

Correlation between DVH parameters and lung function changes before and after radiotherapy and the occurrence of radiation induced lung injury (RILI)

Sha SHA¹, Jigang DONG¹, Maoyu WANG¹, Ziyu CHEN¹, Peng GAO¹

¹ Department of Radiotherapy, Jiaozhou Central Hospital, Qingdao, 266300, China

Correspondence to: Sha Sha, MM
Department of Radiotherapy, Jiaozhou Central Hospital, 29 Xuzhou Road, Jiaozhou City, Qingdao, 266300, China
TEL.: 15966865092; E-MAIL: shashavip325@163.com

Submitted: 2021-02-05 *Accepted:* 2021-07-15 *Published online:* 2021-07-15

Key words: lung function; DVH; radiation-induced lung injury; radiotherapy

Neuroendocrinol Lett 2021;42(5):297-304 PMID: 34506093 NEL420521A06 © 2021 Neuroendocrinology Letters • www.nel.edu

Abstract

OBJECTIVE: To explore the correlation between dose volume histogram (DVH) parameters and lung function changes before and after radiotherapy and the occurrence of radiation-induced lung injury (RILI), and to evaluate its value in predicting the risk of RILI.

METHODS: 120 patients with advanced non-small cell lung cancer diagnosed in Jiaozhou Central Hospital of Qingdao City in the past three years and received chest conformal (intensity modulated) radiation therapy were selected. Before radiotherapy, irradiation of 45-50 Gy, and 1 month after radiotherapy, the patients were tested for lung function. The evaluation of radiation lung injury was based on the RTOG acute radiation lung injury classification standard, and the observation end point was ≥ 2 grade RILI.

RESULTS: There are 34 patients with ≥ 2 grade RILI among all enrolled patients, including 23 cases of grade 2 and 11 cases of grade 3. The difference between FVC, FEV1, FEV1 / FVC, DLCO, V5, V10, V15 before radiotherapy, 45-50 Gy, and 1 month after radiotherapy were statistically significant ($P < 0.05$). The lung function, V5, V10, and V15 before radiotherapy were related factors for RILI ($P < 0.05$). And the risk of RILI was 1.855 times that of patients with higher FEV1 / FVC before radiation therapy ($OR = 1.855 (1.199-1.946)$), patients with V10 $\geq 50\%$ were 3.673 times higher than patients with V10 $< 50\%$ ($OR = 3.673 (1.548-7.582)$).

CONCLUSIONS: V10 $\geq 50\%$ and FEV1 / FVC are high-risk factors for RILI before radiotherapy, which has certain value in predicting the risk of RILI.

Abbreviations:

COPD	- chronic obstructive pulmonary disease
DVH	- dose volume histogram
DLCO	- carbon monoxide dispersion
FEV1	- forced expiratory volume in 1 second
FEV1 / FVC	- forced expiratory volume in 1 second and forced vital capacity ratio
FVC	- forced vital capacity
RILI	- radiation-induced lung injury
RTOG	- Radiation Oncology Collaborative Group

INTRODUCTION

As the number of elderly people in our country continues to increase, the incidence of elderly lung cancer increases year by year (Masraksa *et al.* 2020; Akamatsu *et al.* 2020; Pacheco & Moghanaki, 2019). Lung cancer has become a current global malignant tumor that seriously threatens human health and life safety (Haratani *et al.* 2019; Carter & Erasmus, 2019). Among them, non-small cell

lung cancer accounts for 80% of lung cancer, and the incidence and mortality are relatively high, which has aroused widespread concern among clinicians (Yazgan et al. 2019; Katayama et al. 2019; Thakur et al. 2018). Locally advanced non-small cell lung cancer refers to stage IIIa and IIIb lesions that have not been found to have distant metastases, accounting for 30% to 40% of non-small cell lung cancer, of which about 80% cannot be surgically removed (Zhu et al. 2019; Filippou et al. 2019; Fakhri et al. 2017). Simultaneous radiotherapy and chemotherapy for non-small cell lung cancer that is not suitable for surgery has become the current standard treatment mode (Chatterjee et al. 2017). The emergence of radiation-induced lung injury (RILI) not only poses a great threat to the patient's prognosis and quality of life, but also hinders the clinical application of effective radiation dose, which ultimately leads to the failure of local tumor control (Fountain et al. 2020; Bousabarah et al. 2019).

Postoperative radiotherapy is a necessary treatment, but it is easy to cause radiation pneumonia (Alberts et al. 2020). Volume parameters such as V5, V10, and V15 are commonly used indicators to predict radiation pneumonitis, but recent studies have found that the occurrence of radiation pneumonitis is also related to the lung function before radiotherapy (Uzel et al. 2019; Jassem, 2019). The incidence of chest cancer treated with radiotherapy and chemotherapy is 40% to 60% (Gainor et al. 2020; Quan et al. 2018). Due to its high incidence, insidious onset, delayed onset, and no targeted treatment, once diagnosed, it is often impossible to reverse the condition. This makes research simple and easy to predict indicators attract attention, in order to evaluate the risk of lung injury before and early treatment. At the same time, effective interventions can be implemented early to prevent or avoid the occurrence of RILI (Avancini et al. 2020). The occurrence of RILI is the result of multiple factors. This study retrospectively analyzed the occurrence of RILI in 120 patients with locally advanced non-small cell lung cancer after receiving three-dimensional conformal radiotherapy to explore related factors affecting the occurrence of RILI and provide a reference for further optimizing the 3D-CRT plan for locally advanced non-small cell lung cancer.

MATERIALS AND METHODS

Recruitment of research objects

A total of 120 patients with advanced non-small cell lung cancer who had been diagnosed in Jiaozhou Central Hospital of Qingdao City in the past three years (2017-2019) and undergoing chest conformal (modulated intensity) radiotherapy were selected. This study was approved by the Ethics Committee of Jiaozhou Central Hospital. All patients gave informed written consent to participate in the study. Before radiotherapy, irradiation of 45-50 Gy, and 1 month after the end

of radiotherapy, the patients were tested for lung function, and the ventilation and diffusion function of the patients were tested using the Japanese CHESTAC-8800 lung function tester. Including forced vital capacity (FVC), forced expiratory volume in 1 second (FEV1), forced expiratory volume in 1 second and forced vital capacity ratio (FEV1 / FVC) and carbon monoxide dispersion (DLCO). DVH parameters include V5, V10, and V15. The evaluation of radiation lung injury was based on the RTOG acute radiation lung injury classification standard, and the observation end point was ≥ 2 grade RILI.

Treatment method

The patient took the supine position, fixed the position with a vacuum air cushion, and performed a chest enhancement scan under the US GE16 slice spiral CT simulator. Scanning range: from the level of the cricothyroid membrane to the lower edge of the costal diaphragm angle 3cm, both are 5mm layer spacing. Import the scanned image into the VENUSTPS treatment planning system and perform 3D reconstruction. The outline of the target area is strictly implemented in accordance with the relevant provisions of the ICRU No. 50 and No. 62 report tumors. GTV: CT image shows the tumor area and enlarged lymph nodes (the long diameter of lymph nodes ≥ 1 cm). CTV: On the basis of GTV, the vertical direction is expanded by 10 to 15 mm, and the surrounding directions are each released 5 to 6 mm (squamous cell carcinoma 5 mm, adenocarcinoma 6 mm). The outline of CTV should be noted and appropriately modified according to specific anatomical limits (such as lung tissue, spinal cord, vertebral body, etc.).

PTV: Based on CTV, all directions are expanded by 5mm. The VENUS TPS planning system designs a radiotherapy plan and optimizes the treatment plan using a dose-volume histogram to ensure that 95% of the isodose line covers PTV. The limit for organs at risk is: the exposure dose of 1/3 volume of the heart does not exceed 50Gy, and the exposure dose of 2/3 volume does not exceed 45Gy. The maximum dose of spinal cord irradiation does not exceed 45Gy. According to the specific situation of the patient, the prescribed radiation dose is determined to be 50-70Gy, divided dose 200cGy / time, 1 time / d, 5 times / week, a total of 5-7 weeks of radiotherapy is completed. Chest-enhanced CT was reexamined from radiotherapy to DT30 ~ 40Gy. At the same time, according to the patient's situation to decide whether to chemotherapy, more than chemotherapy before radiotherapy, usually based on platinum-based chemotherapy.

Determination of lung function and DVH parameters

In this study, patients with abnormal lung function group means that the measured value / predicted value of lung function and ventilation function index is less than 70%, and lung function tests are routinely

Tab. 1. Comparison of the general characteristics of the two groups

Variables	RILI group (n=34)	Non-RILI group (n=86)	χ^2	P
Gender			0.477	0.49
Male	24	55		
Female	10	31		
Age (yrs)			0.946	0.331
<65	21	61		
≥65	13	25		
Pathological type			0.005	0.945
Squamous cell carcinoma	20	50		
Adenocarcinoma	14	36		
Clinical stage			0.563	0.453
IIIa	18	39		
IIIb	16	47		
Primary tumor location			0.657	0.418
Central type	15	45		
Peripheral type	19	41		
Has a history of smoking			0.119	0.73
Yes	21	56		
No	13	30		

Tab. 2. Comparison of DVH parameters in different treatment stages

Different treatment stages	DVH parameters	RILI group (n=34)	Non-RILI group (n=86)	t	P
Before radiation	V5 (%)	58.87±5.91	46.18±6.75	9.597	<0.001
	V10 (%)	43.14±6.17	33.57±6.58	7.306	<0.001
	V15 (%)	33.76±5.01	25.49±6.88	7.291	<0.001
Radiation (45-50 Gy exposure)	V5 (%)	57.96±5.39	44.41±5.85	11.669	<0.001
	V10 (%)	41.91±6.19	31.99±5.86	8.195	<0.001
	V15 (%)	33.84±4.61	25.96±4.68	8.349	<0.001
1 month after radiation	V5 (%)	51.01±4.09	39.87±5.01	11.516	<0.001
	V10 (%)	34.04±4.91	28.03±4.70	6.241	<0.001
	V15 (%)	26.84±4.31	22.37±4.69	4.801	<0.001

performed within 1 week before radiotherapy. When FEV1 / FVC <70%, it can be diagnosed as obstructed airway and restricted airflow in the lungs, that is, chronic obstructive pulmonary disease. In the study, the dose-volume histogram in each patient's radiation treatment plan was read to obtain the dosimetric parameters such as V5, V10, V20, and V30 required for the study.

Diagnostic criteria for radioactive damage

In the first year after the end of radiotherapy, general conditions, lung function and chest CT examinations were performed every 1 month to evaluate the efficacy and the occurrence of RILI. The classification

of radiation injury is strictly based on the classification standard of acute radiation injury developed by the American Radiation Oncology Collaborative Group (RTOG). In this study, patients with RILI who occurred ≥ 2 grades were included in the occurrence group. The specific grading standards are as follows, level 1: paroxysmal mild cough or shortness of breath during activity. Level 2: Continuous or repeated coughing, which can be relieved by treatment with narcotic antitussive drugs, or shortness of breath and difficulty in breathing after light activity (forced). Level 3: Use narcotic antitussive drugs to treat severe cough that cannot be controlled or relieved, or shortness of breath at rest, or confirmed

Tab. 3. Comparison of lung function and DVH parameters before radiotherapy between the two groups

Parameters	RILI group (n=34)	Non-RILI group (n=86)	χ^2	P
FVC			7.749	0.021
<70%	4	3		
70-80%	15	23		
>80%	15	60		
FEV1			8.532	0.014
<70%	3	2		
70-80%	17	25		
>80%	14	59		
FEV1/FVC			37.135	<0.001
<70%	2	5		
70-80%	26	16		
>80%	6	65		
DLCO			23.546	<0.001
<70%	2	11		
70-80%	24	20		
>80%	8	55		
V5			8.405	0.004
<75%	23	77		
≥75%	11	9		
V10			5.124	0.024
<50%	21	70		
≥50%	13	16		
V15			4.028	0.045
<25%	25	76		
≥25%	9	10		

Tab. 4. Comparison of lung function and DVH parameters in radiotherapy (radiation 45-50 Gy) between the two groups

Parameters	RILI group (n=34)	Non-RILI group (n=86)	χ^2	P
FVC			7.557	0.023
<70%	4	3		
70-80%	14	21		
>80%	16	62		
FEV1			7.715	0.021
<70%	3	2		
70-80%	16	24		
>80%	15	60		
FEV1/FVC			32.366	<0.001
<70%	2	5		
70-80%	24	15		
>80%	8	66		
DLCO			18.524	<0.001
<70%	2	10		
70-80%	21	18		
>80%	11	58		
V5			5.549	0.018
<75%	24	76		
≥75%	10	10		
V10			4.453	0.035
<50%	22	71		
≥50%	12	15		
V15			6.564	0.01
<25%	24	77		
≥25%	10	9		

by imaging examination. Clinically, symptomatic treatment such as intermittent oxygen inhalation and corticosteroids is needed. Level 4: It has been accompanied by severe respiratory dysfunction or pulmonary insufficiency, and clinical symptomatic treatment such as continuous oxygen inhalation or assisted ventilation is required.

Statistical analysis

SPSS 23.0 statistical software was used to process the data. The measurement data is expressed as mean ± SD, using *t* test. Count data is expressed as a rate (%), using χ^2 test. Chi-square test was used for univariate analysis, and Logistic regression model for multivariate logistic analysis. Inspection level $\alpha = 0.05$. *P* <0.05 indicates that the difference is statistically significant.

RESULTS

The general characteristics of research objects

In the 120 patients in this study, there were 34 cases with ≥2 grade RILI, including 23 cases with grade 2 and 11 cases with grade 3, with an incidence rate of 28.33%. Nine cases occurred during radiotherapy, and the remaining 25 cases occurred within 1 month after radiotherapy. There was no grade 4 RILI in the whole group. The median age was 57.9 years old (31-76 years old). There were 79 males and 41 females, 70 cases of squamous cell carcinoma and 50 cases of adenocarcinoma, 57 cases in stage IIIa and 63 cases in stage IIIb. The primary tumor location was 60 cases in each of central type and peripheral type. There were 77 cases with smoking history. There was no statistical difference in the comparison of general characteristics between the RILI-occurring group and the non-occurring RILI

Tab. 5. Comparison of lung function and DVH parameters 1 month after the radiotherapy in the two groups

Parameters	RILI group (n=34)	Non-RILI group (n=86)	χ^2	P
FVC			6.126	0.047
<70%	3	3		
70-80%	14	20		
>80%	17	63		
FEV1			7.614	0.022
<70%	2	1		
70-80%	16	23		
>80%	16	62		
FEV1/FVC			24.84	<0.001
<70%	2	3		
70-80%	20	13		
>80%	12	70		
DLCO			13.057	<0.001
<70%	1	6		
70-80%	18	17		
>80%	15	63		
V5			4.896	0.027
<75%	25	77		
≥75%	9	9		
V10			4.525	0.033
<50%	23	73		
≥50%	11	13		
V15			5.906	0.015
<25%	25	78		
≥25%	9	8		

group ($P > 0.05$). The statistical analysis results are shown in Table 1.

Comparison of lung function and DVH parameters in different treatment stages

The V5, V10 and V15 of the RILI-occurring group were higher than those of the corresponding non-occurring group, and the DVH parameters were statistically different between the two groups ($p < 0.05$). The statistical analysis results are shown in Table 2.

Tab. 6. Analysis of related influencing factors of RILI in two groups of patients

Variables	B	Wald	OR	P	95%CI
FEV1/FVC	0.032	3.794	1.855	0.037	1.199-1.946
V10≥50%	1.305	3.102	3.673	0.039	1.548-7.582

Taking the median of each DVH parameter in patients with RILI as the cut-off point, the DVH parameters in this study were grouped accordingly. The analysis results showed that the incidence of the DVH parameter RILI was significantly different among different groups ($p < 0.05$). The incidence of RILI at different stages of treatment during before radiotherapy (Table 3), radiotherapy (Table 4) and 1 month after the radiotherapy (Table 5) in the two groups of subjects was in V5≥75% group, V10≥50% group, and V15≥25% group, FVC <70% group, FEV1 <70% group, FEV1/FVC <70% group and DLCO <70% group. The differences between every groups were statistically significant ($p < 0.05$).

Analysis of related influencing factors of RILI in two groups of patients

The variables with single factor analysis $P < 0.1$ in this study were included in the multivariate logistic regression model. The analysis found that the factors closely related to the occurrence of RILI were FEV1 / FVC and V10 ≥50% before radiotherapy, and the OR (95% CI) values were 1.855 and 3.673, respectively. The results suggest that FEV1 / FVC and V10 ≥50% before radiotherapy are independent risk factors associated with RILI. The risk of RILI in patients with higher FEV1 / FVC before radiotherapy was 1.855 times that of relatively lower patients (OR = 1.855, 95% CI = 1.199-1.946, $P = 0.037$). Patients with V10 ≥50% were 3.673 times higher than patients with V10 <50% (OR = 3.673, 95% CI = 1.548-7.582, $P = 0.039$). The statistical analysis results are shown in Table 6.

DISCUSSION

At present, the incidence of lung cancer is showing a rapid upward trend in the world, and the mortality rate ranks first in malignant tumors in the world (Nardone *et al.* 2020; Yu *et al.* 2019). Lung cancer has become the first cause of death from malignant tumors in China, accounting for 22.7% of all malignant tumor deaths (Zhang *et al.* 2019; Wang & Ma, 2019). Synchronous radiotherapy and chemotherapy for non-small cell lung cancer that is not suitable for surgery has become the current standard treatment mode, and RILI is a common complication of lung cancer radiotherapy (Zheng *et al.* 2020; Yang *et al.* 2019). RILI not only poses a great threat to the patient's prognosis and quality of life, but also hinders the clinical application of effective radiation dose (Thrall *et al.* 2019; MacVittie *et al.* 2019). It is one of the dose limiting factors of chest

radiotherapy and an important factor affecting the failure of local tumor control (Izumi *et al.* 2019).

Radiotherapy is one of the main methods for the treatment of lung cancer, but due to serious complications of radiation pneumonitis, the dose of radiotherapy has decreased and the local recurrence rate has increased (Yanik *et al.* 2019). With the emergence of three-dimensional conformal radiotherapy and three-dimensional conformal intensity-modulated radiotherapy, the incidence of radiation pneumonitis has decreased compared with the previous. However, the data show that radiation pneumonitis with clinical symptoms occurred 7% -32%, severe radiation pneumonitis occurred 2.6% -18.0%, and death occurred 0-2% (S N *et al.* 2019; Hisakane *et al.* 2017; Yu *et al.* 2016). The occurrence of radiation pneumonitis reduces the patient's quality of life and even endangers the patient's life.

There are two manifestations of RILI, divided into two stages, namely acute radiation pneumonia and radiation pulmonary fibrosis (Bousabarah *et al.* 2019; Beach *et al.* 2020). Early research believed that these two stages were a gradual process, that is, radiation pulmonary fibrosis was gradually evolved and developed from radiation pneumonia (MacVittie *et al.* 2019). Typical radiation pneumonia generally occurs during radiotherapy or 1 to 3 months after radiotherapy. Congestion and edema of the lung tissue, increased exudation of alveolar fibrin, and thickening of the lung interstitial. Its clinical manifestation is almost no different from that of general pneumonia. Its particularity is that it is not caused by pathogenic microorganism infection, so it is often called radiation lung disease. Once diagnosed, it is often irreversible, and the clinical manifestations are mainly dry cough, less sputum, chest tightness, and chest pain (Zanette *et al.* 2018). Wet rales can be heard in the lung field where the lesion is occurring, and the breath sounds are rough. Severe cases will be accompanied by varying degrees of dyspnea, low fever, and normal blood leukocytes, which are not effective after antibacterial treatment (Ekanayake *et al.* 2020). The incidence of radiation-induced pulmonary fibrosis is more than 6 months after radiotherapy, and the fibrosis of the alveolar septum is accompanied by atrophy of the alveoli. Replaced and filled by fibrous connective tissue, the lung function is severely damaged and eventually leads to dyspnea and death. The percussion of the lung field where the lesion occurred showed dullness, the breathing sound was low or the fine wet rales were heard.

The results of univariate analysis in this study showed that lung function before radiotherapy was associated with RILI. RILI is mostly a latent and hidden development process. RILI with symptoms occurs in 13% to 37% of patients undergoing chest radiotherapy. The incidence of RILI in this study is 26%, which is consistent with it. At present, the research related to the clinical factors affecting RILI is more controversial.

Some researchers believe that low KPS score (Ao *et al.* 2009), smoking (Kim *et al.* 2010), chronic obstructive pulmonary disease (COPD) (Filippou *et al.* 2019), lung function status before radiotherapy (Borst *et al.* 2005), lower lung lobe tumors (Chang *et al.* 2006), and concurrent radiotherapy and chemotherapy can increase the risk of RILI.

So far, the academic community has not formed a unified opinion on whether the basic lung function before radiotherapy is closely related to the occurrence of radiation pneumonia. Some researchers believe that before radiotherapy, there is no direct correlation between basic pulmonary diseases such as COPD and radiation pneumonia (Filippou *et al.* 2019). However, some studies have pointed out that the state of lung function before radiotherapy is the main factor affecting the occurrence of RILI, even its independent risk factor (Hanania *et al.* 2019). The results of this study showed that patients with higher FEV1/FVC before radiation therapy had 1.855 times the risk of RILI than those with relatively lower patients. The results of the study suggest that FEV1/FVC is an independent risk factor affecting the occurrence of RILI before radiotherapy. Vdose (eg V5, V10, V15) refers to the volume of lung tissue that is irradiated above a certain dose (5Gy, 10Gy, 15Gy), as a percentage of the total lung volume. Graham *et al.* (1999) research confirmed that V20 is the only independent factor that affects the occurrence of RILI. Kim *et al.* (2005) reported that V20, V30, V50 and MLD are related to the occurrence of RILI. This study also found that all DVH parameters are involved in the occurrence of RILI. Among them, multivariate analysis confirmed that V15 is the only independent factor that affects the occurrence of RILI. And $V15 \geq 25\%$ may be an independent predictor of dosimetric occurrence of RILI.

In conclusion, the plasma levels of cytokines before and after radiotherapy changed in this study. Among the cytokines, only FEV1 / FVC and $V10 \geq 50\%$ before radiotherapy were found to be independent risk factors related to RILI. Each DVH parameter in the RILI-occurring group was higher than that in the corresponding non-occurring group. Among them, V15 was an independent high-risk factor affecting the occurrence of RILI, and $V15 \geq 25\%$ may be an independent dosimetric predictor of RILI. Before the end of radiotherapy, for the susceptible patients with risk factors, reasonably change or adjust the late radiotherapy plan to minimize the risk of RILI.

ACKNOWLEDGEMENTS

None.

COMPETING INTERESTS

The authors declare that they have no competing interests.

FUNDING

None.

REFERENCES

- 1 Akamatsu H, Murakami E, Oyanagi J, Shibaki R, Kaki T, Takase E, et al. (2020). Immune-Related Adverse Events by Immune Checkpoint Inhibitors Significantly Predict Durable Efficacy Even in Responders with Advanced Non-Small Cell Lung Cancer. *Oncologist*. **25**(4): e679–e683.
- 2 Alberts L, Wolff HB, Kastelijin EA, Schramel F, Coupe VMH. (2020). Patient-reported Outcomes After the Treatment of Early Stage Non-small-cell Lung Cancer With Stereotactic Body Radiotherapy Compared With Surgery. *Clin Lung Cancer*. **21**(3): e231–e232.
- 3 Ao X, Zhao L, Davis MA, Lubman DM, Lawrence TS, Kong FM. (2009). Radiation produces differential changes in cytokine profiles in radiation lung fibrosis sensitive and resistant mice. *J Hematol Oncol*. **2**: 6.
- 4 Avancini A, Sartori G, Gkoutakos A, Casali M, Trestini I, Tregnago D, et al. (2020). Physical Activity and Exercise in Lung Cancer Care: Will Promises Be Fulfilled? *Oncologist*. **25**(3): e555–e569.
- 5 Beach TA, Groves AM, Williams JP, Finkelstein JN. (2020). Modeling radiation-induced lung injury: lessons learned from whole thorax irradiation. *Int J Radiat Biol*. **96**(1): 129–44.
- 6 Borst GR, De Jaeger K, Belderbos JS, Burgers SA, Lebesque JV. (2005). Pulmonary function changes after radiotherapy in non-small-cell lung cancer patients with long-term disease-free survival. *Int J Radiat Oncol Biol Phys*. **62**(3): 639–44.
- 7 Bousabarrah K, Temming S, Hoevels M, Borggreffe J, Baus WW, Ruess D, et al. (2019). Radiomic analysis of planning computed tomograms for predicting radiation-induced lung injury and outcome in lung cancer patients treated with robotic stereotactic body radiation therapy. *Radiomics-Analyse von Planungs-Computertomogrammen zur Vorhersage von strahlen-induzierter Lungenschädigung und onkologischem Ergebnis bei Lungenkrebspatienten nach robotischer stereotaktischer Strahlentherapie*. *Strahlenther Onkol*. **195**(9): 830–42.
- 8 Carter BW, Erasmus JJ. Current Concepts in the Diagnosis and Staging of Lung Cancer. In: *Diseases of the Chest, Breast, Heart and Vessels 2019-2022: Diagnostic and Interventional Imaging*. edn. Edited by Hodler J, Kubik-Huch RA, von Schulthess GK. Cham (CH) 2019. p. 79–93.
- 9 Chang DT, Olivier KR, Morris CG, Liu C, Dempsey JF, Benda RK, et al. (2006). The impact of heterogeneity correction on dosimetric parameters that predict for radiation pneumonitis. *Int J Radiat Oncol Biol Phys*. **65**(1): 125–31.
- 10 Chatterjee K, Bhowmik R, Chattopadhyay B. (2017). Regional reporting of the incidence of Anaplastic Lymphoma Kinase mutation in 379 non-small-cell lung cancer patients from Kolkata: Using immunohistochemistry as the diagnostic modality in a significant subset. *South Asian J Cancer*. **6**(4): 169–70.
- 11 Ekanayake A, Madegedara D, Chandrasekharan V, Magana-Arachchi D. (2020). Respiratory Bacterial Microbiota and Individual Bacterial Variability in Lung Cancer and Bronchiectasis Patients. *Indian J Microbiol*. **60**(2): 196–205.
- 12 Fakhri G, Akel R, Salem Z, Tawil A, Tfayli A. (2017). Pulmonary Sarcoidosis Activation following Neoadjuvant Pembrolizumab plus Chemotherapy Combination Therapy in a Patient with Non-Small Cell Lung Cancer: A Case Report. *Case Rep Oncol*. **10**(3): 1070–5.
- 13 Filippou D, Kleontas A, Tentzeris V, Emmanouilides C, Tryfon S, Baka S, et al. (2019). Extended resections for the treatment of patients with T4 stage IIIA non-small cell lung cancer (NSCLC) (T4N0-1M0) with or without cardiopulmonary bypass: a 15-year two-center experience. *J Thorac Dis*. **11**(12): 5489–501.
- 14 Fountain MD, McLellan LA, Smith NL, Loughery BF, Rakowski JT, Tse HY, et al. (2020). Isoflavone-mediated radioprotection involves regulation of early endothelial cell death and inflammatory signaling in Radiation-Induced lung injury. *Int J Radiat Biol*. **96**(2): 245–56.
- 15 Gainor JF, Rizvi H, Jimenez Aguilar E, Skoulidis F, Yeap BY, Naidoo J, et al. (2020). Clinical activity of programmed cell death 1 (PD-1) blockade in never, light, and heavy smokers with non-small-cell lung cancer and PD-L1 expression \geq 50. *Ann Oncol*. **31**(3): 404–11.
- 16 Graham MV, Purdy JA, Emami B, Harms W, Bosch W, Lockett MA, et al. (1999). Clinical dose-volume histogram analysis for pneumonitis after 3D treatment for non-small cell lung cancer (NSCLC). *Int J Radiat Oncol Biol Phys*. **45**(2): 323–9.
- 17 Hanania AN, Mainwaring W, Ghebre YT, Hanania NA, Ludwig M. (2019). Radiation-Induced Lung Injury: Assessment and Management. *Chest*. **156**(1): 150–62.
- 18 Haratani K, Takeda M, Nakagawa K. (2019). A first attempt to establish a definition of oligometastatic non-small cell lung cancer by a European consensus group. *J Thorac Dis*. **11**(12): 5635–42.
- 19 Hisakane K, Yoh K, Nakamura N, Udagawa H, Kirita K, Umemura S, et al. (2017). Salvage chemoradiotherapy with cisplatin and vinorelbine for postoperative locoregional recurrence of non-small cell lung cancer. *Medicine*. **96**(47): e8635.
- 20 Izumi Y, Nakashima T, Masuda T, Shioya S, Fukuhara K, Yamaguchi K, et al. (2019). Suplatast tosilate reduces radiation-induced lung injury in mice through suppression of oxidative stress. *Free Radic Biol Med*. **136**: 52–9.
- 21 Jassem J. (2019). Adjuvant EGFR tyrosine kinase inhibitors in EGFR-mutant non-small cell lung cancer: still an investigational approach. *Transl Lung Cancer Res*. **8**(Suppl 4): S387–S390.
- 22 Katayama Y, Yamada T, Shimamoto T, Iwasaku M, Kaneko Y, Uchino J, et al. (2019). The role of the gut microbiome on the efficacy of immune checkpoint inhibitors in Japanese responder patients with advanced non-small cell lung cancer. *Transl Lung Cancer Res*. **8**(6): 847–53.
- 23 Kim DR, Laurence B, Jan VM, Wilfried de N, Hubert T. (2010). Association of TGFbeta1 polymorphisms involved in radiation toxicity with TGFbeta1 secretion in vitro. *Cytokine*. **50**(1): 37–41.
- 24 Kim TH, Cho KH, Pyo HR, Lee JS, Zo JI, Lee DH, et al. (2005). Dose-volumetric parameters for predicting severe radiation pneumonitis after three-dimensional conformal radiation therapy for lung cancer. *Radiology*. **235**(1): 208–15.
- 25 MacVittie TJ, Farese AM, Parker GA, Jackson W 3rd. (2019). The Time Course of Radiation-Induced Lung Injury in a Nonhuman Primate Model of Partial-body Irradiation With Minimal Bone Marrow Sparing: Clinical and Radiographic Evidence and the Effect of Neupogen Administration. *Health Phys*. **116**(3): 366–82.
- 26 Masraksa W, Tanasawet S, Hutamekalin P, Wongtawatchai T, Sukketsiri W. (2020). Luteolin attenuates migration and invasion of lung cancer cells via suppressing focal adhesion kinase and non-receptor tyrosine kinase signaling pathway. *Nutr Res Pract*. **14**(2): 127–33.
- 27 Nardone V, Tini P, Pastina P, Botta C, Reginelli A, Carbone SF, et al. (2020). Radiomics predicts survival of patients with advanced non-small cell lung cancer undergoing PD-1 blockade using Nivolumab. *Oncol Lett*. **19**(2): 1559–66.
- 28 Pacheco JM, Moghanaki D. (2019). Local consolidative therapy for oligometastatic patients with stage IV non-small cell lung cancer may improve survival, but unanswered questions remain. *Transl Lung Cancer Res*. **8**(Suppl 4): S407–S411.
- 29 Quan X, Gao H, Wang Z, Li J, Zhao W, Liang W, et al. (2018). Epidermal growth factor receptor somatic mutation analysis in 354 Chinese patients with non-small cell lung cancer. *Oncol Lett*. **15**(2): 2131–8.
- 30 S N SG, Raviraj R, Nagarajan D, Zhao W. (2019). Radiation-induced lung injury: impact on macrophage dysregulation and lipid alteration - a review. *Immunopharmacol Immunotoxicol*. **41**(3): 370–9.
- 31 Thakur C, Rapp UR, Rudel T. (2018). Cysts mark the early stage of metastatic tumor development in non-small cell lung cancer. *Oncotarget*. **9**(5): 6518–35.

- 32 Thrall KD, Mahendra S, Jackson MK, Jackson W 3rd, Farese AM, MacVittie TJ. (2019). A Comparative Dose-response Relationship Between Sexes for Mortality and Morbidity of Radiation-induced Lung Injury in the Rhesus Macaque. *Health Phys.* **116**(3): 354–65.
- 33 Uzel EK, Figen M, Uzel O. (2019). Radiotherapy in Lung Cancer: Current and Future Role. *Sisli Etfal Hastan Tip Bul.* **53**(4): 353–60.
- 34 Wang M, Ma X. (2019). Texture analysis in contrast enhanced CT: New method to predict prognosis of small cell lung cancer treated with platinum-based chemotherapy. *Ann Oncol.* **30** Suppl 1: i10.
- 35 Yang X, Wang X, Wang N, Jiang W, Li Y, Yin B. (2019). A study on the efficacy of recombinant human endostatin combined with chemotherapy intreating advanced non-small-cell lung cancer. *J BUON.* **24**(6): 2260–6.
- 36 Yanik F, Karamustafaoglu YA, Yoruk Y. (2019). Esophageal self-expandable metal stent placement for the palliation of dysphagia due to lung cancer. *Turk Gogus Kalp Damar Cerrahisi Derg.* **27**(1): 88–92.
- 37 Yazgan S, Gursoy S, Ucvet A, Yagci T, Unal M, Samancilar O, et al. (2019). Long-term results of sleeve lobectomy with continuous suture technique in non-small cell lung cancer. *Turk Gogus Kalp Damar Cerrahisi Derg.* **27**(1): 93–100.
- 38 Yu J, Yuan X, Liu Y, Zhang K, Wang J, Zhang H, et al. (2016). Delayed Administration of WP1066, an STAT3 Inhibitor, Ameliorates Radiation-Induced Lung Injury in Mice. *Lung.* **194**(1): 67–74.
- 39 Yu S, Yi M, Xu L, Qin S, Li A, Wu K. (2019). CXCL1 as an Unfavorable Prognosis Factor Negatively Regulated by DACH1 in Non-small Cell Lung Cancer. *Front Oncol.* **9**: 1515.
- 40 Zanette B, Stirrat E, Jelveh S, Hope A, Santyr G. (2018). Physiological gas exchange mapping of hyperpolarized (129) Xe using spiral-IDEAL and MOXE in a model of regional radiation-induced lung injury. *Med Phys.* **45**(2): 803–16.
- 41 Zhang C, Wang L, Li W, Huang Z, Liu W, Bao P, et al. (2019). Surgical outcomes of stage IV non-small cell lung cancer: a single-center experience. *J Thorac Dis.* **11**(12): 5463–73.
- 42 Zheng R, Shen Q, Mardekian S, Solomides C, Wang ZX, Evans NR 3rd. (2020). Molecular profiling of key driver genes improves staging accuracy in multifocal non-small cell lung cancer. *J Thorac Cardiovasc Surg.* **160**(2): e71–e79.
- 43 Zhu Y, Fu L, Jing W, Kong L, Yu J. (2019). Radiotherapy for patients with completely resected pathologic IIIA(N2) non-small-cell lung cancer: a retrospective analysis. *Cancer Manag Res.* **11**: 10901–8.