Video-assisted thoracoscopic resection of a Hattori’s cyst in the posterior mediastinum: a case report

Videotorakoskopska resekcija Hattorijeve ciste v posteriornem mediastinumu: prikaz primera

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Abstract
First reported by Hattori in 2004, the mediastinal cyst with Mullerian differentiation is a rather new finding. We present a case of a 51-year-old woman with a cystic formation in the left paravertebral space at the level of 4th and 5th thoracic vertebrae, which was discovered on an MRI scan. We have performed a video-assisted thoracoscopic (VATS) resection of the cyst. Histological and immunohistochemical staining showed a ciliated cyst with Mullerian differentiation (i.e. Hattori’s cyst). To our knowledge, only 31 cases have been described in the literature so far. Data suggests their prevalence may be higher since they are often found incidentally and are typically asymptomatic.

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1 Introduction

A case of a ciliated cyst with Mullerian differentiation in the posterior mediastinum was first described by Hattori in 2004. He suggested Mullerian origin due to female preponderance of ciliated cysts as well as their morphological resemblance to fallopian tubes (i.e. ciliated epithelium with underlying smooth muscle layer) (1). As similar reports of mediastinal cysts with Mullerian differentiation followed, it became increasingly clear that they are usually found in female patients aged 40 to 60 years, are often asymptomatic and their true incidence is still unknown (2-18). The general treatment recommendation for symptomatic and asymptomatic mediastinal cysts alike is surgical resection (19).

2 Case report

We present a case of a 51-year-old woman referred to our outpatient clinic due to a cystic formation in the left paravertebral space, behind the aorta, at the level of 4th and 5th thoracic vertebrae (Figure 1). This well defined lesion was discovered on an MRI scan while running diagnostics for middle back pain. We opted for a video-assisted thoracoscopic surgery (VATS) resection of the cyst.

Two 5 mm ports were made in the 4th intercostal space and one 11 mm port in the 5th intercostal space. The lung was retracted anteriorly to obtain good visualization of the aorta and the paravertebral cyst. The parietal pleura was opened with a harmonic scalpel and the cyst dissected completely from the chest wall and the aorta.

A 20 Fr chest tube was inserted through the incision in the anterior axillary line. The total duration of the procedure (skin to skin) was 32 minutes, there was no blood loss. Postoperative recovery was uneventful. The chest tube was removed on postoperative day one.

Histologically, a unilocular cyst was present in the specimen. The cyst was lined by a single layer of non-stratified cuboid to columnar ciliated epithelium that included three different cell types, the ciliated, secretory and intercalated. The subepithelial lamina propria was rather scarce, with smooth muscle surrounding the whole of the cyst (Figure 2A). The first morphological impression of the lesion was rather reminiscent of a fallopian tube. Immunohistochemically, epithelial cells were diffusely positive for WT1, PAX8 and ER (Figure 2B-D), and uniformly negative for calretinin. Due to its histomorphological and immunohistochemical characteristics, the lesion was signed-out as mediastinal cyst with Mullerian differentiation, i.e. Hattori’s cyst.

Figure 1: Chest MRI showed a cystic formation in the left paravertebral space at the level of 4th and 5th thoracic vertebrae.
On follow-up visit, the patient appeared to have no remarkable symptoms other than mild pain following the procedure, whereas the back pain persisted and did not change.

3 Discussion

The treatment of choice for mediastinal cysts is complete resection, preferably by VATS. It will provide a definitive diagnosis, exclude the possibility of malignancy and prevent the development of symptoms and possible complications. Morbidity, mortality and recurrence rates associated with resection are very low. In patients unfit for surgery, CT-guided percutaneous fine-needle evacuation is a possible less invasive treatment modality, although its recurrence rates are higher (19,20).

The presence of three types of cells in the cyst epithelium (i.e. ciliated, secretory and intercalated) is typical for a fallopian tube-type epithelium. Mullerian differentiation was demonstrated immunohistochemically with positive WT1, PAX8 and ER stains.

In the differential diagnosis of mediastinal cysts one must also consider simple mesothelial cysts, thymic cysts and bronchogenic cysts. Simple mesothelial cysts are lined by a single layer of cuboidal, non-ciliated epithelium, which is immunohistochemically WT1 and calretinin positive, and PAX8 and ER negative. In contrast to Hattori's cysts, thymic cysts typically present in the anterior mediastinum and include remnants of thymic tissue in the cyst wall. Bronchogenic cysts are lined by ciliated epithelium, just like Hattori's cyst, but in contrast to the latter, bronchogenic cyst epithelium contains goblet cells as well. Immunohistochemical characteristics differ, too, as bronchogenic cyst lack Mullerian differentiation and are therefore negative for WT1, PAX8 and ER.

Reviewing the literature, we have found 31 reports of Hattori's cysts described so far. Its true incidence is still unclear. Most papers on Hattori's cysts are case reports or small case series with only one larger series reported so far by Thomas-de-Montpréville et al. They published a systematic analysis of 163 mediastinal cysts, nine (5.5%) of which showed Mullerian differentiation. Most reported Hattori’s cyst were asymptomatic and accidental findings, suggesting that a vast majority of these lesions probably go unrecognised and that their true incidence is underestimated (1-18).

4 Conclusion

Mediastinal cysts with Mullerian differentiation are a rather newly described entity. Data suggests their prevalence may be higher since they are often
found incidentally and are typically asymptomatic (1-9,11,13,17). With the widespread use of minimally invasive techniques, complete resection is the most sensible treatment of posterior mediastinal cysts since it provides a definitive histological diagnosis and prevents possible symptoms and complications.

References


Conflict of interest
None declared.

Informed consent of the patient
Written informed consent for publication was obtained from the patient.