Stage IIIa cancer of the right lung ingrowing via right lower pulmonary vein to the left heart atrium

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Abstract
Computed tomography is performed in every patient before lung tumour resection. The presented case realises how important it is to perform this study with contrast. In a 75-year-old male we detected a tumour ingrowing from the right lung through the right lower pulmonary vein into the left atrium of the heart. The patient was qualified for primary sternotomy with extracorporeal circulation and resection of the intracardiac part of the tumour. In the second stage, right-sided thoracotomy was performed, and right lower lung lobectomy was done. Mixed heterogeneous lung cancer was diagnosed (squamous cell and non-small cell endocrine) in stage IIIa. The perioperative period was uncomplicated. The patient, due to renal failure, was not eligible for adjuvant chemotherapy. If the patient were qualified for lobectomy based directly on computed tomography without contrast, there would have been a high risk of perioperative death due to embolic incidents and heart failure. Effective multidisciplinary collaboration allowed us to avoid this sort of complication.

Key words: lung cancer, cardiac tumour, heart failure.

Introduction
A 75-year-old male heavy smoker came to our Lung Diseases Centre. He reported chronic non-productive cough, weakness for the preceding three weeks, loss of appetite, and weight loss of about ten kilograms in six months. The interview and the medical history documentation indicated that he was suffering from diabetes type 2, paroxysmal atrial fibrillation, hypertension, chronic renal failure grade 3, and from generalised atherosclerosis. At age 45 he had undergone resection of the bladder papilloma and cholecystectomy due to cholecystitis. We performed a chest X-ray and found a lung tumour (Fig. 1). In computed tomography (CT) it was located in segment 6 of the right lung, with diameter of 4.5 cm and lysis in the centre. In addition, the CT revealed in the enlarged lower pulmonary vein and in left atrium of the heart an hourglass-shaped structure with dimensions 5.4 × 1.6 cm. It extended to the periphery of the pulmonary vein branch and sixth segment. The radiologist suggested the presence of neoplastic spigot, thrombus, or myxoma. There were no enlarged lymph nodes in the chest CT. The results of laboratory tests of serum revealed the elevation of D-dimers (791 ng/ml) and C-reactive protein concentration (37 mg/l). In bronchoscopy we did not notice any anatomical changes. The bristled biopsy taken from RB6 was negative. We performed a transoesophageal ultrasound heart examination.
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In the left atrium light the additional longitudinal echo ranging from right pulmonary vein ostia to the mouth of mitral valve was recorded – the image corresponded to the character of tumour, thrombus, or myxoma.

**Material and methods**

After cardiothoracic consultation the patient was qualified for surgery. In the first stage sternotomy with extracorporeal circulation was performed. We incised the left heart atrium and revealed a pale pink tumour, which we removed completely from the atrium and from the final section of the right lower pulmonary vein.

In intraoperative cytology of the tumour imprints the pathologist found non-small cell lung carcinoma cells. We decided to perform a right-sided thoracotomy in the second stage with right lower lung lobectomy and mediastinal lymphadenectomy. The resected material was sent for histopathological examination. In Figure 2C the incised right pulmonary vein is shown with its wall infiltration forming the beginning section of the cardiovascular part of the tumour.

**Results**

In the final examination of the tumour the pathologist diagnosed mixed heterogeneous lung cancer (squamous cell and non-small cell endocrine) in stage IIIa. Postoperative course was without complication. The patient, due to renal failure, was not eligible for adjuvant chemotherapy. The patient is still alive 18 months after surgery.

**Discussion**

The most common secondary heart tumour is lung cancer [1]. There are very few cases of lung cancer extension via the pulmonary vein into the left atrium of the heart [2, 3]. They can be diagnosed by use of transoesophageal echocardiograms, magnetic resonance imaging, and CT scan [1, 3]. Most of those tumours show rapid growth and comprise a large mass in the lung parenchyma [2]. Resection of the intra-atrial mass seems to be a life-threatening procedure because the patient may die from cardiac inflow obstruction and sudden cardiac arrest or massive tumour embolism involving the major organs [2, 4, 5]. The possibil-
ity of systemic tumour embolisation should be considered in patients with large, central tumours and particularly those that abut the pulmonary veins [4, 5].

Conclusions

Computed tomography with contrast allows the detection of the spread of lung cancer through the vessels into the heart cavity [1]. Malignant lung tumours penetrating into the cavity of the heart can be, in some cases, radically removed. In this case, the tumour did not infiltrate the left atrium of the heart but only grew into its light through the lower pulmonary vein (IIa – T3N1M0 – stage). According to this, we determined the TNM as T3N1M0 – not T4.

Considering the rapid progression of the tumour in the presented case (the floating part of the tumour in the atrium grew 1 cm in a week, as seen by comparing ultrasound

Fig. 2. A) Intracardiac part of the tumour just before the resection and (B) just after the resection. C) Right lower lobe – the pulmonary vein is cut and the infiltration of the vein wall is shown. D) Right lower lobe with the parenchymal part of the tumour.
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Examinations of the heart) rapid multidisciplinary collaboration of the two centres enabled fast and efficient qualification procedure.

Disclosure
Authors report no conflict of interest.

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