A COMBINATION OF PATHOLOGIES - FACET JOINT SYNOVIAL CYST, EPIDURAL LIPOMATOSIS AND CONJOINED NERVE ROOT ANOMALY IN THE LUMBAR SPINE OF A 45-YEAR-OLD PATIENT. A CASE REPORT

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ABSTRACT
Facet joint synovial cysts are abnormal fluid-filled cystic formations that develop from a degenerative process of the facet joints. Spinal epidural lipomatosis is a process of excessive accumulation of fat in the epidural space. Conjoined nerve root (CNR) anomaly refers to an anatomical variation in which two adjacent nerve roots share a common dural sleeve. The aforementioned three rare pathologies cause the same pathological process of stenosis, therefore compressing the nervous structures.

A 45-year-old female patient was admitted to the Department of Neurosurgery with severe low back and radicular pain, and numbness along her left leg. Antalgic posture, painful paravertebral muscle tenderness, L4 radiculopathy, and L5 and S1 dermatome hypoesthesia on the left as well as positive straight leg raising sign (Laseque) were present upon examination. CT and MRI scanning showed stenosis on L4-L5 level caused by bilateral synovial cyst formation.

Microsurgical decompression by foraminotomy at L4-L5 level on the left with dissection of the synovial cyst and epidural lipomatosis was performed. In the course of the operation CNR anomaly was discovered. The complete decompression alleviated the symptoms and after 3 uneventful postoperative days the patient was discharged.

Spinal stenosis causing compression upon nervous structures is a process that could be caused by a number of conditions. The incidence of rare pathologies and anomalies or even the more rarely occurring combination of them must be considered on daily basis in the clinical practice in order to make the correct diagnosis and plan the best possible treatment for the patient.

Keywords: neurosurgery, spinal stenosis, facet joint synovial cyst, epidural lipomatosis, conjoined nerve root anomaly, duplication of nerve root

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INTRODUCTION

Facet joint synovial cysts are abnormal fluid-filled cystic formations that develop from the degenerative process of the facet joints (1,2). The reported incidence of these lesions in magnetic resonance imaging (MRI) and computed tomography (CT) scanning varies in the different groups and imaging modalities with the highest prevalence in symptomatic population reported by Doyle and Merilees - 2.3% for anterior and 7.3% for posterior cysts (3,4,5,6). The symptomatic patients are usually in their sixties and present with low back pain, unilateral radicular pain, and neurogenic claudication, with much rarer incidence of motor and sensory disturbance (6,7,8).

Spinal epidural lipomatosis is a process of abnormal excessive accumulation of fat in the epidural space, which leads to narrowing of the spinal canal and compression of the nervous structures. The mean age of patients is 69.4±10.9 years and the usual manifestation in the symptomatic population is low back pain, myelopathy or radiculopathy due to compression at different spinal levels (9,10,11).

The term conjoined nerve root (CNR) anomaly refers to an anatomical variation in which two adjacent nerve roots share a common dural sleeve at some point in their course before separating to follow individual courses through the same or separate foramina. On its own in normal conditions the CNR anomaly is usually asymptomatic. However, due to overcrowding of the nerve canal, compression of different origin may lead to radiculopathy, neurogenic claudication and neurological deficit (12,13,14).

CASE PRESENTATION

A 45-year-old female patient was complaining of low back, radicular pain and numbness along her left leg that were gradually intensifying to a severe level for few weeks prior to admission to the Department of Neurosurgery. Upon examination antalgic posture and gait were prominent. Paravertebral muscle tenderness and painful trigger points in the lumbar spine were present as well as L4 radiculopathy and L5 and S1 dermatome hypoesthesia for the left leg. On the same side, the straight leg raising sign (Laseque) was positive with maximum height of 20 degrees. As the laboratory test results were normal, the imaging was essential. First a CT scan was performed and it showed degenerative changes in the lumbar spine with central stenosis on L4-L5 level – 14 mm, and L5-S1 – 8 mm (minimum normal width of the vertebral canal in the lumbar region - 15 mm). In addition to that, a lesion causing lateral recess and foraminal stenosis was visible at L4-L5 level on the left (Fig. 1).

As an imaging modality with better visualization of the tissues, an MRI scan was performed in order to aid the correct identification of the pathology of our patient. It showed bulging of the intervertebral disks at L4-L5 and L5-S1 spinal levels and spondyloarthrosis of the facet joints at L4-L5 level with formation of bilateral synovial cysts resulting in central stenosis of the spinal canal (12.7 mm sagittal length), left lateral recess narrowing, and bilateral foraminal stenosis. In addition to that, anterior spondylolisthesis...
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sis at L4-L5 vertebral level was also present (Fig. 2), (Fig. 3).

The medical history of the patient, the physical examination and the imaging results pointed to a diagnosis of spinal stenosis and nerve root compression in the lumbar region on the left due to facet joint synovial cyst formation. After careful reviewing of the case and all of its aspects, a decision for surgical treatment was made in order to alleviate the neurological symptoms.

The surgical procedure started with assessment of the correct vertebral level using a mobile Ziehm Vision FD Vario 3D C-arm X-ray system, followed by linear incision from L4 to L5 level. After dissection of the left paravertebral muscles using Zeiss OPMI PENTERO 900 surgical microscope under optical magnification, a left foraminotomy at L4-L5 level was performed. Compression of the dural sac by the synovial cyst on the left and epidural lipomatosis were prominent. In the course of dissection of the formations, an anatomical variation of duplication of L5 nerve root was discovered. Complete decompression of the nervous structures was performed by excision of the synovial cyst and the epidural lipomatosis. In addition, INCERT-S anti-adhesion gel was applied (Fig. 4), (Fig. 5).

RESULTS
The surgical procedure done in the Department of Neurosurgery - total excision of both the synovial cyst and the epidural lipomatosis, allowed complete decompression of the nervous structures. The identification of the duplicated L5 nerve root did not change the course of the planned intervention as the left foraminotomy and the careful dissection of the pathological formations provided decompression of the two roots in the stenosed left foraminal canal as well. In addition to that, the resected material was sent for histological verification of the clinical diagnosis of facet joint synovial cyst. After uneventful postoperative period of 3 days, the patient was discharged with improvement of the neurological symptoms.

DISCUSSION
The first point that deserves attention in this case is the presence of epidural lipomatosis. This

Fig. 3. Preoperative MRI scan. A. T2 coronal section, B. T1 sagittal section, C. T2 sagittal section, D. STIR sagittal section, E. T2 axial section. At L4-L5 and L5-S1 spine levels there is building of the intervertebral discs. A synovial cyst at L4-L5 level on the right is marked by a red arrow on all images.

Fig. 4. Intraoperative image from Zeiss OPMI PENTERO 900 Surgical Microscope showing a synovial cyst (blue arrow).

Fig. 5. Intraoperative image from Zeiss OPMI PENTERO 900 Surgical Microscope showing double root anomaly (blue arrows) and epidural lipomatosis (green arrow).
pathology is believed to develop either when there is hormonal disturbance with corticosteroid therapy, endocrine disorders and obesity or idiopathically (10,17,18). As it is a rare pathology with incidence of symptomatic cases reported in only 0.1% of all the patients with low back pain, there are no clinical trials assessing the results after conservative and surgical treatment (10,11). Conservative measures, such as termination of corticosteroid therapy, reduction of weight, and treatment for endocrine disorders can be implemented when the etiology of elevated hormonal levels is suspected. For idiopathic cases, such as our patient, surgical decompressive laminectomy and resection of the fat tissue is the treatment of choice (10,18).

Another aspect of this case is the diagnosis of facet joint synovial cyst. As it was mentioned in the introduction, this pathology is a rare cause of symptomatic spinal compression. Because of the similarity of the clinical manifestation of facet joint synovial cysts to that of other vertebral degenerative pathologies causing compression of the nervous structures, it is important for the clinician to consider this lesion in the differential diagnosis (6,8). A meta-analysis comparing the operative treatment methods for facet joint synovial cysts (percutaneous aspiration and surgical decompression with or without fusion) in 870 patients in 50 studies was performed in a study last year. The results showed full resolution of the symptoms in the percutaneous group in 58% compared to 90% in the decompression one. In addition to that, 29% in the percutaneous group underwent more than 1 aspiration and 38% were indicated for surgery in the follow-up period (19). This study and its results are comparable to other sources and published clinical experience so that decompressive surgical approach is recommended as a treatment with best possible outcome for the patients (6,8). The question about spinal stability in patients with facet joint cysts remains disputable though. In a number of studies, the combination of synovial cysts and preoperative spondylolisthesis suggests instability. In these cases, it is recommended that patients underwent decompression and fusion surgery than decompression alone or these patients would probably require additional fusion surgery after the initial decompressive one (20). In our patient, spondylolisthesis was also present but due to the lateral localization of the cyst with lateral recess and foraminal stenosis, the surgical approach was decompression by foraminotomy as in this way the spinal stability would not be compromised.

In our case, attention should also be paid to the presence of double root anatomical variation. Conjoined nerve root anomalies tend to be asymptomatic until a coexisting pathology causes compression of the already narrowed by the duplicated root lateral recess and foramen. The clinical manifestation of CNR is similar to that of the causative lesion and so the underlying pathology is not usually addressed. Without adequate decompression by foraminotomy in addition to the laminectomy, the stenosis and subsequently the symptoms are not relieved. In such cases, an undiagnosed CNR can be considered as a possible cause for failed spinal surgery (13,14,21). As it is said, when it comes to women age is an important issue. Reviewing the literature it appears that facet joint synovial cysts and epidural lipomatosis as diagnosis of symptomatic compression of the spinal cord usually present in patients in their 50s, 60s and older (2,3,9,11). This late onset might be related to the degeneration process of the vertebral column as possible pathogenesis of these lesions (3,8,14). Nonetheless our patient is 15 years younger than the usual occurrence period and the fact that she has a combination of three pathologies, two that cause stenosis - facet synovial cyst and epidural lipomatosis, and a double root anomaly that narrows the lateral recess and the intervertebral foramen seems too much of a coincidence to be ignored.

**CONCLUSION**

We presented you a case with a combination of three rare conditions that cause the same pathophysiological process - stenosis with typical manifestation in our patient. This case was reported in order to show that a careful evaluation of the symptoms and their possible causative pathologies should always be done. The good physician should have a wide range of differential diagnoses as well as think about the mechanism, which caused the given condition. The incidence of rare conditions, anomalies, combination of pathologies and, as in our case, combination of rare pathologies and anomalies should not stay just in the books or articles, but must be considered on a daily basis in the clinical practice in order
to make correct diagnosis and plan the best possible treatment for the patient.

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