Iliopsoas hematoma due to muscular rupture following defibrillation

Artan Jahollari, M.D.,1 Raif Cavolli, M.D.,1 Murat Tavlasoglu, M.D.,2 Ferat Sallahu, M.D.,3 Shkelzen Muriqi, M.D.4
1Department of Cardiovascular Surgery, International Medicine Hospital, Pristina, Kosova
2Department of Cardiovascular Surgery, Diyarbakir Military Hospital, Diyarbakir, Turkey
3Department of General Surgery, International Medicine Hospital, Pristina, Kosova
4Department of Anaesthesia, International Medicine Hospital, Pristina, Kosova

ABSTRACT

We describe a 62 year old patient who presented with acute anterior ischemia and subsequently developed an iliopsoas hematoma. The patient was treated surgically due to rapid progression and femoral neuropathy, and the iliopsoas muscle rupture was diagnosed intraoperatively. The rupture was related to the external electrical defibrillation the patient had on admission. This was a rare case, and we hope the report would help to raise physicians’ awareness regarding this complication and treatment.

Key words: Defibrillation; hematoma; iliopsoas rupture.

INTRODUCTION

Iliopsoas hematoma is a rarely seen form of retroperitoneal hemorrhage. The causes are various and the treatment differs based on the underlying pathology and clinical situations of the patient.

We herein report the case of a patient who was admitted with acute anterior ischemia and afterward developed an iliopsoas hematoma. The patient underwent an operation and identified with an iliopsoas rupture, which was related to external electrical defibrillation. We could not find a similar case reported in the medical literature, and we think that our case report may be the first of its kind to be published.

CASE REPORT

A 62-year-old male patient, who presented with severe retrosternal pain, was diagnosed with acute anteroseptal ischemia. Several minutes after his admission he experienced ventricular fibrillation and was immediately converted with a single 200 joule shock of a biphasic defibrillator. Emergent coronary angiography via the right femoral artery was performed, revealing total thrombotic occlusion of the proximal LAD (left anterior descending artery) which was treated successfully with PCI (percutaneous coronary intervention). 5000 IU of unfractioned heparin was administered intravenously during the procedure and the following treatment included subcutaneous LMWH (low molecular weight heparin, enoxaparine sodium 6000 anti-Xa IU/0.6 ml, twice a day), aspirin 100 mg /day and clopidogrel 75 mg/day. On the 2nd day he complained of left flank pain and a decrease in hemoglobin was observed. An abdominal ultrasound revealed the presence of a left retroperitoneal hematoma. The magnetic resonance imaging (MRI) showed a large hematoma of the left iliopsoas muscle (Figure 1). Anticoagulants and antiagreggants were immediately stopped and the patient was followed conservatively, supplied with fresh frozen plasma and fresh blood. Despite ruling out active gross hemorrhage by a radial angiogram, the progressive decrease of hemoglobin persisted and the pain severity increased, radiating to the left leg. He developed abdominal defense and paresis of the left leg, which was attributed to femoral neuropathy. Urgent surgery consisting of a laparatomic exploration was performed on the 4th day. Approximately 1000 ml of aspirated fluid was collected from the retroperitoneal area. The psoas fascia was incised and a
mixture of 1000 ml of liquefied and clotted blood was removed. Rupture of the muscle fibers was observed and intact muscle parts were sutured together (Figure 2). During the postoperative period he was treated with LMWH (enoxaparine sodium 6000 anti-Xa IU/0.6 ml, daily). He had a rapid recovery and was discharged after 5 days on antiaggregant therapy (aspirin 100 mg/day and clopidogrel 75 mg/day).

DISCUSSION

Hematoma of the iliopsoas muscle is a rare and challenging condition. Anticoagulant use, hemophilia, trauma, alcoholic cirrhosis, etc. appear to be some of the main causes. However, the nature of this pathology is not fully understood and a many cases develop spontaneously. Clinical manifestation includes abdominal and flank pain, abdominal distention and defense, hemorrhagic shock features, and leg or groin pain in presence of femoral nerve compression. Abdominal ultrasound provides a first glance and quick information about the ongoing condition. However, more specific tools, such as MRI and CT-scan, are necessary to confirm the pathology and help to make a proper decision.

There are different treatment modalities for iliopsoas hematomas. Good outcomes of conservative therapy have been successfully reported, and we agree that this is the first choice in stable patients. Patients suffering from hemophilia or that are under anticoagulant therapy seem to benefit the most from this choice. Local puncture can also be applied to drain the hematoma, but it carries the risk of further complications and is not helpful in removing clotted blood. Selective embolization of active branch bleeding is an option in selected patients and requires expertise. However, the definite treatment in complicated and unstable patients remains surgery. The presence of severe pain and the development of neurologic symptoms due to compression and hemorrhagic shock are some of the main indications for surgical decompression. Different sources strongly support the evacuation of the hematoma immediately after the onset of femoral neuropathy, and associate it with a great likelihood of a return to pre-event neurological status. In our case, surgery was obligatory by abdominal defense, femoral neuropathy, and hemodynamic progression; all together compromising the general situation. Despite the delicate heart condition, our patient tolerated the surgery well and recovered quickly.

During the surgery we discovered that rupture of the iliopsoas was the reason for the hematoma. Although it helped to increase the hemorrhage, anticoagulant usage cannot be blamed, as it does not explain the rupture of the muscle. On the other hand, defibrillation was the only physical trauma the patient experienced and it is a well-known fact that it provokes skeletal muscle contractions. The psoas major muscle itself originates at the vertebral bodies of T12 and L1-L5 and their associated intervertebral discs, a region anatomically close to the apical defibrillator pad. External electrical defibrillation is an old and safe treatment modality in ventricular fibrillation, and very few cases with mechanical complications of external defibrillation have been reported. Most complications include skin burns, but bone and muscle involvement have also been described, including sternal and vertebral fractures, myocardial lesions and pectoral muscle necrosis. Iliopsoas rupture due to defibrillation has not been previously reported and although it is a rare complication, it must be kept in mind owing to its potential severity.

As a result, we herein described a rare case of iliopsoas hematoma due to muscular rupture. We strongly believe that the rupture resulted as a complication of external defibrillation. Conservative treatment is the therapy of choice in iliopsoas hematomas. However, surgery should be considered in rapidly progressing, neurologically complicated and hemodynamically unstable cases.

Conflict of interest: None declared.

REFERENCES

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Defibrillation-related iliopsoas hematoma due to muscular rupture following defibrillation

Dr. Artan Jahollari,1 Dr. Raff Cavoli,1 Dr. Murat Tavlasoglu,2 Dr. Ferat Sallahu,3 Dr. Shkelzen Muriqi4

1International Medicine Hospital, Kalp ve Damar Cerrahisi Bölümü, Pristina, Kosova
2Diyarbakir Asker Hastanesi, Kalp ve Damar Cerrahisi Bölümü, Diyarbakir, Türkiye
3International Medicine Hospital, Genel Cerrahi Bölümü, Pristina, Kosova
4International Medicine Hospital, Anestezi Bölümü, Pristina, Kosova

Bu yazida, akut anterior iskemi ile başvuran bir hasta da iliopsoas hematomu gelişen 62 yaşında bir hasta sunuldu. Çabuk ilerleme ve femoral nöropati gelişmesi nedeniyle hastanın cerrahi tedavi edildi ve ameliyat sırasında iliopsoas kas yırtığı tespit edildi. Kas yırtığı, kabul esnasında uygulanan eksternal elektrik defibrilasyonu iliskiliydi. Bu nadir oltu sunumunun komplikasyon ve tedavisinin hakkında hekimlerin farkındalığını artıracağını umuyoruz.

Anahtar sözcükler: Defibrilasyon; hematom; iliopsoas yırtığı.