



MELENEY'S SYNERGISTIC GANGRENE: A CASE SERIES OF FOUR PATIENTS WITH HETEROGENEOUS PRESENTATIONS AND SURGICAL OUTCOMES FROM A TERTIARY CARE CENTRE

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ABSTRACT

Background: Meleney's synergistic gangrene (MSG) is a rare necrotizing soft tissue infection characterized by progressive subcutaneous necrosis, severe pain, and delayed systemic toxicity. Its indolent onset and polymicrobial etiology frequently result in delayed diagnosis and poor outcomes. **Methods:** Over a 12-month period, four patients with MSG involving different anatomical sites were managed at a tertiary care hospital. Clinical features, comorbidities, microbiology, surgical interventions, and outcomes were prospectively analyzed. **Results:** Three patients survived following early resuscitation, broad-spectrum antibiotics, serial surgical debridement, negative pressure wound therapy, and delayed reconstruction. One immunocompromised postoperative patient succumbed to multiorgan failure despite aggressive management. **Conclusion:** MSG requires a high index of suspicion, especially in postoperative or high-risk patients presenting with disproportionate pain and progressive skin changes. Early debridement and multidisciplinary care remain the cornerstones of management. Delayed diagnosis and immunosuppression are associated with poor outcomes.

KEYWORDS : Meleney's Gangrene; Necrotizing Soft Tissue Infection; Surgical Debridement; Negative Pressure Wound Therapy; Postoperative Infection

INTRODUCTION

Meleney's synergistic gangrene (MSG), first described by Meleney in 1924, is an uncommon variant of necrotizing soft tissue infection caused by synergistic interaction between aerobic and anaerobic organisms, classically *Staphylococcus aureus* and microaerophilic streptococci [16–18]. Unlike necrotizing fasciitis, MSG demonstrates a deceptively slow radial spread with extensive subcutaneous undermining and delayed systemic toxicity, often leading to diagnostic delay [9,20].

MSG typically follows surgical procedures, minor trauma, or skin breakdown, particularly in patients with diabetes mellitus, malnutrition, tuberculosis, malignancy, or immunosuppression [7,23]. Clinically, it presents with severe pain out of proportion to cutaneous findings, violaceous margins, foul discharge, and progressive ulceration [1,25]. Mortality ranges from 10–40%, particularly when surgical intervention is delayed [4,15].

We present four cases of MSG with diverse anatomical involvement and host factors, highlighting diagnostic challenges, management strategies, and outcomes.

Methods

Between January 2024 and January 2025, four patients with clinically diagnosed MSG were treated at our institution. Diagnosis was based on clinical features supported by imaging and intraoperative findings. Empirical broad-spectrum antibiotics were initiated immediately and later tailored to culture results. Serial surgical debridement was performed until viable tissue margins were achieved. Negative pressure wound therapy (NPWT) and reconstructive procedures were employed as indicated. Outcomes assessed included survival, need for reconstruction, and recurrence within 90 days.

Case Summaries

Case 1

A 48-year-old male on antitubercular therapy developed progressive necrosis at a postoperative colostomy closure site. Despite emergency debridement and escalation of antibiotics, he developed septic shock and multiorgan failure, succumbing on postoperative day 8. This case highlights the adverse impact of immunosuppression and delayed recognition in postoperative MSG [8,22].



Fig 1- Showing Violaceous Discoloration, Foul-Smelling Discharge, and Progressive Necrosis, Extending to the Perineum and Proximal Left Thigh



Fig 2- Showing Wound Dehiscence Along-With Discoloration and Foul-Smelling Discharge and Progressive Necrosis Along the Closure Site

Case 2

A 55-year-old chronic smoker presented with suprapubic and

bilateral scrotal involvement. Early aggressive debridement, serial wound exploration, NPWT, and delayed skin grafting resulted in complete recovery without testicular loss. Smoking-related tissue ischemia likely contributed to disease severity [21,26].



Fig 3- Extensive Erythema, Crepitus, Foul-Smelling Purulent Discharge, and Blackish Necrotic Patches Over the Bilateral Scrotum With Exposed Testicles



Fig 4- Post Debridement Picture of Bilateral Inguinal Area with Still Patches of Purulent Areas

Case 3

A 32-year-old immunocompetent woman presented with early MSG involving the mons pubis and vulva. Imaging excluded fascial involvement. Conservative management with targeted antibiotics and local wound care led to complete healing without surgical debridement, demonstrating that selected early cases may be managed non-operatively [5,10].



Fig 5- Involvement of Mons Pubis and Vulva with Focal Areas of Superficial Necrosis

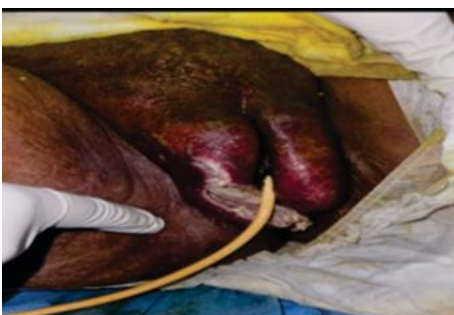


Fig 6- Superficial Necrotic Patches on Vulva and Lower Abdomen

Case 4

A 58-year-old poorly controlled diabetic female developed extensive perineal and thigh involvement. Multiple debridement, NPWT, and delayed reconstruction using split-thickness skin grafts and local flaps resulted in good functional recovery. Diabetes and delayed presentation were major contributors to disease extent [1,4,23].



Fig 8- Extensive Tissue Involvement of Right Sided Perineum Extending Into Medial Aspect of Thigh.



Fig 9- Extensive Tissue Necrosis Involving Almost Whole Thigh

DISCUSSION

MSG remains a diagnostic challenge due to its rarity and indolent early course. Severe pain disproportionate to physical findings, progressive undermining, and polymicrobial infection are key clinical clues [20,25]. Unlike necrotizing fasciitis, early systemic toxicity may be absent, leading to delayed intervention.

Our series demonstrates wide variability in host factors and disease severity. Immunosuppression, diabetes, smoking, and delayed referral were associated with extensive tissue loss and poorer outcomes. Early and repeated surgical debridement remains the most important determinant of survival [4,15]. Clindamycin plays a critical adjunctive role due to its antitoxin effects in streptococcal infections [24].

NPWT facilitated wound healing and simplified later reconstruction in survivors, supporting its role as an adjunct rather than a substitute for surgery [2,19]. Early involvement of plastic surgeons improved functional and cosmetic outcomes.

CONCLUSION

Meleney's synergistic gangrene is a rare but potentially fatal necrotizing infection requiring early recognition and aggressive multidisciplinary management. A high index of suspicion, prompt surgical debridement, broad-spectrum antibiotics, and meticulous supportive care are essential to improve survival and reduce morbidity.

REFERENCES

1. Anaya DA, Dellinger EP. Necrotizing soft-tissue infection: diagnosis and management. Clin Infect Dis. 2007;44(5):705-710.
2. Apelqvist J, Willy C, Fagerdahl AM, et al. Negative pressure wound therapy-overview, challenges and perspectives. J Wound Care.

- 2017;26(Sup3):S1–S43.
3. Brook I. Microbiology and management of soft tissue and muscle infections. *Int J Surg*. 2008;6(4):328–338.
4. Elliott D, Kufera JA, Myers RA. Necrotizing soft tissue infections: risk factors for mortality and strategies for management. *Ann Surg*. 1996;224(5):672–683.
5. Elliott D, Kufera JA, Myers RA. The microbiology of necrotizing soft tissue infections. *Am J Surg*. 2000;179(5):361–366.
6. Giuliano A, Lewis F Jr, Hadley K, Blaisdell FW. Bacteriology of necrotizing fasciitis. *Am J Surg*. 1977;134(1):52–57.
7. Goh T, Goh LG, Ang CH, Wong CH. Early diagnosis of necrotizing fasciitis. *Br J Surg*. 2014;101(1):e119–e125.
8. Gupta RK, Gupta S, Singh G. Adverse effects of antitubercular treatment: an Indian perspective. *Trop Doct*. 2011;41(4):211–213.
9. Hakkarainen TW, Kopari NM, Pham TN, Evans HL. Necrotizing soft tissue infections: review and current concepts in treatment, systems of care, and outcomes. *Curr Probl Surg*. 2014;51(8):344–362.
10. Headley AJ. Necrotizing soft tissue infections: A primary care review. *Am Fam Physician*. 2003;68(2):323–328.
11. Hersh RE, Bhama AR, Shakir A, et al. Vacuum-assisted closure therapy for the treatment of complex wounds in patients with necrotizing fasciitis. *Ann Plast Surg*. 2013;71(4):394–401.
12. Hsiao CT, Weng HH, Yuan YD, Chen CT, Chen IC. Predictors of mortality in patients with necrotizing fasciitis. *Am J Emerg Med*. 2008;26(2):170–175.
13. Lin JN, Chang LL, Lai CH, Chen YH, Tsai SS, Lin HH. Necrotizing fasciitis caused by *Staphylococcus aureus* and *Bacteroides fragilis*: a case report and review. *Kaohsiung J Med Sci*. 2011;27(9):410–414.
14. May AK, Stafford RE, Bulger EM, et al. Treatment of complicated skin and soft tissue infections. *Surg Infect (Larchmt)*. 2009;10(5):467–499.
15. McHenry CR, Piotrowski JJ, Petrinic D, Malangoni MA. Determinants of mortality for necrotizing soft-tissue infections. *Ann Surg*. 1995;221(5):558–563.
16. Meleney FL. Hemolytic streptococcus gangrene. *Arch Surg*. 1924;9(2):317–364.
17. Meleney FL. Bacterial synergism in disease processes: a study of mixed infections in war wounds. *Ann Surg*. 1931;94(5):961–981.
18. Meleney FL. Bacteriology of synergistic infections in the skin and subcutaneous tissues. *Surg Gynecol Obstet*. 1931;53:145–162.
19. Orgill DP, Bayer LR. Negative pressure wound therapy: past, present and future. *Int Wound J*. 2013;10 Suppl 1:15–19.
20. Sarani B, Strong M, Pascual J, Schwab CW. Necrotizing fasciitis: current concepts and review of the literature. *J Am Coll Surg*. 2009;208(2):279–288.
21. Soperi M. Effects of cigarette smoke on the immune system. *Nat Rev Immunol*. 2002;2(5):372–377.
22. Song JY, Park CW, Jeon JH, et al. Necrotizing fasciitis in patients with tuberculosis. *Scand J Infect Dis*. 2005;37(12):918–921.
23. Stevens DL, Bryant AE. Necrotizing soft-tissue infections. *N Engl J Med*. 2017;377(23):2253–2265.
24. Stevens DL, Ma Y, Salmi DB, McIndoo E, Wallace RJ, Bryant AE. Impact of antibiotics on expression of virulence-associated exotoxin genes in *Streptococcus pyogenes*. *J Infect Dis*. 2007;195(5):684–693.
25. Ulu-Kilic A, Karakas A, Erdem H, Turker T. Meleney's gangrene: a forgotten and overlooked diagnosis. *J Infect Dev Ctries*. 2015;9(9):1041–1044.
26. Guidelines for the diagnosis and management of skin and soft-tissue infections. *Clin Infect Dis*. 2005;41(10):1373–406.