

CHAPTER 5

DIAGNOSIS AND TREATMENT OF FOURNIER'S GANGRENE

Cemil KUTSAL¹

Fournier's Gangrene was described in 1883 by Jean Alfred Fournier, a dermatologist, and venerologist from Paris, and is referred to by its specific name (1).

Fournier's Gangrene is a severe disease that affects the genital, perianal, and perineal regions. When the diagnosis and treatment are delayed, it progresses rapidly between the fascial planes and causes widespread soft tissue necrosis. The disease typically spreads aggressively between the fascial planes and involves surrounding soft tissue. The spread of the infection causes microemboli in the arterial vessels, causing blood circulation disorder and tissue necrosis in the surrounding soft tissue and fascial planes (2).

This process spreads rapidly between Dartos, Colles, and Scarpa fascia planes(2). Due to the involvement of the subcutaneous and fascial areas first, doctors may be unable to diagnose it in the early stages of the disease. The overlying skin often appears as uncomplicated cellulitis(2-4).

Urogenital infections, anorectal infections, and trauma are the primary etiologic factors of Fournier's Gangrene. It is a polymicrobial condition usually caused by various aerobic and anaerobic microorganisms(5-7). The most common and cultured organisms are gram-negative bacteria in polymicrobial form. These include Group A Streptococci and Staphylococcus aureus, and E. Coli and Pseudomonas aeruginosa. (8,9).

These bacteria can enter the body from the urinary, intestinal systems, or dermal routes. Sometimes, urinary tract infections and perianal abscesses may also cause infection(2).

In FG, it may first give symptoms as local infection depending on the way of entry into the body. It may start as a local infection around the rectum in the perineum, the urethra, and the scrotum in the genital area (10,11). Although Fournier's Gangrene is more common in men and the elderly, it can affect both sexes and all age groups (12-15).

¹ Dr., İstanbul Şişli Hamidiye Training and Research Hospital, Urology Clinic, kutcem@hotmail.com

Fournier's Gangrene is seen in healthy individuals (26% to 30%) and is more common in immunocompromised patients. It is frequently seen in elderly diabetic male patients with chronic alcohol use (16,17). The prognosis worsens in patients with multiple comorbidities (16).

ETIOLOGY OF FOURNIER'S GANGRENE;

Diabetes (glucosuric drugs used in diabetic patients), Malignant diseases, obesity, chronic alcohol drug addiction, and immunosuppressive chemotherapy are the most common conditions (3,17-20).

Despite advances all over the world in the treatment of sepsis due to the developed antibiotic therapy, the overall mortality rate for Fournier's Gangrene, unfortunately, remained stable at 40% (16,24,26). The reason for worrying death rates is the delay in diagnosis and treatment (8,17,26). The most critical factor affecting mortality is the time to surgery because early surgery can halve the mortality rate (8,26,27).

Although Fournier's Gangrene is an acute and rapidly progressing condition, it can progress slowly over days or weeks. Most studies have shown that the time from the onset of symptoms to hospital admission is approximately five days (28).

FOURNIER'S GANGRENE HAS FIVE STAGES:

Stage 1: Initial complaints include drowsiness, fatigue, and fever. These symptoms can be observed for 2 to 7 days.

Stage 2: The skin in the genital area is painful and edematous.

Stage 3: With increasing skin redness, genital pain, and tenderness are more pronounced.

Stage 4: Subcutaneous crepitation develops and appears dark.

Stage 5: Purulent discharge is observed. There is significant Gangrene in the involved area (29).

LABORATORY:

Evaluation should include a comprehensive metabolic panel (CMP) and a complete blood count (CBC) in suspected Fournier's Gangrene. The CBC will usually show a high white blood count (WBC) with the potential for a left shift. CMP may indicate concomitant renal failure and electrolyte abnormalities such as hyponatremia or metabolic acidosis.

High levels of lactate, c-reactive protein, and procalcitonin in the blood may be helpful in the evaluation of bacteremia and sepsis (30). Arterial blood gases are

used to assess oxygenation and acidosis, base deficit. Blood and wound cultures are required to plan antibiotic therapy. (31,32).

The Fournier Gangrene severity index (FGSI) was found to be inspired by the APACHE 2 score (33). This scoring system includes nine parameters, including vital signs and biochemistry. In most studies, the FGSI score was found to be significantly higher in the deceased group (34).

IMAGING:

The diagnosis of FG is based on clinical findings. In many cases, imaging is either unnecessary or not done because it can delay surgery. Surgical intervention should not be delayed in hemodynamically unstable patients (35).

Imaging can be used to determine the localization of the disease in asymptomatic and non-obvious cases. Simple radiography can show gas formation in soft tissue before the physical examination (36). Gas formation is present in most patients with FG and is specific to this disease (37). Ultrasonography helps detect subcutaneous gas or emphysema developing in the tissue. Gas in the scrotum is pathognomonic for Fournier's gangrene. A hazy appearance caused by infected subcutaneous tissue may appear as dirty shadowing (38).

Computed tomography is the most sensitive and specific imaging modality for FG diagnosis. Thickening in the affected area, fluid collection, subcutaneous emphysema, and abscess development are specific CT is also valuable for excluding perineal abscess, fistula formation, and various intra – and retroperitoneal disease processes (39). Magnetic resonance imaging is excellent for soft tissue imaging (40). Due to cost and time, its use is severely restricted (41).

TREATMENT / MANAGEMENT

When the diagnosis is sure, or the patient is not stable, emergency surgery should be performed immediately without waiting. Laboratory results should not be expected. Delaying intervention may result in the progression of a life-threatening infection. Fournier's Gangrene is a genuine surgical emergency. Since patients will usually be septic and in shock, managing the disease should be done with surgical and medical resuscitation (42). The first step in medical intervention is to start empirical broad-spectrum antibiotics without waiting for culture antibiogram results. Antibiotherapy should include triple therapy involving gram-positive, gram-negative, and anaerobic organisms where Fournier's Gangrene is most common. The most common microorganisms include staphylococci, streptococci, coliforms, Pseudomonas, Bacteroides, Clostridium, and yeast (43).

A combination of broad-generation cephalosporin, aminoglycoside, penicillin, and metronidazole is used as standard. Medical treatment is continued for at least two weeks(43).

HEMODYNAMIC RESUSCITATION AND PATIENT STABILIZATION

Fluid resuscitation plays an essential role in the treatment of FG patients. Patients with FG may present with hypotension and septic shock, and aggressive fluid resuscitation and hemodynamic support are required as this is associated with end-organ failure (44).

The patient's vital signs, urine output, and blood biochemistry should be closely monitored. Vasoactive agents such as norepinephrine should be started in patients with hypoperfusion to protect them from end-organ failure (45). If the patient's hypotension does not respond to fluid resuscitation, vasopressors may be added to the treatment. (43).

Electrolyte voids should be corrected with crystalloid fluids such as Lactated Ringer's solution (9,43,46,47). Diabetic patients with Fournier's Gangrene must correct their blood glucose abnormalities(15).

SURGICAL DEBRIDEMENT

As critical as these antibiotic and resuscitation measures are, these treatments should not delay the definitive and critical treatment for Fournier's Gangrene: urgent, early, and aggressive surgical exploration and debridement. Delaying the surgery will increase patient morbidity and mortality (48,26,49). The principle of surgical debridement is based on resectioning all dead, infected, and necrotic tissue. Removal of unhealthy tissue may be necessary. In this way, premature closure of the wound is prevented before the underlying tissues heal sufficiently. The removal of inanimate tissues in the first intervention is considered the most critical factor in the patient's survival. Ventilation of living tissues by fenestration is recommended in most studies. Close monitoring of the wound and repeated surgical debridements are necessary to control infection. Extensive debridement in FG surgery results in extensive tissue loss at the site. The choice of flap or graft in surgical reconstruction depends on the defect's location and the local tissue's presence (51).

HYPERBARIC OXYGEN THERAPY

Hyperbaric oxygen therapy is a beneficial option in the postoperative period. Occlusion developing in the vascular structure causes necrosis by disrupting tissue

nutrition. Therefore, oxygenation of the tissues will help the treatment. Oxygen therapy stimulates the immune system by increasing fibroblast proliferation and neutrophil functions and accelerating the passage of antibiotics into the cell, which accelerates wound healing (52).

VACUUM ASSISTED CLOSURE

Vacuum-assisted closure (VAC) method accelerates wound healing by reducing edema and increasing blood flow. This system increases angiogenesis and accelerates tissue nutrition and healing. Thanks to this system is the primary mechanism of the system to drain the infected fluid and debris (53).

TREATMENT SUMMARY

- Success in the treatment of Fournier's Gangrene is early diagnosis and surgical debridement.

- Hemodynamic resuscitation and broad-spectrum antibiotics should be added to the treatment.

- Early surgical intervention is essential for survival, imaging and laboratory tests should not delay intervention in critical cases.

- Postop debrided areas should be treated with sterile dressings or vacuum wound pressure systems.

- As the vascular structure of the testicles is not affected, it is usually preserved.

- If there is urethral involvement, a suprapubic catheter should be placed instead of the urethral catheter.

- If the rectum or anus is affected, a temporary colostomy may be required.

- Hyperbaric oxygen therapy can help reduce morbidity and mortality.

Reconstructive surgery should be performed when the debrided wound is completely healed. (35).

REFERENCES

1. Corman ML. Jean-Alfred Fournier 1832-1914. Gangrène foudroyante de la verge (overwhelming gangrene). *Sem Med* 1883. *Diseases of the colon and rectum [Internet]*. 1988 [cited 2022 Jan 12];31(12):984–8. Available from:
- 2 – Mishra SP, Singh S, Gupta SK. Necrotizing Soft Infections: A Surgeon's Prospect. *Int J Enflam*.2013;2013 – 609628
3. Thwaini A, Khan A, Malik A, Cherian J, et al. Fournier's gangrene and emergency management. *Postgrad Med J*. 2006 Agosto; 82 (970):516-9.
4. Voelzke BB, Hagedorn JC. Presentation and Diagnosis of Fournier's Gangrene. *Urology*. 2018

Nis; 114 :8-13

5. Canbaz H, Caglikulekci M, Altun U, et al. Fournier's Gangrene: analysis of risk factors affecting the prognosis and cost of therapy in 18 cases. *Ulus Travma Acil Cerrahi Derg* 2010;16:71-6.
6. Oymaci E, Coskun A, Yakan S, et al. Evaluation of factors affecting mortality in Fournier's Gangrene: Retrospective clinical study of sixteen cases. *Ulusal Cer Derg* 2014;30:85-9.
7. Czymek R, Schmidt A, Eckmann C, Bouchard R, Wulff B, Laubert T, et al. Fournier's Gangrene: vacuum-assisted closure versus conventional dressings. *Am J Surg* 2009;197:168-76.
8. Bjurlin MA, O'Grady T, Kim DY, et al. Causative pathogens, antibiotic sensitivity, resistance patterns, and severity in a contemporary series of Fournier's Gangrene. *Urology*. 2013 Apr;81(4):752-8.
9. Auerbach J, Bornstein K, Ramzy M, et al . Fournier Gangrene in the Emergency Department: Diagnostic Dilemmas, Treatments and Current Perspectives. *Open Access Emerg Med*. 2020;12:353-364.
10. H. J. Jeong, S. C. Park, I. Y. Seo, et al. "Prognostic factors in Fournier gangrene," *Int J Urol*, vol. 12, no. 12, pp. 1041-1044, Dec. 2005, doi: 10.1111/j.1442-2042.2005.01204.x.
11. A. Kiliç, Y. Aksoy, L. Kiliç, "Fournier's gangrene: etiology, treatment, and complications," *Ann Plast Surg*, vol. 47, no. 5, pp. 523-527, Nov. 2001, doi: 10.1097/0000637-200111000-00009.
12. Erdogan A, Aydogan I, Senol K, et al. Simple scoring system for prediction of mortality in Fournier's Gangrene. *Eur J Trauma Emerg Surg* 2016;42:513-8.
13. Smith GL, Bunker CB, Dinneen MD. Fournier's gangrene. *Br J Urol* 1998;81:347-55.
14. Korkut M, Icoz G, Dayangac M, et al. Fournier's gangrene Outcome analysis in patients with Fournier's gangrene: report of 45 cases. *Dis Colon Rectum* 2003;46:649-52.
15. Mallikarjuna MN, Vijayakumar A, Patil VS, et al . Fournier's gangrene: Current Practices. *ISRN Surg* 2012;2012:937-42.
16. Joury A, Mahendra A, Alshehri M, Downing A. Extensive necrotizing fasciitis from Fournier's Gangrene. *Urol Case Rep*. 2019 Sep;26:100943. [PMC free article]
17. Sorensen MD, Krieger JN, Rivara FP, et al. Fournier's Gangrene: population based epidemiology and outcomes. *J Urol*. 2009 May;181(5):2120-6. [PMC free article]
18. Elem B, Ranjan P. Impact of immunodeficiency virus (HIV) on Fournier's Gangrene: observations in Zambia. *Ann R Coll Surg Engl*. 1995 Jul;77(4):283-6.
19. Moussa M, Abou Chakra M. Isolated Penile Fournier's gangrene: A case report and literature review. *Int J Surg Case Rep*. 2019;62:65-68.
20. Mouraviev VB, Pautler SE, Hayman WP. Fournier's Gangrene following penile self-injection with cocaine. *Scand J Urol Nephrol*. 2002;36(4):317-8.
21. Bloomgarden Z, Einhorn D, Grunberger G, et al. Fournier's Gangrene and sodium-glucose cotransporter 2 inhibitors: Is there a causal association? *J Diabetes*. 2019 May;11(5):340-341. [PubMed]
22. Perkins TA, Bieniek JM, Sumfest JM. Solitary Candida albicans Infection Causing Fournier Gangrene and Review of Fungal Etiologies. *Rev Urol*. 2014;16(2):95-8.
23. Serrano Olave A, Bueno Moral AI, Martínez Bañón C, et al . Fournier's Gangrene under Sodium-Glucose Cotransporter-2 Inhibitors Therapy in Gynecological Patients. *Int J Environ Res Public Health*. 2022 May 21;19(10)
24. Joury A, Mahendra A, Alshehri M, Downing A. Extensive necrotizing fasciitis from Fournier's Gangrene. *Urol Case Rep*. 2019 Sep;26:100943.

24. Rudd KE, Johnson SC, Agesa KM, et al. Global, regional, and national sepsis incidence and mortality, 1990-2017: analysis for the Global Burden of Disease Study. *Lancet*. 2020 Jan 18;395(10219):200-211.
25. Radcliffe RS, Khan MA. Mortality associated with Fournier's Gangrene remains unchanged over 25 years. *BJU Int*. 2020 Apr;125(4):610-616.
26. Peetermans M, de Prost N, Eckmann C, et al. Necrotizing skin and soft-tissue infections in the intensive care unit. *Clin Microbiol Infect*. 2020 Jan;26(1):8-17.
27. Sugihara T, Yasunaga H, Horiguchi H, et al. Impact of surgical intervention timing on the case fatality rate for Fournier's Gangrene: an analysis of 379 cases. *BJU Int*. 2012 Dec;110(11 Pt C):E1096-100.
28. Ferreira PC, Reis JC, Amarante JM, et al. Fournier's gangrene: a review of 43 reconstructive cases. *Plast Reconstr Surg*. 2007 Jan;119(1):175-184.
29. Talwar A, Puri N, Singh M. Fournier's Gangrene of the Penis: A Rare Entity. *J Cutan Aesthet Surg*. 2010 Jan;3(1):41-4.
30. Fan SL, Miller NS, Lee J, et al. Diagnosing sepsis – The role of laboratory medicine. *Clin Chim Acta*. 2016 Sep 01;460:203-10.
31. Verma S, Sayana A, Kala S, et al. Evaluation of the Utility of the Fournier's Gangrene Severity Index in the Management of Fournier's Gangrene in North India: A Multicentre Retrospective Study. *J Cutan Aesthet Surg*. 2012 Oct;5(4):273-6.
32. Eke N. Fournier's gangrene: a review of 1726 cases. *Br J Surg*. 2000 Jun;87(6):718-28.
33. E. Laor, L. S. Palmer, B. M. Tolia, et al, "Outcome prediction in patients with Fournier's gangrene," *J Urol*, vol. 154, no. 1, pp. 89–92, Jul. 1995.
34. T.-Y. Lin et al., "Validation and simplification of Fournier's gangrene severity index," *Int J Urol*, vol. 21, no. 7, pp. 696–701, Jul. 2014, doi: 10.1111/iju.12426.
35. Chennamsetty A, Khourdaji I, Burks F, et al. Contemporary diagnosis and management of Fournier's Gangrene. *Ther Adv Urol*. 2015 Aug;7(4):203-15.
36. A. Chennamsetty, I. Khourdaji, F. Burks, et al. "Contemporary diagnosis and management of Fournier's gangrene," *Ther Adv Urol*, vol. 7, no. 4, pp. 203–215, Aug. 2015, doi: 10.1177/1756287215584740.
37. E. P. Misiakos, G. Bagias, P. Patapis, et al. "Current concepts in the management of necrotizing fasciitis," *Front Surg*, vol. 1, p. 36, 2014, doi: 10.3389/fsurg.2014.00036
38. N. Gupta, K. M. Zinn, I. Bansal, et al. "Fournier's gangrene: ultrasound or computed tomography?," *Med Ultrason*. vol. 16, no. 4, pp. 389–390, Dec. 2014.
39. R. B. Levenson, A. K. Singh, R. A. Novelline, "Fournier gangrene: role of imaging," *Radiographics*, vol. 28, no. 2, pp. 519–528, Apr. 2008, doi: 10.1148/rg.282075048
40. K.-T. Kim et al., "Can necrotizing infectious fasciitis be differentiated from nonnecrotizing infectious fasciitis with MR imaging?," *Radiology*. vol. 259, no. 3, pp. 816–824, Jun. 2011, doi: 10.1148/radiol.11101164.
41. B. B. Voelzke and J. C. Hagedorn, "Presentation and Diagnosis of Fournier Gangrene," *Urology*. vol. 114, pp. 8–13, Apr. 2018, doi: 10.1016/j.urology.2017.10.031.
42. Chernyadyev SA, Ufimtseva MA, Vishnevskaya IF, et al. Fournier's Gangrene: Literature Review and Clinical Cases. *Urol Int*. 2018;101(1):91-97.
43. Hakkarainen TW, Kopari NM, Pham TN, et al. Necrotizing soft tissue infections: review and current concepts in treatment, systems of care, and outcomes. *Curr Probl Surg*. 2014

Aug;51(8):344-62.

44. E. B. Benjelloun et al., "Fournier's gangrene: our experience with 50 patients and analysis of factors affecting mortality," *World J Emerg Surg*, vol. 8, no. 1, p. 13, Apr. 2013, doi: 10.1186/1749-7922-8-13
45. A. Rhodes et al., "Surviving Sepsis Campaign: International Guidelines for Management of Sepsis and Septic Shock: 2016," *Intensive Care Med*, vol. 43, no. 3, pp. 304–377, Mar. 2017, doi: 10.1007/s00134-017-4683-6.
46. Toporek AH, Semler MW, Self WH, et al, SMART Investigators and the Pragmatic Critical Care Research Group. Balanced Crystalloids versus Saline in Critically Ill Adults with Hyperkalemia or Acute Kidney Injury: Secondary Analysis of a Clinical Trial. *Am J Respir Crit Care Med*. 2021 May]
47. Benjelloun el B, Souiki T, Yakla N, et al. Fournier's Gangrene: our experience with 50 patients and analysis of factors affecting mortality. *World J Emerg Surg*. 2013 Apr 01;8(1):13.
48. Wong CH, Chang HC, Pasupathy S, et al. Necrotizing fasciitis: clinical presentation, microbiology, and determinants of mortality. *J Bone Joint Surg Am*. 2003 Aug;85(8):1454-60.
49. Kabay S, Yucel M, Yaylak F et al. The clinical features of Fournier's Gangrene and the predictivity of the Fournier's Gangrene Severity Index on the outcomes. *Int Urol Nephrol*. 2008;40(4):997-1004.
50. A. Thwaini et al., "Fournier's gangrene and its emergency management," *Postgrad Med J*, vol. 82, no. 970, pp. 516–519, Aug. 2006, doi: 10.1136/pgmj.2005.042069.
51. P.C.Ferreira et al., "Fournier's gangrene: a review of 43 reconstructive cases," *Plast Reconstr Surg*, vol. 119, no. 1, pp. 175–184, Jan. 2007, doi: 10.1097/01.prs.0000244925.80290.57.
52. A. Janane et al., "[Hyperbaric oxygen therapy adjunctive to surgical debridement in management of Fournier's gangrene: usefulness of a severity index score in predicting disease gravity and patient survival]," *Actas Urol Esp*, vol. 35, no. 6, pp. 332–338, Jun. 2011, doi: 10.1016/j.acuro.2011.01.019.
53. Hong KS, Yi HJ, Lee RA, et al. Prognostic factors and treatment outcomes for patients with Fournier's Gangrene: a retrospective study. *Int Wound J*. 2017 Dec;14(6) :1352-1358.