/0.88 L (55% predicted)/27.0%, respectively. No symptoms relative to radiation pneumonitis (CT thorax 6 weeks and 3 months post-irradiation) and/or worsening of the baseline COPD-related symptoms were observed. The Patient achieved good partial response and she is progression-free after 9 months follow-up. Lung function at 9-month follow-up revealed VC/FEV1/DLCO 1.61 L/0.98 L (59.9% predicted)/22%.

The second patient presented with an acute COPD exacerbation during the first week of Hypo-IGRT. Immediate hospitalization with application of systemic antibiotics and
steroids was necessary. He developed grade 2 esophagitis at the end treatment, which was self-limiting after 3 weeks. Hypo-IGRT was interrupted for 3 days. Convalescence occurred 4 days following hospitalization. Lung function following treatment with bronchodilators, steroids, oxygen und pulmonary rehabilitation therapy was at 1 month VC/FEV1/DLCO 2.22 L/0.93 L (30.8% of predicted)/not measurable and at 3 months with 2.16 L/0.86 L (28.4% predicted)/not measurable. Pulmonary function test at 6 months was refused by the patient. No symptoms of radiation pneumonitis on CT and/or worsening of the baseline COPD-related symptoms were observed. Partial response was achieved and remained stable, but distant progression after 6 months follow-up was documented and second-line chemotherapy was initiated.

**DISCUSSION**

The present case report demonstrates that moderate Hypo-IGRT can be considered as a feasible treatment in locally advanced NSCLC patients with very limited lung function. An important requirement of our radiation treatment was an accurate transfer of SBRT technique for early-stage lung cancer to these high-risk node-positive patients. We have chosen a moderate dose prescription concept with overall duration of 3 weeks (15 fractions at 3 Gy per fraction once daily, 5 times a week) and corresponding biologically effective dose 10 Gy (BED α/β-ratio 10 Gy assumed) of 58.5 Gy according to Amini et al. [3]. Local effectiveness of this moderate accelerated hypofractionated protocol was previously extensively described [3,4]. Important prerequisites referring to the treatment planning were also maximal accepted MLD and V20 (both lungs) of 10 Gy and 20% to significantly reduce the risk of symptomatic pneumonitis [5–7]. Both patients completed Hypo-IGRT with good partial response and complete regression of the tumor-related symptoms. Although we observed a mild decline of the pulmonary function parameters (Table 1) 3 to 6 months after Hypo-IGRT, no worsening of the baseline COPD-related symptoms and radiation pneumonitis were documented.

Chiang et al. [8] reported the effectiveness and feasibility of
Hypofractionated radiotherapy in multimorbid patients with early-stage NSCLC without lymph node involvement but not for locally-advance node-positive NSCLC as described in the above-mentioned cases.

A study by He et al. [9] analyzed a subgroup of patients with limited pulmonary function, who received hypofractionated (60 Gy in 20 fractions) CRT via helical tomotherapy and showed good response without an increase in treatment-related toxicity.

Furthermore, previous studies have shown that conventional radiotherapy was associated with significant alterations in patient pulmonary function. Apart from tumor-related morbidity, residual pulmonary function is a major determinant of patient condition and survival. A previous study in locally advanced NSCLC treated with definitive CRT found only 8 months survival in patients with pronounced lung function deficits (VC <60%, FEV1 <80% predicted, DLCO <60%) vs. 16 months in patients with adequate pulmonary function [10]. An analysis from Gopal et al. [11] in 100 lung cancer patients treated with CRT revealed a 1.3% DLCO decline for each 1% of total lung volume that received >20 Gy. Borst et al. [12] also found a significant decrease in the lung function parameters starting at 3 months after definitive CRT.

In the last decade, several groups investigated poor baseline pulmonary function in patients with early-stage NSCLC treated with SBRT and demonstrated that this condition should not be used to exclude patients from treatment [13]. Moderate decline in lung function parameters were revealed 6 to 24 months after SBRT. In addition, Guckenberger et al. [14] reported an improvement of parameters in a subgroup of patients with worse pre-treatment pulmonary function. Another analysis has failed to demonstrate any dose-response relationship for the reduction of function parameters after SBRT.

In conclusion, our case report demonstrates that moderate Hypo-IGRT can be considered in locally advanced node-positive NSCLC patients with very limited lung function but careful patient selection and monitoring as well as experience with SBRT technique are important prerequisites. Prospective studies for the evaluation of safety and efficacy of this approach are urgently necessary; we are currently planning a prospective study pending approval from the ethics committee of the Ludwig-Maximilians University of Munich.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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Table 1. Treated volumes and lung function parameters pre- and 3- (patient #2) to 6-month (patient #1) post-Hypo-IGRT

| Patient no. | ITV (mL) | PTV (mL) | VC pre Hypo-IGRT (L) | FEV1 L | % | DLCO | FEV1 L | % | DLCO |
|-------------|---------|---------|---------------------|-------|---|------|-------|---|------|
| 1           | 128.7   | 255.5   | 2.02                | 1     | 60| 35   | 1.65  | 56| 2.16 |
| 2           | 167.4   | 314.28  | 2.66                | 0.84  | 56| N/M  | 2.16  | 86| 28.4 |

Hypo-IGRT, hypofractionated image-guided radiotherapy; ITV, internal target volume; PTV, planning target volume; VC, vital capacity; FEV1, forced expiratory volume in 1 second; DLCO, diffusion capacity for carbon monoxide; N/M, not measurable.
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