

Metastases of lung cancer to such organs as the liver, bones or to the central nervous system appear to be a frequent complication of this disease. At the same time, metastases to the adrenal gland are found less frequently. Metastases of lung cancer to the spleen are a great rarity and they are described sporadically. Our report presents a unique case of left lung cancer with simultaneous metastases to the adrenal gland and to the spleen. All the presented lesions were diagnosed by ultrasound guided biopsy and confirmed by histopathological examination.

The patient received combined chemoradiotherapy. She was closely monitored over an 18-month observation period following treatment. No new metastases were reported.

Key words: lung cancer diagnosis, ultrasound guided fine-needle biopsy, metastasis to the adrenal gland, metastasis to the spleen.

Atypical dissemination of lung cancer to the adrenal gland and to the spleen

Marek Chorąży¹, Marta Majcher¹, Katarzyna Fedyszyn-Urbanowicz¹, Grażyna Bierzyńska-Macyszyn², Robert Kwiatkowski³

¹Department of Clinical Oncology and Internal Medicine, St. Leszczyński Hospital, Katowice, Poland

²Histopathological Department, Silesian Medical University, Katowice, Poland

³Radiotherapy Department, St. Leszczyński Hospital, Katowice, Poland

Introduction

Metastases to organs such as the liver, bones or central nervous system appear to be a frequent complication of lung cancer, whereas metastases to the suprarenal glands are found less frequently [1]. Metastases of lung cancer to the spleen are a great rarity and they are described sporadically [2].

An adrenal gland tumor detected incidentally during imaging tests is described as an incidentaloma [3–7].

Splenic lesions are most often incidentally detected on imaging tests requested for other conditions. Primary spleen tumors are extremely rare [8]. Primary cysts acquiring enormous proportions and hemangiomas are classified as benign tumors [9, 10]. Metastatic lesions and inflammatory pseudotumors may also be seen, but only very rarely and usually as casuistry [11, 12]. Splenic lesions may be observed in the course of malignant lymphoma [13, 14]. Lesions characteristic of sclerosing angiomatoid nodular transformation (SANT) have also been described [15].

In most cases, the typical characteristics of splenic tumors are established on the basis of histopathological findings, which are obtained by the surgical removal of the tumor or by post-mortem examination [8, 10, 16–18].

Metastases to the adrenal gland are also rare. This work presents a case of simultaneous dissemination of lung cancer to the adrenal gland and to the spleen.

Material and methods

A female patient (age 74) was sent from a hospital in Zawiercie for further investigations and management of a left lung tumor lesion discovered during X-ray examination. Chest surgeons had rejected her from an invasive therapy. However, bronchoscopy was performed and revealed no evidence of pathological bronchial lesions. In this situation the patient was sent to our hospital for the purpose of making the histopathological diagnosis (History No. 16735/877/09).

Computed tomography (CT) scan showed chest infiltration situated peripherally in the left lung. After establishing the distance, place and depth of the puncture by using CT (Fig. 1), the parietal tumor was visualized by ultrasound and a biopsy was performed. We performed an ultrasound-guided (free hand technique) fine-needle biopsy of the lesion using a Hitachi EUS 515 sonographic machine (Fig. 2). The procedure was performed under local anesthesia; no complications were recorded.

The ultrasound examination of the abdomen revealed a pathological mass in the spleen and in the left adrenal gland (Fig. 3). We also performed in local anesthesia an ultrasound-guided (free hand technique) fine-needle biopsy of these lesions.

Results and discussion

In our case small cell lung cancer was detected in the percutaneous biopsy of the left lung. The same type of cancer as in the left lung was observed in both the adrenal gland and in the spleen (metastases of small cell cancer).

Imaging methods available to us showed no evidence of cancer metastases in other organs.

In the existing literature, we found only a few cases of lung cancer metastases to the spleen [2, 17, 18]. There are also some descriptions of metastases isolated in spleen from other organs [16]. Simultaneous metastases of lung cancer to the adrenal gland and the spleen have never been described.

The case presented above shows that the metastatic lesion can sometimes be an accessible place to collect tissue for diagnosing the cancer pattern of the primary cancer site. The case is exceptional because the spleen is an organ where lung cancer metastases are not frequently found, while metastases to the adrenal gland alone are common. More often, metastases are observed in the liver. The case is also unique because the adrenal gland and the spleen are organs where finding concurrent metastases of lung cancer is very rare.

The patient received combined chemoradiotherapy. She was closely monitored over an 18-month observation period following treatment. No new metastases were reported.

The authors declare no conflict of interest.

References

1. Bilimoria KY, Shen WT, Elaraj D, Bentrem DJ, Winchester DJ, Kebebew E, Sturgeon C. Adrenocortical carcinoma in the United States: treatment utilization and prognostic factors. *Cancer* 2008; 113: 3130-6.
2. Van Hul I, Cools P, Rutsaert R. Solitary splenic metastasis of an adenocarcinoma of the lung 2 years postoperatively. *Acta Chir Belg* 2008; 108: 462-3.
3. Al-Hawary MM, Francis IR, Korobkin M. Non-invasive evaluation of the incidentally detected indeterminate adrenal mass. *Best Pract Res Clin Endocrinol Metab* 2005; 19: 277-92.
4. Hennings J, Hellman P, Ahlström H, Sundin A. Computed tomography, magnetic resonance imaging, and 11C-metomidate positron emission tomography for evaluation of adrenal incidentalomas. *Eur J Radiol* 2009; 69: 314-23.
5. Johnson PT, Horton KM, Fishman EK. Adrenal mass imaging with multidetector CT: pathologic conditions, pearls, and pitfalls. *Radiographics* 2009; 29: 1333-51.
6. Grumbach MM, Biller BM, Braunstein GD, et al. Management of the clinical inapparent adrenal mass ("incidentaloma"). *Ann Intern Med* 2003; 138: 424-9.
7. Terzolo M, Bovio S, Pia A, Reimondo G, Angeli A. Management of adrenal incidentaloma. *Best Pract Res Clin Endocrinol Metab* 2009; 23: 233-43.
8. Kochar K, Vijayasekar C, Pandey U, Bhogal R, Brown L, Mathew G. Primary carcinosarcoma of spleen: case report of a rare tumor and review of the literature. *Int J Surg Pathol* 2009; 17: 72-7.
9. Lee H, Maeda K. Hamartoma of the spleen. *Arch Pathol Lab Med* 2009; 133: 147-51.
10. Orawczyk T, Ćwik P, Ziaja D, Kazibudzki M. Familia lymphangioma – a rare form of splenic cysts. *Chir Pol* 2002; 4: 187-91.
11. Bhatt S, Simon R, Dogra VS. Radiologic-pathologic conferences of the University of Rochester School of Medicine: inflammatory pseudotumors of the spleen. *AJR Am J Roentgenol* 2008; 191: 1477-9.

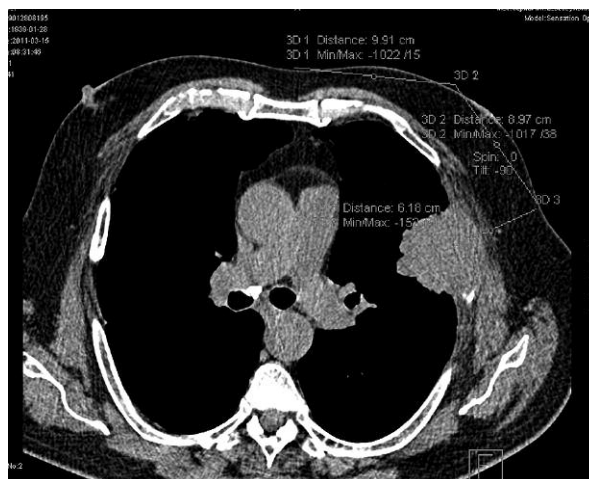


Fig. 1. Computed tomography imaging scan – establishing the distance, place and depth of the puncture

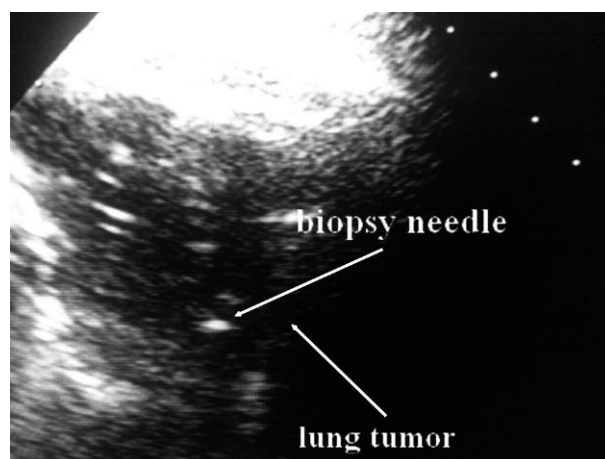


Fig. 2. Ultrasound imaging scan – wall located left lung tumor – biopsy needle



Fig. 3. Ultrasound imaging scan – small cell cancer metastasis to left adrenal gland and spleen

12. Tee M, Vos P, Zetler P, Wiseman SM. Incidental littoral cell angioma of the spleen. *World J Surg Oncol* 2008; 6: 87-92.
13. Gupta R, Naseem S, Sukumaran S, Kashyap R, Kaur S, Paul L. Splenic lymphoma with villous lymphocytes. *Indian J Pathol Microbiol* 2008; 51: 113-5.
14. Takata F, Kaida H, Ishibashi M, et al. Primary splenic lymphoma detected by F-18 FDG PET. *Clin Nucl Med* 2008; 33: 204-7.
15. Koreishi AF, Saenz AJ, Fleming SE, Teruya-Feldstein J. Sclerosing angiomatoid nodular transformation (SANT) of the spleen: a report of 3 cases. *Int J Surg Pathol* 2010; 18: S136-41.
16. Showalter SL, Hager E, Yeo CJ. Metastatic disease to the pancreas and spleen. *Semin Oncol* 2008; 35: 160-71.
17. Kinoshita A, Nakano M, Fukuda M, et al. Splenic metastasis from lung cancer. *Neth J Med* 1995; 47: 219-23.
18. Dias AR, Pinto RA, Ravanini JN, Lupinacci RM, Cecconello I, Ribeiro U Jr. Isolated splenic metastasis from lung squamous cell carcinoma. *World J Surg Oncol* 2012; 10: 24.

Address for correspondence**Marek Chorąży** MD, PhD

Department of Clinical Oncology and Internal Medicine

S. Leszczyński Hospital

Raciborska 27

40-074 Katowice, Poland

tel. +48 601 51 38 87

fax +48 32 251 45 33

e-mail: marekchorazy@wp.pl

Submitted: 13.10.2011**Accepted:** 12.09.2012