Awake Uniportal VATS for the Evacuation of an Extensive, Superinfected Hemothorax in a Patient with Advanced Mediastinal SMARCA4-Deficient Tumor

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Introduction

SMARCA4-deficient thoracic malignancies are rare sarcomas, identified by certain genetic mutations, particularly in the SMARCA4 gene, encoding important ATP-dependent transcription and chromatin reconfiguration mechanisms.1 They were described for the first time by transcriptome analysis and gene profiling among unclassified sarcomas only a few years ago and can be located in the mediastinum, pleura, or lung. On histopathology, the tumors showed rhabdoid to epithelioid patterns with high proliferation and mitotic activity.2,3 Beside the typical poorly differentiated morphology, diagnosis is confirmed by the loss of SMARCA4 expression and further special immunohistochemistry staining procedures. In the literature, the affected patients are described as middle-aged, mostly between 40 and 50 years, and smoking is considered a possible risk factor. Despite novel and innovative therapeutic approaches, including systemic chemotherapy and immunotherapies, as well as radiotherapy, possibly by particle therapy, SMARCA4-deficient tumors show limited treatment response and are associated with relatively poor prognosis.3,4

During recent years, minimally invasive procedures by video-assisted thoracoscopic surgery (VATS), and more recently robotic-assisted thoracoscopic surgery (RATS), have become the new standard for many indications in thoracic surgery.5 VATS is most often performed via three access sites but can also be done via one single thoracentesis, called uniportal-VATS (uVATS).6 Besides minimizing the pure surgical trauma, anesthesiological adoptions may also contribute to reduced overall procedural invasiveness and faster recovery. Nonintubated (NI) VATS are performed on nearly awake, spontaneously breathing patients.7

Case Description

A male patient in his 50’s was referred to our thoracic surgery department for further treatment of an extensive left-sided pleural effusion. The underlying diagnosis of an undifferentiated SMARCA4-deficient mediastinal tumor was made by

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► thoracic surgery
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Abstract

Background The so-called nonintubated or awake video-assisted thoracoscopic surgery (NIVATS) is performed on spontaneously breathing patients, which was shown to reduce postoperative complications and shorten hospital stay.

Case Description Awake uniportal VATS was indicated for the evacuation of an extensive, superinfected hemothorax with symptomatic mediastinal shift in a patient with advanced mediastinal SMARCA4-deficient tumor and declined condition, who did not allow a general anesthetic procedure and was not a candidate for extensive surgery.

Conclusion This short microinvasive intervention was a prerequisite to stabilize the threat to the patient’s life and thus potentially enable any further tumor-specific therapy.
radiologically guided biopsy during workup of extensive weight loss several months before. Initial systemic chemotherapy included carboplatin and etoposide; later pembro-lizumab and paclitaxel were included. In addition, simultaneous radiation therapy with protons up to a cumulative dose of 50 Gy was performed at a particle irradiation center. On restaging, the mediastinal tumor showed minimal regression in size; a newly emerged left-sided pleural effusion was treated by insertion of a small chest tube, and the patient was discharged in a stable condition. Two days after an outpatient follow-up in cardiopulmonary stable status at our emergency unit, the patient re-presented himself in a very poor clinical condition, with tachycardia (~150 beats per minute), hypotension, severe dyspnea, and increased inflammation parameters (leukocytes 23.3/nL, C-reactive protein [CRP] 71 mg/L). On emergency computed tomography (CT) scan, an extensive, loculated left-sided pleural effusion with intense mediastinal shift was shown (►Fig. 1A). No immediate chest tube was reapplied due to effusion morphology with distinct loculations, solid tissue organizations, and potential postinterventional major bleeding. Suspecting an acute, potentially superinfected hemothorax (Hb level decreased from 8.8 to 5.7 g/dL) with progressive, hemodynamically effective mediastinal shift, we chose a minimally invasive thorascopic evacuation. Because of the patient’s poor condition, the procedure was performed in close collaboration with the department of anesthesiology under local anesthesia in the awake and spontaneously breathing patient. After disinfection according to the WHO standards and sterile draping, a multilevel intercostal nerve infiltration with bupivacaine was performed. One single thoracocentesis was placed in the fifth intercostal space at the anterior axillary line. After blunt opening of the pleura, a soft-tissue retractor was placed and the hemothorax was evacuated; approximately 2 L of mostly coagulated, organized effusion was aspirated. No active bleeding was detected. Pneumopleural adhesions were taken down and the lung was carefully mobilized. Only then, could the mediastinal tumor be exposed and approached, and several biopsies were taken (histological picture; see ►Fig. 1D). After insertion of a (24-Ch) chest drain and extensive irrigation, the wound was closed in several layers. The operative time was 30 minutes, during which the patient was breathing spontaneously and communicating. Postoperative CT scan and X-ray (►Fig. 1B and C) confirmed significantly decreased left-sided opacity with a better-expanded left lung and regression of the mediastinal shift. Postoperatively, the vital signs, especially the heart rate, normalized immediately and leukocytes decreased to 13.6/nL within 5 days. In addition, the patient’s dyspnea improved gradually, allowing him to take short walks around the ward. The patient showed a clinically significantly improved condition after surgery, which allowed us to consider further tumor-specific therapies (including tumor debulking surgery) shortly after. Eleven days after the initial microinvasive uniportal VATS approach in the awake and spontaneously breathing patient, a second elaborate surgery was performed under general anesthesia for evacuation of residual hemothorax and tumor debulking. After 20 days following the awake VATS intervention, the patient was able to leave the hospital in a reduced but stable general condition. The patient was seen again by his primary oncologist to consider further tumor-specific therapy.

Discussion

Uniportal VATS on spontaneously breathing patients without any relaxation, with only minimal sedation and adequate local anesthesia (“awake VATS”), is for now the least invasive concept in thoracic surgery. Although different types of thoracic procedures including extensive anatomic pulmonary resections have been shown to be feasible by this approach, awake VATS is not an established standard approach and it remains unclear which patient groups benefit most from it.8 It might be effective for operating on a specific cohort of severely comorbid patients with restricted cardiorespiratory function, which otherwise would not qualify for any type of surgery. Our oncological end-stage patient tolerated the procedure well and quickly recovered from the procedure despite his devastated condition. This case suggests that awake VATS should be considered for short interventions in high-risk patients and could be a link between purely conservative and more invasive therapeutic strategies. Although awake VATS may never be the routine standard approach in thoracic surgery, it is a very helpful tool in challenging situations and thus should be a part of the portfolio of every advanced thoracic unit.

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Conflict of Interest
None declared.

References