The most frequent locations are the vessels of the head and neck, fingers, trunk and cutaneous veins. Intra-abdominal lesions are rare, and even rarer are those in the liver.

Epidemiologically, a slightly higher incidence has been observed in women than in men, with a ratio of 1.2: 1, with no predilection for age. Cases have been reported in patients ranging from 7 months to 81 years of age.<sup>1</sup>

Intravascular papillary endothelial hyperplasia is an uncommon disease that can simulate other diseases. It is therefore necessary to carry out a detailed differential diagnosis, and the definitive diagnosis is often reached by the pathology study.

Treatment of IPEH depends on its location and, considering that it is a benign disease, the symptoms that it causes. In general, surgical resection is the treatment of choice for IPEH as it provides a very good prognosis and very low recurrence rate (mainly in the mixed and extravascular forms).<sup>10</sup> Surgery is considered curative in the pure forms when the margins of the surgical specimen are free, with low reported rates of recurrence in the mixed and extravascular forms.

### Funding

The authors have received no public or private funding for the completion of this article.

### REFERENCES

- Hashimoto H, Daimaru Y, Enjoji M. Intravascular papillary endothelial hyperplasia: a clinicopathologic study of 91 cases. Am J Dermatopathol. 1938;5:539–46.
- 2. Clearkin KP, Enzinger FM. Intravascular papillary endothelial hyperplasia. Arch Pathol Lab Med. 1976;100:441–4.

- Salyer WR, Salyer DC. Intravascular angiomatosis: development and distinction from angiosarcoma. Cancer. 1975;36:995–1001.
- 4. Masson P. Hemangioendotheliome vegetant intravasculaire. Bull Soc Anat Paris. 1923;93:517–23.
- Henschen P. Lendovasculite proliferante thrombopoietique dans la lesion vasculaire locale. Ann Anat Pathol. 1932;9:113– 21.
- Albrecht S, Kahn HJ. Immunohistochemestry of intravasclar papillary endothelial hyperplasia. J Cutan Pathol. 1990;17:16–21.
- Hong SG, Cho HM, Chin HM, Park IY, Yoo JY, Hwang SS, et al. Intravascular papillary endothelial hyperplasia (Masson's hemangioma) of the liver: a new hepatic lesion. J Korean Med Sci. 2004;19:305–8.
- Kuo T, Sayers CP, Rosai J. Masson's vegetant intravascular hemangioendothelioma: a lesion often mistaken for angiosarcoma. Study of seventeen cases located in the skin and soft tissues. Cancer. 1976;38:1227–36.
- 9. Weiss SW, Goldblum JR. Benign tumors and tumor-like lesions of blood vessels. In: Enzingers & Weiss's soft tissue tumors5th ed. New York: Mosby; 2008: 668–71.
- Meadows MC, Sun X, Dardik M, Tarantino DR, Chamberlain RS. Intraabdominal intravacular papillary endothelial hyperplasia (Masson's tumor): a rare and novel cause of gastrointestinal bleeding. Case Rep Gastroenterol. 2010;4:124–32.

Irene Ramallo Solís<sup>•</sup>, José Tinoco González, Ana Senent Boza, Carmen Bernal Bellido, Miguel Ángel Gómez Bravo

Unidad de Cirugía Hepatobiliopancreática y Trasplante Hepático, Hospital Universitario Virgen del Rocío, Sevilla, Spain

\*Corresponding author.

E-mail address: Irene.ramallo@gmail.com (I. Ramallo Solís).

### 2173-5077/

© 2016 AEC. Published by Elsevier España, S.L.U. All rights reserved.

# Spontaneous Pulmonary Hernia: Presentation of Four Cases and Literature Review



## Hernia pulmonar espontánea: presentación de 4 casos y revisión de la literatura

Pulmonary hernias are characterized by a protrusion of the lung through a defect of the chest wall. Herniations of the lung can be either acquired or spontaneous. Predisposing factors have been described, such as obesity, chronic obstructive pulmonary disease (COPD) or the chronic use of corticosteroids. We present the cases of 4 patients with spontaneous lung hernias treated at our hospital in 2015. The first case is that of a 60-year-old patient who was overweight and had a history of COPD. In February, he reported the appearance of a hematoma on the right thoracic wall, and a CT scan diagnosed a right pulmonary herniation between the 8th and 9th ribs. By thoracotomy, primary closure was performed with non-absorbable sutures. In April, he was diagnosed with pleural effusion compatible with empyema

<sup>\*</sup> Please cite this article as: Laguna Sanjuanelo S, López Sanz I, Zabaleta Jimenez J, Aguinagalde Valiente B, Izquierdo Elena JM. Hernia pulmonar espontánea: presentación de 4 casos y revisión de la literatura. Cir Esp. 2017;95:237–239.

<sup>\*\*</sup> The data published in this article have been previously presented as an oral communication at the Spanish National Thoracic Surgery Conference, (SECT), May 5, 2016.

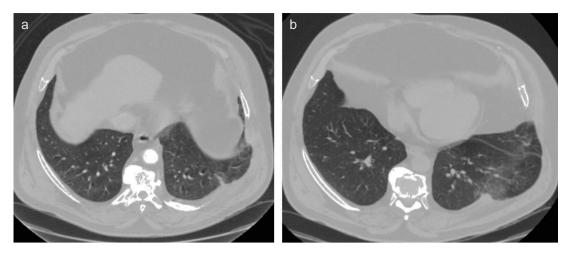


Fig. 1 – (a) Initial thoracic CT scan demonstrating the protrusion of the lung parenchyma between the left 9th and 10th ribs; (b) follow-up CT, with the image of the lung hernia recurrence between the left 8th and 9th ribs.

due to S. *aureus*. A chest drain was inserted, and the treatment was complemented with the intrapleural instillation of urokinase for 4 days, leading to resolution of the condition. After outpatient follow-up, the patient was discharged in May 2015 after showing no signs of relapse.

The second case is a 56-year-old patient with a history of COPD and grade III obesity who consulted in February for left chest pain accompanied by chest wall hematoma. CT scan revealed a left pulmonary hernia between the 9th and 10th ribs (Fig. 1a). Primary closure with non-absorbable sutures was performed by means of thoracotomy. In May, the patient again consulted for left rib pain; CT scan demonstrated an image compatible with left pulmonary herniation between the 8th and 9th ribs (Fig. 1b). The defect was closed with placement of polytetrafluoroethylene mesh. In February 2016, the patient presented with a respiratory infection and recurrence of the pulmonary hernia between the 8th and 9th ribs. The patient was treated conservatively with the use of a rib binder and referred to the endocrinology department for a weight loss regimen. The patient's progress is being followed in the outpatient clinic.

The third case is a 68-year-old overweight patient, who consulted in February for intense rib pain accompanied by the appearance of a hematoma. Thoracic CT scan detected a hematoma that was actively bleeding from the left 9th to 10th ribs, which ceased spontaneously. After 10 days, an induration was observed in the area of the hematoma, and the presence of a pulmonary herniation was confirmed. In the month of March, primary closure of the defect was performed with nonabsorbable sutures by means of thoracotomy. The last followup visit was in the month of April 2016, at which time no signs of relapse had been observed.

The fourth case is that of a 60-year-old patient who was overweight and had a history of asthma. He presented with right chest pain in the month of August, and CT scan revealed a lung hernia between the 8th and 9th ribs. Scheduled surgery was performed in the month of November using thoracotomy with primary closure of the defect with non-absorbable sutures. The last follow-up visit was in the month of March 2016, at which time no signs of recurrence were observed.

Pulmonary herniations are a rare entity and were described for the first time in 1499 by Roland. In 1845, Morel-Lavallée provided a detailed description of lung hernias, classifying them according to their etiology and anatomical location.<sup>1,2</sup>

Lung herniations are defined as protrusions of the lung parenchyma through the chest wall. They can be classified as spontaneous, traumatic and postoperative. Spontaneous lung hernias account for 30% of cases, and most do not present complications.<sup>1,3</sup>

Lung hernias are usually asymptomatic and present as an inconsistent mass that protrudes through the intercostal spaces with the elevated pressure maneuvers of the thoracic cavity.<sup>1,2</sup> Thoracic CT is a very sensitive test for the diagnosis of lung hernias and is able to assess the hernial orifice and sac as well as the relationship with the pectoral and intercostal muscles.<sup>2</sup>

The purpose of surgical treatment is to close the defect of the intercostal muscles by means of primary closure with intercostal sutures or with the placement of synthetic mesh material.<sup>2</sup> Currently, there are no studies indicating which type of surgical repair is the best for the treatment of lung hernias.<sup>3</sup>

In summary, we have presented 4 cases of spontaneous pulmonary hernias from the year 2015. In all patients, primary closure of the defect was conducted with non-absorbable sutures, taking into account the size of the defect of the intercostal space and the possibility for rib approximation with sutures. The only case of hernia recurrence was observed in the patient with type III obesity, and it occurred despite the placement of a synthetic mesh. In our experience, obesity is a factor that negatively influences patient progress after the surgical repair of a pulmonary hernia. Primary closure with non-absorbable stitches, without mesh placement, may be adequate surgical treatment for simple spontaneous lung herniations.

### Authorships

Stephany Laguna Sanjuanelo: main writer of the case review and manuscript structure.

Iker López Sanz: main reviewer of the manuscript.

Jon Zabaleta Jiménez and Borja Aguinagalde Valiente: surgeons of the operations conducted, participants in the review of the manuscript. José Miguel Izquierdo Elena: critical review and approval of the final version.

### REFERENCES

- Wani AS, Kalamkar P, Alhassan S, Farrell MJ. Spontaneous intercostal lung herniation complicated by rib fractures: a therapeutic dilemma. Oxf Med Case Reports. 2015;2015:378–81.
- 2. Detorakis EE, Androulidakis E. Intercostal lung herniation. The role of imaging. J Radiol Case Rep. 2014;8:16–24.
- 3. Akkas Y, Peri NG, Kocer B, Kaplan T. Repair of lung herniation with titanium prosthetic ribs and Prolene mesh. Asian Cardiovasc Thorac Ann. 2016;24:280–2.

Stephany Laguna Sanjuanelo<sup>\*</sup>, Iker López Sanz, Jon Zabaleta Jimenez, Borja Aguinagalde Valiente, Jose Miguel Izquierdo Elena

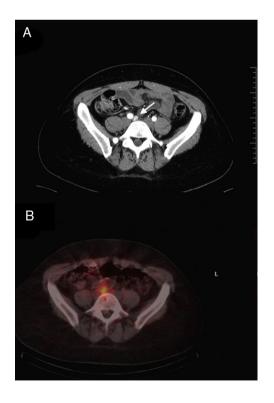
Servicio de Cirugía Torácica, Hospital Universitario Donostia Donostia-San Sebastián, Gipuzkoa, Spain

Corresponding author.
E-mail address: lagunastephy@gmail.com
(S. Laguna Sanjuanelo).

2173-5077/ © 2016 AEC. Published by Elsevier España, S.L.U. All rights reserved.

### 

Resección laparoscópica de paraganglioma en el órgano de Zuckerkandl guiado por sonda gamma con <sup>123</sup>I-metayodobencilguanidina



Pheochromocytomas and paragangliomas (PGL) are rare neuroendocrine tumors. 53% of paragangliomas are located in the organ of Zuckerkandl, situated in the distal abdominal aorta, between the inferior mesenteric artery and the aortic bifurcation.<sup>1</sup> Most sympathetic paragangliomas and pheochromocytomas are catecholamine-secreting tumors, so determining plasma and/urine levels is the initial study used in these patients.<sup>2</sup> After confirming the diagnosis, the initial imaging study should be computed tomography (CT), which can determine the tumor location in most cases. In cases of suspected paragangliomas that are not detected on CT, nuclear medicine studies can be used. Metaiodobenzylguanidine (MIBG) is an analog of guanidine, which is similar to noradrenalin. When used in clinical practice and radio-labeled with <sup>123</sup>I, it accumulates in the adrenergic cells present in pheochromocytomas and paragangliomas. Used in conjunction with CT, this technique provides exact tumor localization and is an essential tool in difficult-to-diagnose tumors.3 Treatment for paragangliomas of the organ of Zuckerkandl

Fig. 1 – Abdominal computed tomography identifying tumor lesion in the area of L5 (white arrow), medial to the bifurcation of the right iliac artery (A); SPECT/CT showing uptake of <sup>123</sup>I-MIBG, confirming the diagnosis of paraganglioma (B).

<sup>\*</sup> Please cite this article as: Navarrete A, Almenara R, Momblán D, Lacy A. Resección laparoscópica de paraganglioma en el órgano de Zuckerkandl guiado por sonda gamma con <sup>123</sup>I-metayodobencilguanidina. Cir Esp. 2017;95:239–241.

CrossMark