Concomitant Abdominal Aortic Aneurysm and L2-L3 Disk Herniation: Is There a Relationship Between Them?

Introduction

Low back pain (LBP) is most commonly caused by mechanical dysfunction. However, 1-2% of cases are caused by nonmechanical spinal disorders or visceral disease, such as abdominal aortic aneurysm (AAA) (1). Factors potentially involved in chronic LBP in patients with AAA include the size of the aneurysm, increased pressure on the vascular wall and surrounding neural elements, and erosion of the vertebral body (2). Several cross-sectional studies have suggested that vascular diseases in the lumbar area are associated with disk degeneration (DD) and LBP (3-5). The correlation of atherosclerotic changes to DD was stronger in the upper levels of the lumbar spine than in the lower ones (5). AAA can cause lumbar DD and clinically mimic upper lumbar disk herniation. The incidence of herniated upper lumbar disks is less than that of lower disks (6). Pathological fractures due to osteoporosis, bone metastasis, and spinal tumors must be considered.

ABSTRACT

Abdominal aneurysm (AAA) is a critical disease and is often asymptomatic. It can cause diminished lumbar blood flow. Impaired blood flow in the lumbar arteries is significantly associated with decreased diffusion in lumbar disks and may lead to progressive disk damage. AAA can clinically mimic upper lumbar disk herniation. Disk herniation is rarely seen in the upper lumbar region since the upper part of the lumbar spine is less mobile than the lower region. To the best of our knowledge, no reports have presented concomitant AAA and L2-L3 disk herniation. We describe a 48-year-old man treated for symptomatic AAA followed by L2-L3 disk herniation at an interval of 3 months.

Keywords: Low back pain, abdominal aortic aneurysm, disk herniation, disk degeneration


© Copyright 2021 by the Bezmialem Vakıf University

Address for Correspondence: Yakup Erden, İzzet Baysal Physical Treatment and Rehabilitation Training and Research Hospital, Physical Medicine and Rehabilitation, Bolu, Turkey
E-mail: yakuperden@hotmail.com ORCID ID: orcid.org/0000-0003-0788-7532

Received: 11.06.2020
Accepted: 19.02.2021
considered alternatively when assessing upper lumbar pathology. AAA and L2-L3 disk herniation are rarely concomitant, as they were in the present case.

Case Report

A 48-year-old man, a current smoker, with a previous diagnosis of hypertension first presented with severe localized LBP after walking 200 m. The pain made walking difficult but was relieved by resting for at least 5 min. It persisted for two months and was not ameliorated by analgesics or myorelaxant agents. The pain was rated 9/10 on a visual analog scale (VAS), and the patient reported no night pain or morning stiffness. The patient is a maintenance technician and had moderate physical activity levels. He had no trauma history. At physical examination, the range of motion testing of the lumbar spine was normal but painful, and palpation of the lumbar spinous process segments and paravertebral muscles also elicited pain. The straight leg raise test was negative. The range of motion of the hip joints was normal, and no pain and restriction were detected at the hip joints in the flexion-adduction-external rotation test and flexion-adduction-internal rotation test. Sacroiliac joint provocation tests were negative. Gait analysis was normal, and no pelvic asymmetry was detected. Neurological examination was unremarkable, and the patellar reflex was normoactive. Arterial pulses in the lower limbs were normal. Edema, leg diameter difference, and discoloration were not observed in the lower extremities. The patient did not report systemic symptoms such as fever, muscle pain, fatigue, or weight loss. Results of the systemic examination were normal. The patient had not sought medical assistance for any systemic disease, except for hypertension. Results of laboratory investigations (such as complete blood count, erythrocyte sedimentation rate, and C-reactive protein) were within the normal limits. Magnetic resonance imaging (MRI) revealed a thrombosed aneurysmatic dilatation (68x60 mm in the widest part) of the wall in an approximately 12 cm segment of the abdominal aorta and L2-3 left paracentral extruded disc herniation with lateral recess stenosis. Loss of signal intensity due to degeneration was also detected in the intervertebral discs and more markedly in the upper lumbar disks (Figure 1). The patient was referred to the cardiovascular surgery department where an endovascular stent graft was inserted in the aortoiliac position (Figure 2). No complication was observed during or after the intervention, and the clinical symptoms improved significantly. Following surgery, the patient started walking 30 min to 1 h daily. At six weeks following surgery, the patient did not report any episodes of low back or abdominal pain.

Three months later, the patient was readmitted to our clinic again with LBP extending to the left lower extremity. Physical examination revealed tenderness over the lumbar spinous process segments, and the femoral nerve stretch test was positive on the left side. The lumbar lordosis had decreased, and antalgic gait was present. The cardiovascular surgery department was consulted in terms of contraindication before treatment. Since the case was compatible with discogenic LBP, 15 sessions of physiotherapy including superficial and deep heating and burst-type transcutaneous electrical nerve stimulation were applied. McKenzie and stabilization exercises were performed. The radiating LBP improved, and the VAS score decreased from 8 to 2 points.

Discussion

The pathogenesis of LBP is multifactorial. AAA can induce LBP. It is mainly caused by arterial stenosis, which results in the reduction of blood flow in the affected area. Initially, no symptoms were observed at rest, but patients may experience claudication during exercise. It may also play an important role in the mechanism of painful degenerative disc diseases. In this case, the patient presented early with claudication and lately with lower radiating back pain.

AAA is defined as a weakening of the aortic wall involving progressive, full-thickness local dilation >150% of the normal diameter. Diameters exceeding 3 cm are clinically significant. Larger aneurysms expand at a faster rate than smaller aneurysms. The risk of spontaneous rupture increases exponentially with...
the aortic diameter. Rupture accounts for more than 40% of mortality cases (7). Thus, surgery is generally recommended when the AAA exceeds >5.5 cm in diameter in men and >5.0 cm in women, in the presence of symptoms, and in the case of a rapid increase in size (8). In the present case, endovascular aneurysm repair was performed since the risk of rupture was high and symptoms developed. Previous studies have suggested that arterial surgery (when possible) is effective for LBP in patients with AAA (9).

Although the etiology of an AAA is still not completely understood, major risk factors for an AAA include hypertension, age >60 years, male gender, smoking history, atherosclerosis, coronary artery disease, and a genetic/family history of AAA (10). AAA is not normally screened for unless the patient has these specific risk factors. In a male patient aged >60 years with one or more additional risk factors, abdominal ultrasonography is initially recommended. In the present case, AAA had been diagnosed incidentally during the investigation of other spinal pathologies.

Autopsy and radiology studies have documented a correlation between vascular disease and DD. Lumbar artery stenosis is significantly associated with decreased diffusion of the lumbar disks. Intervertebral discs receive nutrients by diffusion mainly through the vertebral endplates. Diminished blood flow to the vertebrae may therefore reduce diffusion of nutrients and oxygen to the disk and hamper the disposal of harmful metabolites. Beckworth et al. (11) also found an association with other degenerative changes such as facet arthritis and lumbar canal stenosis. Interestingly, the correlation was strongest in patients aged ≥61 years. Various contributing factors may emerge with advancing age. Similar to vascular diseases, DD begins early in life. Asymptomatic individuals therefore commonly exhibit degenerative changes (12). Trompeter and Paremain reported a case series of four patients who had incidental AAA detected on lumbosacral MRI. All patients had degenerative spinal diseases. Kim et al. (13) reported a case in which AAA with mural thrombus coexisted with aggravated L3-L4 intervertebral disc extrusion. Surgery was performed for both pathologies in that case (14).

The upper three segments of the lumbar spine are supplied only by the segmental lumbar arteries, originating from the posterior wall of the abdominal aorta. The L1, L2, and L3 arteries mainly go to the L1-2, L2-3, and L3-4 disk spaces, respectively. Stenosis of the ostia of the lumbar arteries may trigger degenerative changes. With its location at the end of the nutrient chain, the disk is affected earliest by poor circulation. Insufficient blood supply may render the disks more vulnerable to mechanical stress and thus result in disk damage (5). AAA is frequently associated with diminished lumbar blood flow, and 95% of AAAs are located in the infrarenal region (L1-4 level). The prevalence of complicated lesions associated with atheromatous plaques (i.e., plaques with necrosis, thrombus, and calcification) increases rapidly at age 44-65 years. Either atheromatous plaques or thrombotic layers accumulate gradually inside the aneurysm. This then gradually obstructs the corresponding lumbar arteries but allows sufficient time for collateral pathways to develop. Therefore, ischemic symptoms in the lumbar region are rarely found even after AAA surgery. Symptoms may be observed only during exertion when reduced blood supply fails to meet demand (15). In the present case, degeneration was particularly marked in the upper lumbar disks, and symptoms developed following exertion.

The upper lumbar disks are generally defined as the L1-L2 and L2-L3 levels. Disk herniation is rarely seen in this region since the upper part of the lumbar spine is less mobile than the lower region. Upper lumbar disk herniations constitute 1-2% of all herniated lumbar disks and are mostly seen in the older population (16). Clinical signs associated with upper lumbar disk herniation are nonspecific, with most patients reporting low back and leg pain. The radicular pattern of back pain may not be clear and not reflect the true level of the disease. The femoral nerve stretch test, also known as reverse straight raise test, is positive in 84-95% of the patients with upper lumbar disk herniation, as in the present case (17). The nerve root in the upper lumbar spine does not innervate any specific muscles. In some cases, the clinical signs of radiculopathy do not match the electromyographic signs, so electromyography has no absolute diagnostic value. Accurate diagnosis is often difficult without radiological findings. Surgical outcomes of upper lumbar disk herniation are less satisfactory than those involving lower disks (18). Therefore, conservative treatment was applied in the present case. In the management of patients with upper lumbar degenerative disk diseases, the possibility of comorbidities such as an AAA should not be overlooked because a delayed diagnosis of AAA puts the patient at risk of aneurysmal rupture.

AAA is a rare entity that may be asymptomatic. An advanced disease has been correlated with LBP and DD, in which a degenerative “cascade” of events is triggered. A gradual decrease in disk flexibility and increased ruptures in annulus fibrosis may lead to herniation of the nucleus pulposus. AAA can therefore present with localized back pain or radiculopathy. Further studies are needed to confirm the causal association.

**Ethics**

**Informed Consent:** An informed consent was obtained from the patient.

**Peer-review:** Externally peer reviewed.

**Authorship Contributions**


**Conflict of Interest:** No conflict of interest was declared by the authors.

**Financial Disclosure:** The authors declared that this study received no financial support.
References


