Can we differentiate minimally invasive adenocarcinoma and non-invasive neoplasms based on high-resolution computed tomography features of pure ground glass nodules?

Xiaoye Wang¹,², Lihua Wang³, Weisheng Zhang¹*, Hong Zhao², Feng Li²

¹ Department of Radiology, First Affiliated Hospital of Dalian Medical University, Dalian, China, ² Department of Radiology, Second People’s Hospital of Zhuhai, Zhuhai, China, ³ Medical Imaging Department, Qingdao Women and Children’s Hospital, Qingdao, China

These authors contributed equally to this work.

*11111210027@fudan.edu.cn

Abstract

Objective

The purpose of our study was to assess the differentially diagnostic value of radiographic characteristics of pure ground glass nodules (GGNs) between minimally invasive adenocarcinoma and non-invasive neoplasm.

Methods

Sixty-seven pure GGNs (28 minimally invasive adenocarcinomas (MIA) and 39 pre-invasive lesions) were analyzed from June 2012 to June 2015. Pre-invasive lesions consisted of 15 atypical adenomatous hyperplasia (AAH) and 24 adenocarcinomas in situ (AIS). High-resolution computed tomography (HRCT) features and volume of MIA and pre-invasive lesions were assessed. Fisher exact test, independent sample t-test, Mann-Whitney U test and receiver operating characteristic (ROC) curve analysis were performed.

Results

Inter-observer agreement indexes for the diameter, mean HRCT attenuations and volume of pure GGNs were all high (ICC>0.75). Univariate analyses showed that lesion diameter, mean HRCT attenuation, and volume value differed significantly between two groups. Among HRCT findings, GGN shape as round or oval (F = 13.456, P = 0.002) and lesion borders as smooth or notched (F = 15.742, P = 0.001) frequently appeared in pre-invasive lesions in comparison with MIA. Type II and type III of the relationship between blood vessels and pure GGNs suggested higher possibility of malignancy than type I.
Conclusions
HRCT features of pure GGNs can help to differentiate MIA from non-invasive neoplasms.

Introduction
Pure ground-glass nodules (GGN) are defined as focal nodular areas of increased lung attenuation on high-resolution computed tomography (HRCT), in which the pulmonary vessels and bronchia structures through the nodule can be observed [1]. Most of persistent pure GGNs are diagnosed as focal interstitial fibrosis, adenomatous hyperplasia (AAH), adenocarcinoma in situ (AIS), minimally invasive adenocarcinoma (MIA), or even invasive adenocarcinoma [2, 3]. Previous studies have classified the treatment of pure GGNs into two groups: one for appropriate treatment and the other for following-up [4]. Which kind of pure GGNs should be resected or followed up? Can we predict the histologic subtypes and prognosis of pure GGNs? CT has been implemented on regular health check-up and HRCT enables these nodules to be investigated in more details, which are helpful for detecting malignant tumors at earlier stage. Most pure GGNs are indolent and often diagnosed as pre-invasive adenocarcinoma or MIA [5].

Malignancy rate of pure GGNs has been reported ranging from 18% to 48% [6, 7]. The rate of 5-year disease-free survival of low grade adenocarcinoma has been evaluated as 100% of AIS and near 100% disease-specific survival of MIA [8]. Thus, differential diagnosis of pre-invasive adenocarcinoma and MIA before surgery is important and it can help clinicians to determine appropriate managing strategies for pure-GGNs, which one should be resected or which one can be followed-up according to imaging findings. Pre-invasive lesions are significantly smaller and more frequently non-lobulated than invasive adenocarcinoma in pure GGNs [9]. The initial mean CT attenuation and tumor size of ≥7 mm of pure GGNs are risk factors for tumor growth [10]. The sphericity is seen more frequently with AAH and the presence of internal air bronchograms is more often seen in adenocarcinoma according to a report [11]. Several investigations reported that the notched signs, spiculations, bubbly lucencies, and rapid volume expansion are more common in cases of invasive adenocarcinomas [12–14]. The diagnostic value of radiographic characteristics of GGN has become a major focus. However, the result of above previous studies was notable to differentiate pre-invasive lesions (AAH and AIS) from MIA when they appear as pure GGNs at HRCT. We hypothesized that HRCT imaging and image-data processing techniques would allow us to detect the characteristics of pre-invasive adenocarcinoma and MIA, which appear as small and pure GGNs.

Therefore, we aimed to analyze HRCT features of pure GGNs pathologically diagnosed as MIA and pre-invasive adenocarcinoma for patient stratification and different treatment in clinic.

Materials and methods
Subjects
This retrospective study was approved by the institutional review board of First Affiliated Hospital of Dalian Medical University.

Sixty-seven pure GGNs of 67 Patients (13 men and 54 women; age range, 17–79 years; mean age ± standard deviation, 55.8 years ± 1.4) from June 2012 to June 2015 were selected retrospectively. All patients underwent a preoperative HRCT examination.
The lesions of size between 5mm and 30mm were included while the others of size less than 5mm or greater than 30mm were excluded. Fifteen lesions were diagnosed as AAH, 24 as AIS, and 28 as MIA. The differential diagnosis of AAH, AIS and MIA were based on novel classification criteria (15) proposed by International Association for the Study of Lung Cancer (IASLC), the American Thoracic Society (ATS), and the European Respiratory Society (ERS) in 2011. Pre-invasive lesions included AAH and AIS, which were defined as lesions showing no stromal, vascular, or pleural invasion [15]. MIA was defined as a predominantly lepidic lesion lacking of necrosis and any invasion of lymphatics, blood vessels, or pleura and measuring 30 mm or smaller [15] (Table 1).

### Image acquisition

All images were obtained using 64-detector (GE Discovery HD 750) scanners. The protocol parameters for each HRCT scanner were as follows: section width, 1.25-mm; reconstruction interval, 1.25-mm; tube voltage, 120KVp; tube current, 220 mA; beam pitch, 0.938; rotation time, 0.5s; and acquisition matrix, 512 × 512. Unenhanced spiral acquisitions were obtained with breath-hold from thoracic inlet to lung bases with images reconstructed at a slice thickness and interval of 1.25 mm. Images were reconstructed using a standard algorithm (GE bone algorithm). All images were sent to GE ADW 4.5 workstation and underwent multi-planar reconstruction (MPR) (including coronal and sagittal) with a thickness of 1.25 mm. HRCT Histogram was obtained using volume rendering (VR) method and was utilized to analyze the volume of lesions. All studies were reviewed on a PACS workstation (GE Healthcare, Milwau-kee, WI) with window level −500 to −700 HU and width 1400 HU.

### Image analysis

HRCT images were analyzed jointly by 2 radiologists with 11 and 8 years of experience in thorax imaging. Both readers were blinded to the patients’ clinical data, and a consensus was reached for each patient. HRCT images obtained in same patients were assessed separately in random order, with an interval of at least 2 weeks. Data analysis was based on source images and images of MPR. The imaging characteristics and volume of pure GGNs were recorded. The pure GGN was considered as spherical when its diameter was almost the same on axial,

### Table 1. Characteristics of pure GGNs (n = 67).

| Characteristics | Frequency |
|-----------------|-----------|
| **Sex, n (%)**  |           |
| Male            | 13 (19.4) |
| Female          | 54 (80.6) |
| **Age, mean (range), years** | 55.81 (17–19) |
| **Location, n (%)** |       |
| RUL/LUL         | 28 (41.8) / 18 (26.9) |
| RML             | 7 (10.4) |
| RLL/LLL         | 4 (0.6) / 10 (14.9) |
| **Pathological findings** |     |
| AAH             | 15       |
| AIS             | 24       |
| MIA             | 28       |

LLL, left lower lobe; LUL, left upper lobe; RLL, right lower lobe; RML, right middle lobe; RUL, right upper lobe.

https://doi.org/10.1371/journal.pone.0180502.t001
coronal, and sagittal views [11]. GGN shape was classified as nodular (round, oval or polygonal in shape) or irregular (shape different from above all) [16]. Lesion borders were classified as smooth, notched, lobulated, spiculated. Smooth margins were clear-cut; A notched was defined as V-shaped which the border deeper than 3 mm; lobulated margins were irregular; and spiculated margins were irregular with disordered profiles [17, 18]. The relationship between blood vessels and pure GGN was classified into III types (I, vessels passing by or through pure GGN; II, intact vessels passing through pure GGN with tiny branches; III, the lesion vessels were wider, distorted or tortuous compare to other vessels at the same image).

For each pure GGN, diameter was measured on axial images (the diameter was defined as the average length of long and short diameters [19]). The mean HRCT attenuation value was measured using region-of-interest (ROI) cursors, which traced the edge of the tumor using automatic tracing juts along its internal edge, on the slices containing the region of the lesion with greatest diameter. Large vessels and pulmonary arteries were excluded from ROI [20]. The volume was performed by automatically outlining the lesion perimeter on all axial images on which the pure GGN was visible using software integrated in the workstation. After the nodule segmentation, the workstation created a density histogram of the entire nodule using a volume rendering approach. Two dimensional color mapping image showed different colors in the segmented nodule, which reflected the density (Fig 1). All measurements were performed by two radiologists retrospectively. The average values of the data were used for statistical analysis. If the data differed between two radiologists by >2 mm in diameter, >50 HU in HRCT attenuation or >0.05 cm$^3$ in volume, the HRCT findings were reviewed and re-examined jointly by two radiologists.

Statistical analysis
All data regarding of continuous variables were expressed as mean ± SD. For HRCT variables, means of values were recorded, and inter-observer variability was calculated using repeated measure data analysis for the intra-class correlation coefficient (ICC). Statistical differences between pre-invasive lesions and MIA were analyzed using independent sample $t$ test for differences in diameter and HRCT attenuation. The volume data was analyzed using Mann-Whitney U test. The Fisher’s exact was used for assessment of lesion location and radiographic characteristics. For the differentiation of pre-invasive lesions from MIA, the optimal cut-off values of lesion diameter, HRCT attenuation and volume values were calculated using receiver operating characteristic (ROC) curve analysis. The optimal cut-off values were determined as the point closest to the upper left hand corner of ROC curve. Statistical analysis was performed using SPSS (version 17.0, SPSS, Chicago, III). $p$ value less than 0.05 was considered as statistically significant.

Results
HRCT findings and volume of pure GGN
Inter-observer agreement index for diameter, mean HRCT attenuations and volume of pure GGNs was high (ICC>0.75). Intra-class correlation coefficients were 0.97 (95% confidence interval [CI]: 0.94–0.98) for size, 0.77 (95% CI: 0.68–0.81) for HRCT attenuation, and 0.79 (95% CI: 0.68–0.81) for tumor volume. Univariate analyses showed that lesion diameter, mean HRCT attenuation, and volume value differed significantly between two groups ($p<0.001$, $p = 0.03$ and $p<0.001$, respectively).

The location of lesion showed no significant difference between pre-invasive group and MIA group ($F = 4.464$, $p = 0.34$). Among HRCT findings, pre-invasive lesions appeared as
round or oval (F = 13.456, P = 0.002) and frequently smooth or notched (F = 15.742, P = 0.001) in comparison to MIA (Figs 2–7).

The relationships between GGNs and vessels were analyzed based on relative position. Among of 67 GGNs, the type I, II, III relationships of pure GGNs and vessels were observed in 17, 30, 20 cases, separately (Figs 2–7). Obviously, the number of type II was the largest group with amount of 44.8%. Significant difference was shown between pre-invasive group and MIA group (F = 31.444, P < 0.010). The relationships of Type II and III of pure GGNs and the characteristics of vessels passing through pure GGN strongly suggested high likelihood of malignancy (Table 2).

Fig 1. Nodule contouring process. (A) The edge of the nodule is traced automatically. (B) The segmented nodule shows different colour in the two-dimensional colour mapping.

https://doi.org/10.1371/journal.pone.0180502.g001

Fig 2. Radiologic findings of pure GGN. A 58-year-old woman with an AAH in the upper lobe of right lung. Axial (A) and coronal (B) HRCT images show a pure GGN with a round shape and a smooth border, and a vessel through pure GGN (type I) (coarse arrow).

https://doi.org/10.1371/journal.pone.0180502.g002
ROC analysis of pure GGN

ROC analysis revealed that the area under ROC curve (AUC) for lesion diameter was 0.760, the AUC for mean HRCT attenuation 0.793, and the AUC for lesion volume 0.898. The optimal cut-off value used for lesion diameter, mean HRCT attenuation and volume to differentiate pre-invasive lesions from MIA was 8.18 mm (sensitivity, 75.0%; specificity, 71.8%; PPV, 44.8%; and NPV, 52.7%), -602 HU (sensitivity, 71.4%; specificity, 76.9%; PPV, 49.1%; and NPV, 50.6%), and 0.33 cm³ (sensitivity, 78.6%; specificity, 87.2%; PPV, 51.9%; and NPV, 48.8%), separately (Fig 8).

Discussion

Pure GGNs detected using HRCT are commonly identified abnormalities with different pathological results. Currently, MIA and pre-invasive lesions as pure GGNs garner more attention in clinical practice.
The optimal size cut-off value for discriminating MIA from pre-invasive lesions as pure GGNs was 8.18 mm according to this research. The larger pure GGN is, the more likely to be malignant. It correlates well with a report [10], which showed that pure GGNs with size greater than 8 mm was associated with higher rate of malignancy. However, the other study [9] showed that 10 mm was the optimal cut-off value for distinguishing invasive adenocarcinoma from pre-invasive pure GGNs. With longer life expectancy and for new adenocarcinoma classification, people and clinicians pay much attention to health check-up, Therefore, much more pure GGNs are early detected. It may also due to the different classification. For example, some studies included invasive pulmonary adenocarcinomas as malignant pure GGN, which were not analyzed in this assortment.

Moreover, HRCT attenuation of lesions could be a useful method for distinguishing pre-invasive adenocarcinoma from MIA. High HRCT attenuation and volume values of pure GGNs reflect more tumor cells that grow along alveolar septa [21]. A study showed that the pure GGN with a mean HRCT attenuation value of $\geq -670$ HU was an independent risk factor for malignancy.

**Fig 5.** A 55-year-old woman with an AIS in the upper lobe of left lung. Axial (A) and colour mapping (B) images show a pure GGN. The 6mm nodule shows an oval shape, a smooth border, and vessels through with tiny branches (coarse arrow).

https://doi.org/10.1371/journal.pone.0180502.g005

**Fig 6.** A 55-year-old man with a MIA in the upper lobe of left lung. Axial (A) and coronal (B) HRCT images show a 13.2mm irregular nodule with spiculated margin (arrow), and intact vessels through lesion with tiny branches (coarse arrow).

https://doi.org/10.1371/journal.pone.0180502.g006
for tumor growth [10]. The value of -602HU was an optimal cut-off value for differentiating pre-invasive adenocarcinoma from MIA in this study. Due to less or even none of invasive neoplastic cells growth, pre-invasive adenocarcinoma displayed lower attenuation values than MIA.

On the other hand, the optimal cut-off value of volume using to differentiate pre-invasive lesions from MIA was 0.33 cm$^3$ in this study. And the ROC analysis of volume implied an AUC value of 0.898. The proposed histogram analysis of HRCT volume provides higher discriminatory ability, and aids in improved classification of pre-invasive adenocarcinoma and MIA of pure GGNs. One report also showed that pure GGO also showed rapid volume expansion if it changed to mix GGO [14].

Table 2. HRCT morphologic features of lung pre-invasive lesion and MIA.

| Morphologic features   | Pre-invasive lesion (n = 39) | MIA (n = 28) | p-value$^a$ |
|------------------------|-----------------------------|--------------|-------------|
| Diameter (mm)          | 7.03 $\pm$ 1.74             | 10.16 $\pm$ 3.75 | 0.001$^b$   |
| Mean HRCT value (HU)   | -656.10 $\pm$ 50.71         | -606.00 $\pm$ 67.37 | 0.03$^b$   |
| Volume (cm$^3$)        | 0.18 $\pm$ 0.09             | 0.69 $\pm$ 0.66  | 0.001$^c$   |
| Shape                  |                             |               |             |
| Round/oval             | 20 (51.3%) / 16 (41.0%)     | 7 (25%) / 8 (28.6%) | 0.002       |
| Polygonal/irregular    | 2 (5.1%) / 1 (2.6%)         |               |             |
| Border                 |                             |               |             |
| Smooth/Notch           | 13 (33.3%) / 20 (51.3%)     | 7 (25%) / 3 (10.7%) | 0.001       |
| Lobulated/Spiculated   | 5 (12.8%) / 1 (2.6%)        | 10 (35.7%) / 3 (10.7%) |          |
| Pure GGN-vessels       |                             |               |             |
| Type I                 | 16 (41.0%)                  | 1 (3.6%)      | 0.001       |
| Type II                | 21 (53.8%)                  | 9 (32.1%)     |             |
| Type III               | 2 (5.1%)                    | 18 (64.3%)    |             |

Pure GGN-vessels, the relationship between pure GGN and the blood vessels.

$^a$ Fisher’s exact test.

$^b$ The independent sample t test.

$^c$ Mann-Whitney U test.

https://doi.org/10.1371/journal.pone.0180502.t002
Fig 8. Receiver operating characteristic (ROC) curve for HRCT variables. The area under ROC curve (AUC) for lesion diameter was 0.760 (A), the AUC for mean HRCT attenuation 0.793 (B), and the AUC for lesion volume 0.898 (C). The optimal cut-off value used for lesion diameter, mean HRCT attenuation and volume to differentiate pre-invasive lesions from MIA was 8.18 mm (sensitivity, 75.0%; specificity, 71.8%; PPV, 44.8%; and NPV, 52.7%), -602 HU (sensitivity, 71.4%; specificity, 76.9%; PPV, 49.1%; and NPV, 50.6%), and 0.33 cm³ (sensitivity, 78.6%; specificity, 87.2%; PPV, 51.9%; and NPV, 48.8%), separately.

https://doi.org/10.1371/journal.pone.0180502.g008
Additionally, there are other significant HRCT morphologic characteristics of pure GGNs. The pre-invasive lesions often indicate round or oval shape, and smooth or notched margin, while MIA shows oval, polygonal or irregular shape, together with shallow notch, lobular or speculated margin. The presence of central bubbly lucencies is another feature described in adenocarcinoma [18]. Air bronchograms and bubble-like lucencies are not significantly predictive in some investigations [13,22,23], which reported the notched signs, spiculations, and pleural tags to be more common in cases of invasive adenocarcinomas. The difference of HRCT features in pure GGNs is probably due to histopathologic dissimilarity in the amount of alveolar airspace and cellular components contained within the nodule or the thickness of alveolar walls [12].

As to the relationship between pure GGNs and supplying blood vessels, it was categorized into three types in this study. AAH was defined as a small focal proliferation of mild to moderately atypical type II pneumocytes and/or Clara cells lining alveolar walls. Therefore, type II is more common in AAH. The type III relationship exists more in MIA which resulted from an increased neoplastic cells. Interstitial fiber hyperplasia and malignant tissues may become more severe, in which the vessels present as tinier branches, wider or converging toward a lesion. Furthermore, tumor metabolism is faster than it in normal tissues and needs more blood supply [24]. It was found [25] that estrogen receptor alpha and beta expression distinguished a subset of NSCLC that had defined clinic-pathologic and genetic features. Therefore, the relationships between pure GGNs and supplying blood vessels may indicate whether or not this nodule needs surgical treatment.

There are some limitations which could not be overlooked. The study was retrospective, which may be affected by the bias of design. For example, Pure GGNs of benign lesions were excluded. Furthermore, discrete cutoffs may affect values for diagnosis. A small sample size was the other limitation of this study.

Conclusions

HRCT characteristics of pure GGNs, including diameter, volume, shape, margin of lesions, mean HRCT attenuation, together with the relationship between pure GGNs and supplying blood vessels, can help to differentiate MIA from pre-invasive lesions to some extent.

Acknowledgments

This study was supported by Dalian Municipal Science and Technology Plan Project (grant number 2015E12SF163) and Liaoning Science and Technology Project (grant number 201602245).

Author Contributions

Conceptualization: XW LW WZ HZ FL.
Data curation: XW LW WZ HZ FL.
Formal analysis: XW LW WZ HZ FL.
Funding acquisition: WZ.
Investigation: XW LW HZ FL.
Methodology: XW LW WZ HZ FL.
Project administration: WZ.
Resources: XW LW HZ FL.
Software: XW LW HZ FL.
Supervision: WZ.
Validation: XW LW WZ HZ FL.
Visualization: XW LW WZ HZ FL.
Writing – original draft: XW LW.
Writing – review & editing: WZ HZ FL.

References
1. Godoy MCB, Naidich DP. Overview and strategic management of subsolid pulmonary nodules. Journal Of Thoracic Imaging. 2012; 27(4):240–8. https://doi.org/10.1097/RTI.0b013e31825d515b PMID: 22847591
2. Lee HY, Lee KS. Ground-glass opacity nodules: histopathology, imaging evaluation, and clinical implications. Journal of thoracic imaging. 2011; 26(2):106–18. https://doi.org/10.1097/RTI.0b013e1811b9a64 PMID: 21508733
3. Cho S, Yang H, Kim K, Jheon S. Pathology and prognosis of persistent stable pure ground-glass opacity nodules after surgical resection. The Annals Of Thoracic Surgery. 2013; 96(4):1190–5. https://doi.org/10.1016/j.athoracsur.2013.05.062 PMID: 23968760
4. Haro A, Yano T, Kohno M, Yoshida T, Okamoto T, Maehara Y. Ground-glass opacity lesions on computed tomography during postoperative surveillance for primary non-small cell lung cancer. Lung Cancer (Amsterdam, Netherlands). 2012; 76(1):56–60.
5. Lee HY, Choi Y-L, Lee KS, Han J, Zo Ji, Shim YM, et al. Pure ground-glass opacity neoplastic lung nodules: histopathology, imaging, and management. AJR American journal of roentgenology. 2014; 202(3):W224–W33. https://doi.org/10.2214/AJR.13.111819 PMID: 24555618
6. Henschke CI, Yankelevitz DF, Miettinen OS, et al. CT screening for lung cancer: frequency and significance of part-solid and nonsolid nodules. AJR American Journal of Roentgenology. 2002; 178(5):1053–7. https://doi.org/10.2214/AJR.07.3101 PMID: 11959700
7. Nakata M, Sawada S, Saeki H, Takashima S, Mogami H, Teramoto N, et al. Prospective study of thoracoscopic limited resection for ground-glass opacity selected by computed tomography. Annals Of Thoracic Surgery. 2003; 75(5):1601–5; discussion 5–6. PMID: 12735586
8. Yoshizawa A, Moti N, Rieyi GJ, Sima CS, Gerald WL, Kris MG, et al. Impact of proposed IASLC/ATS/ERS classification of lung adenocarcinoma: prognostic subgroups and implications for further revision of staging based on analysis of 514 stage I cases. Mod Pathol. 2011; 24(5):653–64. https://doi.org/10.1038/modpathol.2010.232 PMID: 21252855
9. Lee SM, Park CM, Goo JM, Lee H-J, Wi JY, Kang CH. Invasive pulmonary adenocarcinomas versus preinvasive lesions appearing as ground-glass nodules: differentiation by using CT features. Radiology. 2013; 268(1):265–73. https://doi.org/10.1148/radiol.13120949 PMID: 23468575
10. Eguchi T, Kondo R, Kawakami S, Matsushita M, Yoshizawa A, Hara D, et al. Computed tomography attenuation predicts the growth of pure ground-glass nodules. Lung Cancer (Amsterdam, Netherlands). 2014; 84(3):242–7.
11. Oda S, Awai K, Liu D, Nakaura T, Yanaga Y, Nomori H, et al. Ground-glass opacities on thin-section helical CT: differentiation between bronchioalveolar carcinoma and atypical adenomatous hyperplasia. AJR American Journal of Roentgenology. 2008; 190(5):1363–8. https://doi.org/10.2214/AJR.07.3101 PMID: 18430856
12. Takashima S, Maruyama Y, Hasegawa M, Yamada T, Honda T, Kadoya M, et al. CT findings and progression of small peripheral lung neoplasms having a replacement growth pattern. AJR American journal of roentgenology. 2003; 180(3):817–26. https://doi.org/10.2214/AJR.180.3.1800817 PMID: 12591704
13. Honda T, Kondo T, Murakami S, Saito H, Oshita F, Ito H, et al. Radiographic and pathological analysis of small lung adenocarcinoma using the new IASLC classification. Clinical Radiology. 2013; 68(1):e21–6. https://doi.org/10.1016/j.crad.2012.09.002 PMID: 23146553
14. Lee SW, Leem CS, Kim TJ, Lee KW, Chung JH, Jheon S, et al. The long-term course of ground-glass opacities detected on thin-section computed tomography. Respiratory Medicine. 2013; 107(6):904–10. https://doi.org/10.1016/j.rmed.2013.02.014 PMID: 23514949

15. Travis WD, Brambilla E, Noguchi M, Nicholson AG, Geisinger KR, Yatabe Y, et al. International association for the study of lung cancer/americanc thoracic society/european respiratory society international multidisciplinary classification of lung adenocarcinoma. Journal of Thoracic Oncology: Official Publication of the International Association for the Study of Lung Cancer. 2011; 6(2):244–85.

16. Koo CW, Miller WT, Kucharczuk JC. Focal ground-glass opacities in non-small cell lung carcinoma resection patients. European Journal of Radiology. 2012; 81(1):139–45. https://doi.org/10.1016/j.ejrad.2010.07.001 PMID: 20674206

17. Jeong YJ, Yi CA, Lee KS. [Solitary pulmonary nodules: detection, characterization, and guidance for further diagnostic workup and treatment]. Radiologia. 2008; 50(3):183–95. PMID: 18471382

18. Lee H, Goo J, Lee C, Park C, Kim K, Park E-A, et al. Predictive CT findings of malignancy in ground-glass nodules on thin-section chest CT: the effects on radiologist performance. European radiology. 2009; 19(3):552–60. https://doi.org/10.1007/s00330-008-1188-2 PMID: 18925404

19. MacMahon H, Austin JHM, Gamsu G, Herold CJ, Jett JR, Naidich DP, et al. Guidelines for Management of Small Pulmonary Nodules Detected on CT Scans: A Statement from the Fleischner Society. Radiology. 2005; 237(2):395–400. https://doi.org/10.1148/radiol.2372041887 PMID: 16244247

20. Yang ZG, Sone S, Takashima S, Li F, Honda T, Maruyama Y, et al. High-resolution CT analysis of small peripheral lung adenocarcinomas revealed on screening helical CT. AJR American journal of roentgenology. 2001; 176(6):1399–407. https://doi.org/10.2214/ajr.176.6.1761399 PMID: 11373200

21. Noguchi M. Stepwise progression of pulmonary adenocarcinoma—clinical and molecular implications. Cancer Metastasis Rev. 2010; 29(1):15–21. https://doi.org/10.1007/s10555-010-9210-y PMID: 20108111

22. Farooqi AO, Cham M, Zhang L, Beasley MB, Austin JHM, Miller A, et al. Lung Cancer Associated With Cystic Airspaces. American Journal of Roentgenology. 2012; 199(4):781–6. https://doi.org/10.2214/ AJR.11.7812 PMID: 22997368

23. Ikeda K, Awai K, Mori T, Kawanaka K, Yamashita Y, Nomori H. Differential diagnosis of ground-glass opacity nodules: CT number analysis by three-dimensional computerized quantification. Chest. 2007; 132(3):984–90. https://doi.org/10.1378/chest.07-0793 PMID: 17573486

24. Gao F, Li M, Ge X, Zheng X, Ren Q, Chen Y, et al. Multi-detector spiral CT study of the relationships between pulmonary ground-glass nodules and blood vessels. Eur Radiol. 2013; 23(12):3271–7. https://doi.org/10.1007/s00330-013-2954-3 PMID: 23832317

25. Raso MG, Behrens C, Herynk MH, Liu S, Prudkin L, Ozburn NC, et al. Immunohistochemical expression of estrogen and progesterone receptors identifies a subset of NSCLCs and correlates with EGFR mutation. Clinical Cancer Research: An Official Journal Of The American Association For Cancer Research. 2009; 15(17):S359–68.