Case Presentation

The Management of a Patient Presented with Fournier’s Gangrene Associated with Undiagnosed Ulcerative Colitis: A Case Report

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Abstract

Introduction: Fournier’s gangrene is a rare polymicrobial necrotizing fasciitis of the perineal and perianal regions, found to be associated with colorectal diseases and urogenital operations. We present here a difficult management of a patient presented with Fournier’s gangrene associated with undiagnosed Ulcerative colitis.

Case Presentation: A 25 year-old man presented with scrotal edema and pain was diagnosed with Fournier’s gangrene. The patient was hospitalized and treated with wide-spectrum antibiotics and surgical debridement of the necrotic tissue. During follow up the patient developed a toxic megacolon which was successfully treated by colectomy and end-leocestomy. The pathological findings of the surgical specimen were resulted as a diffuse active colitis supporting ulcerative colitis.

Conclusion: Fournier’s gangrene is an aggressive infection with high mortality rates which requires a multidisciplinary approach.

Keywords: Fournier’s Gangrene; Ulcerative Colitis; Toxic Megacolon

Introduction

Fournier’s gangrene is a rare form of necrotizing fasciitis localized in the perineal and perianal regions, with a mortality rate ranging from 15% to 50% [1]. There are multiple causes of this polymicrobial infection including urological diseases and procedures, colorectal diseases and superficial trauma by foreign bodies. Predisposing factors are including Diabetes mellitus, chronic alcoholism, immunosuppressive diseases such as HIV infection and myeloproliferative diseases and malnutrition [2-4]. The most common isolated microorganisms are Escherichia coli and Bacteriodes [5]. The standard treatment of Fournier’s gangrene is surgical debridement of the necrotic tissue and wide-spectrum antibiotics.

Fournier’s gangrene complicating inflammatory bowel disease has been reported in four patients so far, two with Ulcerative colitis and two with Crohn’s disease [6-9].

We present here an exceptional case management of a patient presented with Fournier’s gangrene associated with undiagnosed Ulcerative colitis.

Case Presentation

A 25 year-old man complaining of the new onset of rectal pain, swelling and tenderness in the scrotum applied to the emergency service. The patient denied having abdominal pain, nausea, constipation or diarrhea. The patient had experienced rectal bleeding and mucus with defecation for three months. The patient was seen by a general practitioner for abdominal pain and mucus stool. He was started on antibiotics. The patient denied any other pathological findings detected in the abdomen.

In laboratory tests, leukocyte levels were 16.500 (normal limit: 4200-10.600), CRP (C - Reactive Protein) level was 300mg/L and billuribine levels were 3.7mg/dl (total billuribine) and 1.7mg/dl (direct billuribine). AST (Aspartate Aminotranspharase) and ALT (Alanine Aminotranspharase) levels were found to be mildly elevated. Our differential diagnosis included scrotal abscess, perianal abscess and Fournier’s gangrene. We decided to proceed with CT (Computer Tomography) scan of the abdomen.

On physical examination bilateral scrotal congestion was significant with subcutaneous emphysema and hyperemia. Subcutaneous emphysema was reaching to the right groin and lower quadrant of the abdomen. In digital examination tenderness and the fluctuance was present in the right ischiorectal fossa.

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CT scan of the abdomen revealed subcutaneous emphysema in the right lower quadrant reaching out the right inguinal region, right scrotum and ischiorectal fossa. Although the significant increased density and thickening in the subcutaneous soft tissue of the abdominal wall, the peritoneum was intact (Figure 1-2). There were not any other pathological findings detected in the abdomen.

The patient was hospitalized with the presumptive diagnosis of Fournier’s gangrene and underwent operation. The incision was started in the lower crease of the right scrotum and reached out to the right inguinal and the lower quadrant region. The opening of Fournier’s gangrene was made and was found to be associated with Fournier’s gangrene. The patient was treated with a wide-spectrum antibiotics and debridement of the necrotic tissue. During follow up the patient developed a toxic megacolon which was successfully treated by colectomy and end-leocestomy. The pathological findings of the surgical specimen were resulted as a diffuse active colitis supporting ulcerative colitis.

Summary

Fournier’s gangrene is a rare polymicrobial necrotizing fasciitis of the perineal and perianal regions, found to be associated with colorectal diseases and urogenital operations. We present here a difficult management of a patient presented with Fournier’s gangrene associated with undiagnosed Ulcerative colitis. The patient was treated with a wide-spectrum antibiotics and debridement of the necrotic tissue. During follow up the patient developed a toxic megacolon which was successfully treated by colectomy and end-leocestomy. The pathological findings of the surgical specimen were resulted as a diffuse active colitis supporting ulcerative colitis.
of the scrotum foul-smelling brown purulence discharged. The testicles were not involved. The extensive necrotic tissue debridement of the perianal and scrotal region were done including the skin and the fascia. It was also revealed that the external anal sphincters were damaged and the sphincters were primarily repaired. Necrotic tissue samples were taken for aerobic and anaerobic cultures. Vacuum assisted closure (VAC) technique was chosen as the wound dressing (Figure 3-4).

The patient was started on broad-spectrum antibiotics with vancomycin and meropenem. The routine follow-up was done with CBC (Complete Blood Count), liver and kidney function tests. Although leukocyte levels decreased for first three days, then elevated to 19,500 (normal limit: 4200-10,600). Total and direct billirubine levels were elevated up to 4.9 and 2.8 mg/dl. AST and ALT levels were 226 and 72 U/L, ALP (Alkaline Phosphatase) and GGT (Gamma Glutamine Transpharase) levels were 666 and 441 U/L. CRP was measured 147mg/L.

On postoperative 8th day, the patient started having abdominal pain, distention and fever with septic appearance. Another CT scan of the abdomen was performed revealing distention of transverse segment of the colon up to 8-9 cm with extensive fluid surrounding the colonic segments. With the presumptive diagnosis of toxic megacolon, explorative laparatomy was performed (Figure 5-6). Operative findings were extensive distention of the ascending, transverse and the descending segments of the colon. A perforated area was located in the transverse segment of the colon. The sigmoid colon and the rectum were intact. We decided to proceed with subtotal colectomy and end-ileostomy.

The patient was taken to the intensive care unit and the same antibiotic regimen with vancomycin and meropenem were continued for 13 days. CRP and procalcitonin levels were followed regularly. Cultures of the necrotic tissue samples were resulted as Acinetobacter baumanii and Pseudomonas aeroginosa. The antibiotic regimen was changed to linezolid, colistine and tazocin. The regimen was continued for 24 days.

The patients’ debrided wound was continued to be dressed with VAC for 34 days. The VAC dresses were changed every 48 to72 hours. During follow-up, septic appearance of the patient regressed. On postoperative 36th day, the wound was partially closed by secondary suturing and the remaining part was left for spontaneous epithelialisation.

The pathological findings of the surgical specimen were mucosal ischemia and necrosis of the resected segments of the colon. There were multiple mucosal ulcerated lesions, cryptic distortions, and significant lymphoid infiltration with aggregate formation consistent with inflammatory bowel disease. All of the dissected lymph nodes were found to be reactive (Figure 7a-7c).

Also the pathological findings of the colonoscopic biopsies were found to be similar, with increasing cryptitis formation and lymphoid infiltration in mucosa of the distal segments of the colon and the rectum. The granuloma formation was not detected. All these results
suggested diffuse active colitis supporting ulcerative colitis. The patient was started on salofalk and discharged on postoperative 42th day.

**Discussion**

Fournier’s gangrene is an aggressive necrotizing fasciitis of perianal, genitourinary and perianal regions. The source and the entrance of the infective microorganisms are commonly found to be anorectal lesions and cutaneous trauma by foreign bodies. The main predisposing factors are considered to be immunosuppressive diseases, Diabetes Mellitus and chronic alcoholism.

The case report we present here, distinguishes itself with the patients’ younger age and the underlying undiagnosed ulcerative colitis which can easily mislead. Although the disease mostly affects elder patients, with recent studies the age range was found to be between 22 to 88 years old [10]. Various colorectal diseases and procedures have been associated with Fournier’s gangrene; such as iatrogenic colon perforation [11], sigmoid colon perforation via dislocated pancreatic stent [12], perforation of a malignant tumor located in rectum [13].

The possible triggering factors for our patient are undergoing inflammatory bowel disease and performed colonoscopy. So, a thorough evaluation for underlying disease including any previous symptoms such as rectal bleeding, loose stools and weight loss, and previous procedures should be overtaken.

The main standard treatment is surgical debridement of the necrotic tissue and wide-spectrum antibiotics, regardless of associated disease. However, in patients with inflammatory bowel disease, anti-inflammatory regimens such as mesalamine, salofalk and 6-MP can also be effective in alleviating the disease. Hyperbaric oxygen therapy, various skin grafts, and vacuum-assisted closure techniques are also found to be effective in reducing the hospital stay and patients’ discomfort [14,15].

**Conclusion**

We present here an exceptional case management of a patient presented with a Fournier’s gangrene associated with undiagnosed Ulcerative colitis. Fournier’s gangrene is an aggressive infection with high mortality rates despite of new treatment modalities. A multidisciplinary approach is crucial for this devastating disease.

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