

# Impact of primary tumor SUV<sub>max</sub> on PET accuracy in mediastinal lymph node staging of non-small cell lung cancer (NSCLC)

Adam Kuźdzał<sup>1,A–F</sup>, Karolina Gambuś<sup>2,C,E,F</sup>, Konrad Moszczyński<sup>3,B,E,F</sup>, Sofia Popovchenko<sup>4,B,E,F</sup>, Monika Bryndza<sup>4,B,E,F</sup>,  
Lucyna Rudnicka<sup>5,B,E,F</sup>, Janusz Warmus<sup>6,B,E,F</sup>, Jolanta Hauer<sup>6,B,E,F</sup>, Katarzyna Żanowska<sup>6,B,E,F</sup>, Łukasz Trybalski<sup>6,B,E,F</sup>, Piotr Kocouń<sup>7,A–F</sup>

<sup>1</sup> Department of Radiology, Maria Skłodowska-Curie National Institute of Oncology, National Research Institute, Cracow, Poland

<sup>2</sup> Department of Ophthalmology, Ludwik Rydygier Hospital, Cracow, Poland

<sup>3</sup> Department of Coronary Artery Disease and Structural Heart Disease, Institute of Cardiology, Warsaw, Poland

<sup>4</sup> Students' Scientific Society, Jagiellonian University Medical College, Cracow, Poland

<sup>5</sup> Department of Pathology, John Paul II Hospital, Cracow, Poland

<sup>6</sup> Department of Thoracic Surgery, John Paul II Hospital, Cracow, Poland

<sup>7</sup> Department of Thoracic Surgery, Jagiellonian University Medical College, John Paul II Hospital, Cracow, Poland

A – research concept and design; B – collection and/or assembly of data; C – data analysis and interpretation;

D – writing the article; E – critical revision of the article; F – final approval of the article

Advances in Clinical and Experimental Medicine, ISSN 1899–5276 (print), ISSN 2451–2680 (online)

Adv Clin Exp Med. 2026;35(6):979–984

## Address for correspondence

Adam Kuźdzał

E-mail: adam.kuzdzal@krakow.nio.gov.pl

## Funding sources

None declared

## Conflict of interest

None declared

Received on January 13, 2025

Reviewed on July 3, 2025

Accepted on September 5, 2025

Published online on June 25, 2026

## Cite as

Kuźdzał A, Gambuś K, Moszczyński K, et al. Impact of primary tumor SUV<sub>max</sub> on PET accuracy in mediastinal lymph node staging of non-small cell lung cancer (NSCLC).

Adv Clin Exp Med. 2026;35(6):979–984.

doi:10.17219/acem/210368

## DOI

10.17219/acem/210368

## Copyright

Copyright by Author(s)

This is an article distributed under the terms of the Creative Commons Attribution 3.0 Unported (CC BY 3.0) (<https://creativecommons.org/licenses/by/3.0/>)

## Abstract

**Background.** Positron emission tomography (PET) using 18-fluorodeoxyglucose (<sup>18</sup>-FDG) is the primary imaging modality for mediastinal staging of lung cancer. Higher standardized uptake value (SUV<sub>max</sub>) reportedly correlates with more aggressive and more readily detectable tumors. However, high-quality scientific evidence regarding the correlation between primary tumor SUV<sub>max</sub> and PET performance in detecting mediastinal lymph node metastases remains limited.

**Objectives.** To analyze the correlation between <sup>18</sup>-FDG SUV<sub>max</sub> and the diagnostic yield of PET, as well as the impact of clinical factors such as age, sex, body mass index (BMI), primary tumor lobar location, and histological type on this relationship.

**Materials and methods.** This retrospective analysis utilized an institutional database and included a consecutive cohort of patients who underwent lung cancer surgery.

**Results.** In the overall cohort of 774 patients, PET demonstrated sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV) of 59%, 76%, 27%, and 92%, respectively. Significant differences in sensitivity were observed between the SUV<sub>max</sub> < 10 and SUV<sub>max</sub> > 15 groups (p = 0.007), as well as between the SUV<sub>max</sub> 10–15 and SUV<sub>max</sub> > 15 groups (p = 0.001). Significant differences in NPV were found between the SUV<sub>max</sub> < 10 and SUV<sub>max</sub> 10–15 groups (p = 0.011). Logistic regression analysis revealed no association between the risk of false negative (FN) PET results for detecting N2 disease and patient characteristics, primary tumor lobar location, or histological type.

**Conclusions.** Our study confirms that higher SUV<sub>max</sub> values of the primary tumor correlate with improved PET diagnostic performance in assessing mediastinal lymph node metastasis. A novel finding of this study is that clinical variables such as age, BMI, sex, tumor location, and histological type did not significantly influence the risk of FN results. The main limitations of this study are its retrospective design and single-institution cohort, which may not fully represent the diversity of patient demographics, disease characteristics, and treatment practices across different healthcare settings and populations.

**Key words:** lung neoplasms, neoplasm staging, lymph nodes, positron emission tomography, standardized uptake value

## Highlights

- Limited data are available regarding the correlation between primary tumor  $SUV_{max}$  and positron emission tomography (PET) accuracy in detecting mediastinal lymph node metastases in non-small cell lung cancer (NSCLC).
- Higher  $SUV_{max}$  of the primary tumor is strongly associated with improved PET sensitivity for detecting mediastinal lymph node metastases in patients with NSCLC.
- PET sensitivity significantly increases in NSCLC tumors with  $SUV_{max} > 15$  compared with tumors in lower  $SUV_{max}$  groups, thereby enhancing diagnostic performance.
- Clinical factors such as age, body mass index (BMI), sex, tumor location, and histology show no significant effect on false negative PET results in N2 lymph node assessment.

## Background

Lung cancer is one of the leading causes of cancer-related mortality worldwide.<sup>1,2</sup> Positron emission tomography (PET) using 18-fluorodeoxyglucose (<sup>18</sup>-FDG) is the primary imaging modality for mediastinal staging of lung cancer. In particular, the negative predictive value (NPV) of PET is high.<sup>3–7</sup> False positive PET results are common due to the glucose-dependent metabolism of leukocytes and granulation tissue. These mainly affect the specificity of PET and necessitate pathological confirmation of PET-positive lesions. However, as the primary goal of pretreatment evaluation is to exclude N2 disease, NPV remains the most important parameter.

Higher standardized uptake value ( $SUV_{max}$ ) values reportedly correlate with more aggressive and more easily detectable tumors.<sup>8,9</sup> However, this does not directly translate into the risk of mediastinal lymph node involvement. Reliable assessment of the correlation between specific clinical characteristics and the probability of N2 disease requires analysis of large patient cohorts treated according to a uniform protocol. Such reports are lacking. Therefore, high-quality scientific data regarding the correlation between primary tumor  $SUV_{max}$  and PET performance in detecting mediastinal lymph node metastases remain limited. In addition, our understanding of the influence of different clinical variables on the diagnostic yield of PET is still incomplete.

Current guidelines from the European Society of Thoracic Surgeons (ESTS) indicate a low risk of N2 disease in cases of PET-negative mediastinum, tumor diameter <3 cm, and tumor location within the outer 1/3 of the lung. These guidelines do not consider primary tumor  $SUV_{max}$  as a risk factor for N2 involvement.<sup>10</sup> From a practical perspective, it is important to determine whether similar diagnostic performance of PET can be expected across sexes, body mass index (BMI) categories, primary tumor lobar locations, and histological cancer types.

## Objectives

The primary objective of this study was to analyze the correlation between the  $SUV_{max}$  of <sup>18</sup>-FDG in primary

tumors and the effectiveness of PET in diagnosing mediastinal lymph node involvement. The secondary objective was to evaluate the impact of clinical factors such as age, sex, BMI, primary tumor lobar location, and histological type on the rate of false negative (FN) PET results.

## Materials and methods

### Study design

This study employed a retrospective cohort design.

### Clinical questions

1. Is there a correlation between the  $SUV_{max}$  of primary lung cancer and the diagnostic yield of PET in detecting mediastinal lymph node involvement?
2. What is the impact of specific clinical variables on the risk of obtaining FN results in N2 node assessment using PET?

### Patients

The study included a consecutive group of surgically treated patients with non-small cell lung cancer (NSCLC). The inclusion criteria were: 1) age 18–90 years, 2) clinical stages I–IVA, and 3) curative-intent pulmonary resection with systematic lymph node dissection. The exclusion criteria were: 1) neoadjuvant treatment, 2) a final diagnosis other than NSCLC, and 3) incomplete mediastinal lymph node dissection.

### Intervention

Positron emission tomography imaging was performed in all patients using a Discovery 690 scanner (GE Healthcare Chicago, USA). The upper limit of the allowed blood glucose concentration was 11 mmol/L. A weight-based formula for FDG dose calculation of 3–4 MBq/kg was used. The imaging protocol included computed tomography (CT) attenuation-correction imaging and lung

window reconstruction, with specific data acquisition parameters (80–210 mA, 3.75 mm section thickness, and 0.8-s gantry rotation speed). Whole-body PET scans were conducted with a 2.5 mm section thickness. Both non-attenuation-corrected (NAC) and measured attenuation-corrected (MAC) images were acquired using the Q.Clear algorithm (GE Healthcare). The interval between FDG injection and data acquisition was 45–60 min. The maximum SUV values were calculated using Pet Odyssey software (GE Healthcare), adjusted for patient weight and administered isotope dose. Positron emission tomography positivity was defined with reference to the SUV value of the aortic arch. The PET images were analyzed by the same team. A radiologist and a nuclear medicine specialist independently reviewed each PET scan with access to the clinical data.

Next, endobronchial ultrasound (EBUS) and endoscopic ultrasound (EUS) imaging of the mediastinum were performed with fine-needle biopsy of all PET-positive lymph nodes. Patients with no evidence of mediastinal spread were scheduled for curative-intent surgery. Pathological N stage, determined by examination of the surgical specimen and generally considered the gold standard, was used as the reference test. During surgery, anatomical lung resection and systematic lymph node dissection were performed according to the ESTS guidelines.<sup>11</sup> The consultant thoracic surgeon meticulously dissected lymph nodes from all stations and categorized them according to the 7<sup>th</sup> edition of the TNM classification.<sup>8</sup>

Postoperatively, tissue specimens were examined by an experienced lung pathologist with access to the clinical data. Standard light microscopy images with hematoxylin and eosin staining were used to diagnose nodal metastases (Olympus BX51; Olympus Corp., Tokyo, Japan).

## Data extraction and handling

Relevant data, including demographics (age and sex), clinical details (cancer stage and histological type), treatment specifics (surgery type and PET scan parameters), and outcomes (pathological results and PET scan findings), were extracted from the hospital database. Extracted data were anonymized and securely stored to maintain patient confidentiality. Artificial intelligence (AI) was not used for the production of images or graphical elements in the paper, nor for data collection or analysis.

## Statistical analyses

Statistica v. 13.5 PL (StatSoft Inc., Tulsa, USA) was used for statistical analysis. Logistic regression models were used to identify factors associated with FN PET results. Variables were grouped as follows: age (<60, 60–70,

and >70 years), BMI (18.5–25 kg/m<sup>2</sup>, >25–30 kg/m<sup>2</sup>, and >30 kg/m<sup>2</sup>), and SUV<sub>max</sub> (<10, 10–15, and >15). Both univariate and multivariate models were constructed. Initially, the forward stepwise selection method was applied to identify promising predictor variables efficiently. The results were then compared with those obtained using best subset selection, which examines all possible predictor combinations but is more computationally intensive. Odds ratios (ORs) with 95% confidence intervals (95% CIs) calculated using the Clopper–Pearson exact method were used to interpret important factors. Sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV) were calculated for the entire patient cohort and for subgroups based on sex, age, histological type, BMI, SUV<sub>max</sub>, and primary tumor lobar location. The 95% CIs for sensitivity and specificity were also calculated and compared across specific SUV<sub>max</sub> categories. Statistical significance was set at  $p < 0.05$ .

## Results

In total, 884 patients were included. Of these, 110 met the exclusion criteria (89 received neoadjuvant chemotherapy, 17 had a final diagnosis other than NSCLC, and in 4 patients, complete mediastinal lymph node dissection was not performed due to intraoperative cardiopulmonary instability). Thus, data from 774 patients were analyzed. The characteristics of the study groups are presented in Table 1.

For the entire cohort, the sensitivity, specificity, PPV, and NPV of PET were 59%, 76%, 27%, and 92%, respectively. Detailed values according to sex, age, and BMI are presented in Table 2.

Separate analyses were performed for subgroups according to SUV<sub>max</sub> categories: <10, 10–15, and >15. The results are summarized in Table 3.

A significant difference in sensitivity was found between the SUV<sub>max</sub> < 10 and SUV<sub>max</sub> > 15 groups ( $p = 0.007$ ) and between the SUV<sub>max</sub> 10–15 and SUV<sub>max</sub> > 15 groups ( $p = 0.000$ ). For NPV, a significant difference was found between the SUV<sub>max</sub> < 10 and SUV<sub>max</sub> 10–15 groups ( $p = 0.011$ ). Logistic regression analysis did not show an association between the risk of FN PET results in detecting N2 disease and the following factors: age ( $p = 0.769$ ), BMI ( $p = 0.430$ ), sex ( $p = 0.231$ ), primary tumor lobar location ( $p = 0.312$ – $0.757$ ), upper vs lower lobe location ( $p = 0.126$ ), and histological type ( $p = 0.623$ – $0.999$ ). Simplified categorization into squamous and nonsquamous histology also showed no significant association ( $p = 0.275$ ). Multivariate logistic regression analysis, which further investigated potential associations between FN results and clinical factors, also did not reveal significant relationships ( $p = 0.079$ – $0.469$ ).

**Table 1.** Characteristics of the study group

Variable		Value
Age [years]		30–87 (range) (65) (median)
Sex	F/M (n)	248/526
	F/M (%)	32/68
Tumor location, n (%)	RUL	209 (27)
	RML	31 (4)
	RLL	126 (16.3)
	RCE	56 (7.2)
	LUL	199 (25.7)
	LLL	115 (14.9)
	LCE	38 (4.9)
Histology, n (%)	ADC	234 (30.2)
	SCC	377 (48.7)
	ASC	63 (8.1)
	LCC	26 (3.5)
	Other	74 (9.5)
Stage, n (%)	IA	193 (25)
	IB	139 (17.9)
	IIA	49 (6.3)
	IIB	124 (16)
	IIIA	158 (20.5)
	IIIB	66 (8.5)
	IIIC	11 (1.4)
	IVA	34 (4.4)

ADC – adenocarcinoma; SCC – squamous cell carcinoma; ASC – adenosquamous carcinoma; LCC – large cell carcinoma; RCE – right central; LCE – left central; RUL – right upper lobe; RML – right middle lobe; RLL – right lower lobe; LUL – left upper lobe; LLL – left lower lobe.

## Discussion

There is a large body of evidence demonstrating the strong prognostic value of N status.<sup>12</sup> The prognostic value of primary tumor SUV<sub>max</sub> has been confirmed in several studies.<sup>13–15</sup> Our analysis of data from 774 patients revealed that the sensitivity, specificity, PPV, and NPV of PET were 59%, 76%, 27%, and 92%, respectively. These values are consistent with previous studies that also demonstrated a high NPV of PET in evaluating mediastinal lymph node metastasis.<sup>3–7</sup>

Our findings revealed significant differences in sensitivity among the primary tumor SUV<sub>max</sub> categories of <10, 10–15, and >15. The group with SUV<sub>max</sub> > 15 exhibited significantly higher sensitivity, suggesting that higher glucose metabolism in the primary tumor was associated with improved detectability of lymph node metastases using PET. This may be attributable to a higher likelihood of nodal metastasis arising from less differentiated tumor cell clones. These findings are consistent with previous research indicating that higher SUV<sub>max</sub> values correlate with more aggressive and more easily detectable tumors.<sup>8,9</sup> According to some authors, higher primary tumor SUV is associated with a higher rate of FN results regarding N2 node metastasis.<sup>5–7,16</sup> Our previous study demonstrated a correlation between primary tumor SUV<sub>max</sub> and histological grade, further supporting the present findings.<sup>17</sup>

Logistic regression analysis did not show a significant association between the risk of FN PET results and factors such as age, BMI, sex, lobar tumor location, or histological type. This suggests that the diagnostic yield of PET for detecting N2 disease is consistent across different patient

**Table 2.** Diagnostic yield of positron emission tomography (PET) in detecting N2 disease in the whole cohort

Variable	Sensitivity (95% CI)	Specificity (95% CI)	PPV (95% CI)	NPV (95% CI)
All patients	58.82% (48.64–68.48%)	75.60% (72.17–78.80%)	26.79% (22.87–31.10%)	92.36% (90.52–93.87%)
Male	59.02% (45.68–71.45%)	72.90% (68.62–76.89%)	22.22% (18.10–26.97%)	93.13% (90.90–94.85%)
Female	58.54% (42.11–73.68%)	81.64% (75.69–86.67%)	38.71% (30.04–48.18%)	90.86% (87.29–93.50%)
Age <60 years	79.31% (60.28–92.01%)	76.03% (68.27–82.70%)	39.66% (31.79–48.09%)	94.87% (90.02–97.43%)
Age 60–70 years	46.94% (32.53–61.73%)	77.78% (72.93–82.13%)	23.71% (17.83–30.80%)	90.88% (88.38–92.88%)
Age >70 years	58.33% (36.64–77.89%)	71.50% (64.58–77.75%)	20.29% (14.51–27.63%)	93.24% (89.50–95.72%)
BMI 18.5–25 kg/m <sup>2</sup>	60.87% (45.37–74.91%)	71.77% (65.73–77.29%)	28.57% (22.77–35.18%)	90.82% (87.24–93.46%)
BMI 25–30 kg/m <sup>2</sup>	59.46% (42.10–75.25%)	79.69% (74.30–84.40%)	29.33% (22.48–37.27%)	93.27% (90.33–95.37%)
BMI > 30 kg/m <sup>2</sup>	52.94% (27.81–77.02%)	72.97% (65.06–79.94%)	18.37% (11.79–27.47%)	93.10% (88.98–95.76%)

PPV – positive predictive value; NPV – negative predictive value; 95% CI – 95% confidence interval.

**Table 3.** Diagnostic yield of positron emission tomography (PET) for detecting N2 disease in subgroups according to SUV<sub>max</sub>

SUV <sub>max</sub>	Sensitivity (95% CI)	Specificity (95% CI)	PPV (95% CI)	NPV (95% CI)
SUV <sub>max</sub> < 10	57.14% (37.18–75.54%)	85.33% (80.42–89.40%)	29.63% (21.42–39.41%)	94.85% (92.29–96.59%)
SUV <sub>max</sub> 10–15	51.28% (34.78–67.58%)	74.75% (68.18–80.59%)	28.17% (21.03–36.61%)	88.82% (85.08–91.72%)
SUV <sub>max</sub> > 15	68.57% (50.71–83.15%)	64.45% (57.59–70.91%)	24.24% (19.34–29.93%)	92.52% (88.24–95.32%)

95% CI – 95% confidence interval.

populations. To the best of our knowledge, this is the only study to analyze this relationship. Our findings are important from a practical perspective, as they demonstrate that the diagnostic performance of PET is similar across sexes, age groups, BMI categories, tumor locations, and histological tumor types. Moreover, these findings suggest that greater caution should be exercised when assessing N2 disease in patients with primary tumor SUV<sub>max</sub> < 15, as sensitivity for detecting N2 disease is considerably lower in this group. The lower sensitivity of PET in these patients may indicate the need for more extensive use of invasive mediastinal lymph node diagnostics.

The strength of our study lies in its large cohort size, the largest published to date, and the use of standardized imaging and pathological protocols across all analyzed patients. Moreover, the distribution of histological cancer types, with 79% adenocarcinoma (AC) and squamous cell carcinoma (SCC), is representative of that commonly observed in thoracic surgical practice.

## Limitations of the study

The main limitation of this study is its retrospective design. In addition, the study cohort consisted of patients from a single institution, which may not fully represent the diversity of patient demographics, disease characteristics, and treatment practices observed across different healthcare settings and populations. Further prospective multicenter studies with larger patient cohorts are warranted to confirm our findings and identify potential factors affecting PET performance.

## Conclusions

Our study confirms that higher SUV<sub>max</sub> of the primary tumor correlates with improved PET diagnostic performance in assessing mediastinal lymph node metastasis. However, when the primary tumor SUV<sub>max</sub> values are below 15, N2 assessment should be approached with greater caution, as sensitivity for detecting N2 disease in this group is significantly lower. Positron emission tomography demonstrated a high NPV, making it valuable for ruling out N2 metastasis. Additionally, clinical variables such as age, BMI, sex, tumor location, and histological type did

not significantly influence the risk of FN results. Further research is needed to better understand these phenomena and optimize imaging protocols, thereby improving clinical outcomes in patients with lung cancer.

## Supplementary data

The supplementary materials are available at <https://doi.org/10.5281/zenodo.16977760>. The package contains the following files:

Supplementary Table 1. Box–Tidwell test.

Supplementary Table 2. Variance inflation factor (VIF).

Supplementary Table 3. Cook's distance.

## Data Availability Statement

The datasets supporting the findings of this study are not publicly available due to ethical and privacy restrictions related to sensitive patient information. Although the data have been anonymized, they contain health information protected under privacy laws and unrestricted public sharing could risk patient confidentiality and lead to misuse by unauthorized third parties. The data are available from corresponding author upon reasonable request.

## Consent for publication


Not applicable.


## Use of AI and AI-assisted technologies


Not applicable.


## ORCID iDs

Adam Kuźdżał  <https://orcid.org/0009-0008-9370-7967>

Karolina Gambuś  <https://orcid.org/0000-0001-9675-1623>

Konrad Moszczyński  <https://orcid.org/0009-0004-6012-6470>

Sofia Popovchenko  <https://orcid.org/0009-0000-0713-8713>

Katarzyna Żanowska  <https://orcid.org/0000-0003-4863-681X>

Piotr Kocoń  <https://orcid.org/0000-0002-4685-279X>

## References

- Chaitanya Thandra K, Barsouk A, Saginala K, Sukumar Aluru J, Barsouk A. Epidemiology of lung cancer. *Wspolczesna Onkol.* 2021; 25(1):45–52. doi:10.5114/wo.2021.103829
- Siegel RL, Miller KD, Jemal A. Cancer statistics, 2020. *CA Cancer J Clin.* 2020;70(1):7–30. doi:10.3322/caac.21590

3. Farrell MA, McAdams HP, Herndon JE, Patz EF. Non-small cell lung cancer: FDG PET for nodal staging in patients with stage I disease. *Radiology*. 2000;215(3):886–890. doi:10.1148/radiology.215.3.r00jn29886
4. Ventura E, Islam T, Gee MS, Mahmood U, Braschi M, Harisinghani MG. Detection of nodal metastatic disease in patients with non-small cell lung cancer: Comparison of positron emission tomography (PET), contrast-enhanced computed tomography (CT), and combined PET-CT. *Clin Imaging*. 2010;34(1):20–28. doi:10.1016/j.clinimag.2009.03.012
5. Bustos García De Castro A, Ferreirós Domínguez J, Delgado Bolton R, et al. PET-CT in presurgical lymph node staging in non-small cell lung cancer: The importance of false-negative and false-positive findings. *Radiologia*. 2017;59(2):147–158. doi:10.1016/j.rx.2016.12.001
6. Kaseda K, Watanabe K, Asakura K, Kazama A, Ozawa Y. Identification of false-negative and false-positive diagnoses of lymph node metastases in non-small cell lung cancer patients staged by integrated <sup>18</sup>F-fluorodeoxyglucose-positron emission tomography/computed tomography: A retrospective cohort study. *Thorac Cancer*. 2016;7(4):473–480. doi:10.1111/1759-7714.12358
7. Li S, Zheng Q, Ma Y, et al. Implications of false negative and false positive diagnosis in lymph node staging of NSCLC by means of <sup>18</sup>F-FDG PET/CT. *PLoS One*. 2013;8(10):e78552. doi:10.1371/journal.pone.0078552
8. Cengiz A, Aydın F, Sipahi M, et al. The role of F-18 FDG PET/CT in differentiating benign from malignant pulmonary masses and accompanying lymph nodes. *Tuberk Toraks*. 2018;66(2):130–135. PMID:30246656.
9. Takenaka T, Yano T, Morodomi Y, et al. Prediction of true-negative lymph node metastasis in clinical IA non-small cell lung cancer by measuring standardized uptake values on positron emission tomography. *Surg Today*. 2012;42(10):934–939. doi:10.1007/s00595-012-0277-7
10. De Leyn P, Dooms C, Kuzdzal J, et al. Revised ESTS guidelines for preoperative mediastinal lymph node staging for non-small-cell lung cancer. *Eur J Cardiothorac Surg*. 2014;45(5):787–798. doi:10.1093/ejcts/ezu028
11. Lardinois D, DeLeyn P, Van Schil P, et al. ESTS guidelines for intraoperative lymph node staging in non-small cell lung cancer. *Eur J Cardiothorac Surg*. 2006;30(5):787–792. doi:10.1016/j.ejcts.2006.08.008
12. Asamura H, Chansky K, Crowley J, et al. The International Association for the Study of Lung Cancer Lung Cancer Staging Project. *J Thorac Oncol*. 2015;10(12):1675–1684. doi:10.1097/JTO.0000000000000678
13. Imamura Y, Azuma K, Kurata S, et al. Prognostic value of SUV<sub>max</sub> measurements obtained by FDG-PET in patients with non-small cell lung cancer receiving chemotherapy. *Lung Cancer*. 2011;71(1):49–54. doi:10.1016/j.lungcan.2010.04.004
14. Li Y, Wu X, Huang Y, Bian D, Jiang L. <sup>18</sup>F-FDG PET/CT in lung adenocarcinoma and its correlation with clinicopathological features and prognosis. *Ann Nucl Med*. 2020;34(5):314–321. doi:10.1007/s12149-020-01450-1
15. Qiu X, Liang H, Zhong W, et al. Prognostic impact of maximum standardized uptake value on <sup>18</sup>F-FDG PET CT imaging of the primary lung lesion on survival in advanced non-small cell lung cancer: A retrospective study. *Thorac Cancer*. 2021;12(6):845–853. doi:10.1111/1759-7714.13863
16. Damirov F, Büsing K, Yavuz G, et al. Preoperative hilar and mediastinal lymph node staging in patients with suspected or diagnosed lung cancer: Accuracy of <sup>18</sup>F-FDG-PET/CT. A retrospective cohort study of 138 patients. *Diagnostics (Basel)*. 2023;13(3):403. doi:10.3390/diagnostics13030403
17. Kuźdzał B, Moszczyński K, Żanowska K, et al. Correlation between <sup>18</sup>F-FDG standardized uptake value and tumor grade in patients with resectable non-small cell lung cancer. *Transl Cancer Res*. 2023;12(12):3530–3537. doi:10.21037/tcr-23-798