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A Novel Endoscopic Technique for Biopsy and Tissue Diagnosis for a Paraspinal Thoracic Tumor in a Pediatric Patient: A Case Report

SANJAY KONAKONDLA, MD,1 JONATHAN NAKHLA, MD,1 JIMMY XIA, BSC,1 SEAN M. BARBER, MD,1 JARED S. FRIDLEY, MD,1 ADETOKUNBO A. OYELESE, MD, PHD,1 ZIYA L. GOKASLAN, MD,1 NIKOLAI G. RAINOV, MD,2 DIMITAR G. HARITONOV, MD, PHD,3 RALF WAGNER, MD,4 ALBERT E. TELFEIAN, MD, PHD1

1Warren Alpert Medical School of Brown University, Rhode Island Hospital, Providence, RI, 2MVZ Wirbelsäulenzentrum Taufkirchen b. München, Munich, Germany, 3University Hospital Heart and Brain, Pleven, Bulgaria, 4Ligamenta Spine Centre, Frankfurt am Main, Germany

ABSTRACT

Background: Conventional approaches to the thoracic spine can require extensive tissue dissection, bony disruption, and instability that may warrant the need for instrumentation and fusion. Furthermore, anterior approaches may require the involvement of various surgeons from multiple disciplines to ensure a successful operation and mitigate complications. Currently, available minimally invasive approaches still require bony removal and usually rely heavily on computed tomography (CT)-guided imaging without direct gross visualization. Endoscopic spinal procedures have provided an ultra–minimally invasive alternative to access many areas in and around the spinal column.

Methods: We present a 12-year-old boy with a right-sided 2.0 \times 3.2-cm paravertebral lesion at the level of T5. The patient successfully underwent an endoscopic approach to the lesion with minimal tissue and bony disruption for tissue diagnosis and tumor resection.

Results: At initial and 6-month follow-up, the patient remained asymptomatic and without issues.

Conclusions: We demonstrate here the feasibility and suggest the safety of a posterior ultra–minimally invasive endoscopic spinal approach to obtain a tissue biopsy of an incidentally found ventrolateral paraspinal tumor in the thoracic region in a pediatric patient. This minimal approach can prove to achieve similar results as other approaches that may otherwise necessitate more extensive or transthoracic procedures.

Minimally Invasive Surgery

Keywords: endoscopic spine surgery, thoracic tumor, paraganglioma

INTRODUCTION

Access to paraspinal tumors in the thoracic region requires thorough planning and discussion to select the most suitable approach. Generally, the available approaches can be categorized into posterior, anterior, lateral, or a combination of these approaches. Furthermore, we can subclassify these approaches into open or minimally invasive. Open posterior approaches to paraspinal tumors in the thoracic region often demand a wide exposure and significant muscle disruption and require bony removal of the posterior spinal elements with partial removal of unilateral or bilateral joints and ribs. Extensive bony removal can cause concern for iatrogenic spinal instability, which may necessitate the need for a more involved operation requiring instrumentation and fusion. Open anterior or lateral transthoracic approaches can frequently require the involvement of multiple surgeons with expertise in various disciplines for both approach and tumor resection.1 Minimally invasive or computed tomography (CT)-guided approaches to these areas can be transpedicular posteriorly or transthoracic from an anterior or lateral approach, though these also often involve extensive bony removal, poor visualization, additional radiation, and tumor contamination, among other limitations.2 These CT-guided needle approaches, moreover, do not allow for maximally safe tumor resection.

As advancements in endoscopic equipment and visibility progress and evolve, the indications for endoscopic techniques to treat various spinal pathologies are rapidly increasing.3,4 The requirement of precise localization, nuanced operative techniques, and familiarization of surgical anatomy,
however, makes endoscopic spine procedures uniquely challenging. Herein we describe the application of an endoscopic spine technique for tissue biopsy and diagnosis of a paraspinal thoracic tumor in a pediatric patient that avoids all issues of conventional approaches.

METHODS

Case Report

The patient is a 12-year-old boy with a history of a T5-T10 spinal cord arteriovenous malformation status post-embolization. At the time of follow-up, a surveillance MRI was completed that incidentally revealed a right-sided 2.0 × 3.2-cm paravertebral lesion at the level of T5 with moderate enhancement with gadolinium (Figures 1 and 2). The patient did not have any complaints or symptoms localizing to this lesion and was neurologically intact. The decision was made to obtain a tissue specimen to establish a diagnosis.

Endoscopic Operative Technique and Localization

The procedure was performed after the patient was induced under general endotracheal anesthesia without complication. The patient was placed in a prone position onto a Wilson frame. All pressure points were accounted for and padded appropriately. Fluoroscopic imaging was used for level localization and precise targeting. The incision was planned by measuring 3 cm lateral to the spinous process of T4 on the right side. Local anesthetic was given prior to incision. A spinal needle was navigated to the costovertebral joint at T5. Dilators were navigated and advanced sequentially to this target to facilitate placement of the endoscopic working channel (Figure 3A). A 6.8-mm, 30° TESSYS Joimax (Karlsruhe, Germany) endoscope was subsequently passed through the working channel. The tumor was immediately visualized upon entry, and multiple samples could be obtained.

Figure 1. MRI findings. T2 weighted sequences revealing a paraspinal enhancing lesion at the level of T5 (A: coronal plane; B: sagittal plane), ventral to the costovertebral joint on the right side (C).

Figure 2. Tumor location. Artist representation of right ventrolateral tumor at the costovertebral joint at the level of T5.
with 3.5-mm Joimax grasper forceps (Figures 3B and 4). Hemostasis was obtained with both bipolar electrocautery under direct visualization and free surface hydraulics with irrigation. Joimax grasper forceps were used to continue with tumor resection, and hemostasis techniques were applied as necessary until adequate tumor was removed. The safety of the tumor resection was based on the tumor’s proximity to the superior vena cava.

Diagnosis and Postoperative Course

The specimen histologically and immunohistochemically revealed a lobular architecture containing giant cells with abundant bright eosinophilic cytoplasm. A characteristic Zellballen structure was demonstrated (Figure 5A). The nuclei were found with mild polymorphism with chromatin and anastomosing blood vessels. Pan cytokeratin stains were mildly positive and S100 was positive in the supporting cells making up the lobular structure. The tumor was CD30 negative and CD117 positive. Synaptophysin and chromogranin A were diffusely positive (Figure 5B and 5C). The Ki-67 index was 4.8%. Final pathological diagnosis was noted to be paraganglioma. The patient did not have any family history of paraganglioma and genetic testing was pursued. Immediate postoperative MRI was completed during the initial hospital stay to assess resection and to aid in surveillance. Significant tumor resection was noted with expected residual along the superior vena cava. The patient did well and was discharged on the third postoperative day without any issues or complications. On the first postoperative follow-up visit, the patient was asymptomatic, pain free, and without complications. At 7 months, the patient was asymptomatic without any issues. Follow-up MRI at this time was completed for surveillance and did not reveal any tumor recurrence or growth.

DISCUSSION

Spinal paragangliomas are rare, slow-growing, potentially secretory tumors that rarely involve the spine. These tumors are even more rare in the thoracic region, which is evidenced by case reports of paraspinal thoracic paragangliomas.1,5 The definitive diagnosis of spinal paragangliomas is made upon histopathological examination.6 Due to non-specific radiographic findings, preoperative differential diagnosis of paraspinal tumors must include schwannoma, neurofibroma, and malignant nerve sheath tumors as well as paraganglioma. Though classified by the World Health Organization as grade I benign tumors, malignant transformation has been reported in the literature up to 14%.6 Primary treatment involves complete resection to prevent recurrence and improve progression-free survival.

Along with tumor size and location, tissue diagnosis can assist with subsequent surgical planning and multidisciplinary discussions. Various approaches for the complete resection of paraspinal tumors of the thoracic spine have been described.7
Minimally invasive techniques to access intradural lesions in the thoracic spine have been described, though traditional techniques to access the paraspinal thoracic region require both a different approach and more bony removal. Dumbbell tumors resected through minimally invasive approaches may still warrant bony removal and fusion. These open and minimally invasive techniques, however, have been primarily used only for tumor resection rather than for obtaining tissue for diagnosis, because these approaches demand more extensive exposures and arduous procedures.

CT-guided biopsy techniques are feasible alternatives to an endoscopic biopsy to access lesions in this area. The advantage of this technique allows for high-quality CT images and identification of bony landmarks for needle guidance. Disadvantages of CT-guided techniques include the lack of real-time needle passing, radiation exposure, tumor contamination, and high cost. With the endoscopic biopsy, the tissue is under direct visualization throughout the procedure on a high-definition screen. Fluoroscopic guidance can be used sparingly to minimize radiation exposure during localization. Traditional biopsy techniques can create concern for seeding tumor cells along the biopsy tract. Because of these concerns, with biopsies there can be significant clinical relevance and can have implications on future surgical planning for complete tumor resection. With the endoscopic biopsy, a tissue dilator is used to facilitate the use of a working channel. This may help mitigate contamination concerns because the surrounding tissue along the tract is protected. Though the concern of sampling error in all biopsy techniques exists, the larger working channel also allows maneuvering to create a larger surface area for more potential regions to biopsy. After the biopsies are taken under direct endoscopic visualization, areas can be inspected for obvious bleeds. Hemostasis can be immediately and effectively obtained with certainty, which is a unique advantage to the endoscopic technique compared with conventional image-guided biopsy techniques.

Ultra-minimally invasive endoscopic techniques gain entry to the areas of interest by accessing natural corridors such as the Kambin triangle, for example, during transforaminal endoscopic discectomies. Studies have also shown favorable complication rates with endoscopic spine surgery when compared with minimally invasive or open spine surgeries. In a multi-institutional study comprising 553 consecutive patients, the rate of intraoperative and postoperative complications was reported to be 2.7%. Complications reported included durotomies, epidural hematomas, recurrent disc herniations, and systemic complications.

Endoscopic procedures have been described for the treatment of thoracic discs and epidural thoracic lesions; however, these were all completed using a transforaminal technique to access the neural foramen and the epidural space. To conclusively demonstrate safety, the outcomes of more cases should be reviewed, and perhaps, cadaveric studies may be completed to assess postprocedural collateral damage and complications. Herein we describe the first totally posterior paraspinal approach for targeting, biopsy, and tissue diagnosis of a ventrolateral paraspinal thoracic tumor.

CONCLUSION

Ultra-minimally invasive endoscopic spinal surgery from a posterior approach may be a safe alternative for tissue biopsy and histological diagnosis of paraspinal lesions, which may otherwise necessitate more extensive or transthoracic procedures.
REFERENCES


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Corresponding Author: Albert E. Telfeian, MD, PhD, Department of Neurosurgery, Norman Prince Neurosciences Institute, Warren Alpert Medical School of Brown University, Rhode Island Hospital, 593 Eddy St, Providence, RI 02903. Phone: (401) 793-9132; Email: atelfeian@lifespan.org.

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