

BIBLIOGRAPHIC INFORMATION SYSTEM

JOURNAL FULL TITLE: Journal of Biomedical Research & Environmental Sciences

ABBREVIATION (NLM): J Biomed Res Environ Sci **ISSN:** 2766-2276 **WEBSITE:** <https://www.jelsciences.com>

SCOPE & COVERAGE

- ▶ **Sections Covered:** 34 specialized sections spanning 143 topics across Medicine, Biology, Environmental Sciences, and General Science
- ▶ Ensures broad interdisciplinary visibility for high-impact research.

PUBLICATION FEATURES

- ▶ **Review Process:** Double-blind peer review ensuring transparency and quality
- ▶ **Time to Publication:** Rapid 21-day review-to-publication cycle
- ▶ **Frequency:** Published monthly
- ▶ **Plagiarism Screening:** All submissions checked with iThenticate

INDEXING & RECOGNITION

- ▶ **Indexed in:** [Google Scholar](#), IndexCopernicus (**ICV 2022: 88.03**)
- ▶ **DOI:** Registered with CrossRef (**10.37871**) for long-term discoverability
- ▶ **Visibility:** Articles accessible worldwide across universities, research institutions, and libraries

OPEN ACCESS POLICY

- ▶ Fully Open Access journal under Creative Commons Attribution 4.0 License (CC BY 4.0)
- ▶ Free, unrestricted access to all articles globally

GLOBAL ENGAGEMENT

- ▶ **Research Reach:** Welcomes contributions worldwide
- ▶ **Managing Entity:** SciRes Literature LLC, USA
- ▶ **Language of Publication:** English

SUBMISSION DETAILS

- ▶ Manuscripts in Word (.doc/.docx) format accepted

SUBMISSION OPTIONS

- ▶ **Online:** <https://www.jelsciences.com/submit-your-paper.php>
- ▶ **Email:** support@jelsciences.com, support@jbresonline.com

[HOME](#)[ABOUT](#)[ARCHIVE](#)[SUBMIT MANUSCRIPT](#)[APC](#)

 **Vision:** The Journal of Biomedical Research & Environmental Sciences (JBRES) is dedicated to advancing science and technology by providing a global platform for innovation, knowledge exchange, and collaboration. Our vision is to empower researchers and scientists worldwide, offering equal opportunities to share ideas, expand careers, and contribute to discoveries that shape a healthier, sustainable future for humanity.

CASE REPORT

Cutaneous Penile Metastases from Prostate Adenocarcinoma: A Rare Case Report

Navan van Jaarsveld^{1*}, Litha Chimusoro¹, Chiara Ilett¹ and Jeff John^{1,2}

¹Division of Urology, Department of Surgery, Frere Hospital and Walter Sisulu University, East London, South Africa

²Division of Urology, Department of Surgery, Groote Schuur Hospital and University of Cape Town, Cape Town, South Africa

Abstract

Introduction: While prostate cancer is the most common malignancy in men, cutaneous penile metastases are rare and indicate a poor prognosis.

Case Presentation: A 63-year-old man with painful non-healing penile lesions was referred to our unit for workup. Clinical examination revealed indurated ulcerative lesions with bilateral inguinal lymphadenopathy. Incisional biopsy of the penile lesions confirmed metastatic adenocarcinoma with NKX3.1 positivity, indicating prostatic origin. Further workup showed a blood serum PSA level of 489 ng/mL; prostate biopsy confirmed Gleason Grade 8 prostate cancer, and bone metastases were seen on plain skeletal radiography. A bilateral orchidectomy was performed, and he was started on docetaxel chemotherapy and zoledronic acid.

Conclusion: The multidisciplinary management of this case resulted in regression of the metastatic lesions at his six-month follow-up, with no new cutaneous lesions observed.

Keynote message: Cutaneous penile metastases from prostate adenocarcinoma are a rare manifestation of advanced disease and are linked with a poor prognosis. This case highlights the importance of careful clinical evaluation of persistent penile lesions, as early recognition and multidisciplinary management are essential for oncological treatment and symptom control.

*Corresponding author(s)

Navan van Jaarsveld, Division of Urology, Department of Surgery, Frere Hospital and Walter Sisulu University, East London, South Africa

ORCID: 0000-0003-1039-1562

Email: navanvanj@gmail.com

DOI: 10.37871/jbres2314

Submitted: 15 July 2026

Accepted: 23 July 2026

Published: 24 July 2026

Copyright: © 2026 van Jaarsveld N, et al. Distributed under Creative Commons CC-BY 4.0

OPEN ACCESS

Keywords

- Prostate cancer
- Dermal lesions
- Advanced malignancy
- Oncology

Introduction

Prostate cancer (PCa) is the commonest malignancy in men and a leading cause of cancer-related mortality worldwide [1]. It typically metastasizes to the bones, lymph nodes, liver, and lungs. Penile metastases from PCa are an exceedingly rare occurrence

VOLUME: 7 ISSUE: 7 - JULY, 2026



How to cite this article: van Jaarsveld N, Chimusoro L, Ilett C, John J. Cutaneous Penile Metastases from Prostate Adenocarcinoma: A Rare Case Report. J Biomed Res Environ Sci. 2026 July 24; 7(7): 5. Doi: 10.37872/jbres2314

of secondary involvement, with an incidence of less than 0.5%, often signifying advanced disease and a poor prognosis. Owing to their nonspecific appearance, these lesions often mimic benign inflammatory or infectious penile conditions, causing delayed diagnosis and treatment [2,3]. We report a rare case of cutaneous penile metastases secondary to prostate adenocarcinoma and discuss the clinical implications.

Case Presentation

A 63-year-old male was referred to our urology unit with an eight-month history of painful, non-healing penile lesions that responded poorly to antimicrobial therapy. On further history, he reported no Lower Urinary Tract Symptoms (LUTS), urethral discharge, gross hematuria, or any constitutional symptoms. He had no comorbidities, history of Sexually Transmitted Infections (STIs), or family history of genitourinary malignancies.

Clinical examination revealed multiple indurated ulcerative lesions, measuring 2cm x 3cm in diameter, involving the penile glans and external urethral meatus (Figure 1). Hard bilateral inguinal lymph nodes were palpable, and no urethral discharge was noted. The scrotum was uninvolved, and normal testes were palpable bilaterally. Abdominal and neurological examination were unremarkable. An incisional biopsy of the penile lesions was performed, and microscopic examination demonstrated dermal infiltration by a malignant gland-forming epithelial neoplasm composed of irregular glands, poorly formed acinar structures, and tumour cells with enlarged, hyperchromatic nuclei and moderate eosinophilic cytoplasm. Histopathology revealed ulcerated skin infiltrated and undermined by an invasive moderately differentiated adenocarcinoma with MNKX3.1-positive immunohistochemical staining, supporting a prostatic origin for the cutaneous penile metastatic deposits. Based on

this result, a prostate cancer workup was initiated. Digital rectal examination revealed a hard, irregular prostate consistent with a clinical T4 lesion with markedly elevated serum Prostate-Specific Antigen (PSA) levels of 489 ng/mL. A skeletal survey via plain radiography showed extensive sclerotic bony lesions involving the vertebral column and pelvic bones, consistent with spinal bone metastases (Figure 2), despite the patient being asymptomatic. A transrectal ultrasound-guided prostate biopsy confirmed the diagnosis of invasive adenocarcinoma

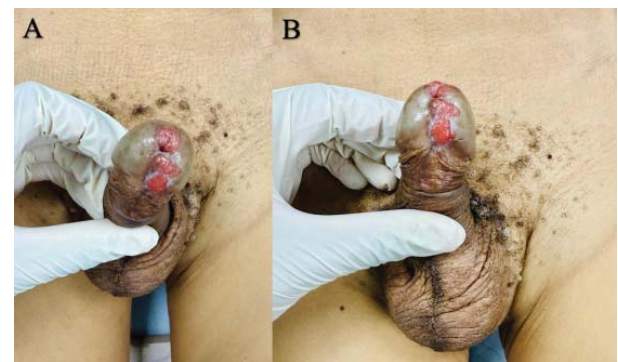


Figure 1 Superior (A) and anterior frontal (B) views of the penis in the supine position show multiple irregular, exophytic, erythematous ulcerative nodules on the dorsal aspect of the glans penis extending into the external urethral meatus.



Figure 2 Anteroposterior pelvic (A) and lateral lumbar spinal (B) view radiographs demonstrating diffuse osteoblastic metastases involving the ilium, sacrum, pubic rami, proximal femur, L1-L5 vertebrae, and the sacrum. The bone trabeculae are obliterated by patchy, sclerotic areas, resulting in a “mottled ivory pelvis” appearance. The lateral thoracolumbar radiograph (C) shows multilevel sclerotic vertebral lesions extending from the mid-thoracic region to the lumbar vertebrae.



of the prostate, Gleason Grade 8 (4+4), with no evidence of perineural or lymphovascular invasion. A multidisciplinary (MDT) meeting ensued, and the patient decided to undergo a bilateral orchidectomy, with consensus to proceed with surgical castration. Following the orchidectomy, he was referred to our oncology unit, where docetaxel chemotherapy and zoledronic acid were initiated. At his most recent follow-up visit, six months later, the penile metastatic lesions regressed, pain improved, partial epithelial healing occurred, and no new cutaneous lesions were noted.

Discussion

Penile metastases from prostate cancer carries a poor prognosis, with a mean survival reported at less than one year after diagnosis [4]. It can present with nodules, ulceration, penile pain, induration, priapism, hematuria, obstructive or irritative urinary symptoms [2]. Unlike the typical metastatic pattern of prostate cancer, which is predominantly skeletal and nodal [5], penile metastases represent an atypical cutaneous manifestation that often leads to diagnostic delay [6].

We present a rare case of a metastatic adenocarcinoma of the prostate presenting as cutaneous lesions of the penile glans and shaft. This phenomenon is believed to result from systemic venolymphatic dissemination, either via retrograde venous spread through the valveless pelvic venous plexus or via lymphatic spread through pelvic and inguinal lymphatics [7]. In our case, the presence of bilateral inguinal lymphadenopathy and skeletal metastases supports a systemic venolymphatic route of spread, consistent with mechanisms described in prior case reports [2,4,8]. Furthermore, the diagnostic challenge was highlighted by a longstanding history of painful penile lesions initially attributed to primary penile pathology. This is echoed in the literature describing misdiagnosed penile lesions as Peyronie's

disease, balanitis, Sexually Transmitted Diseases (STIs), or primary penile carcinoma [4]. Definitive diagnosis was achieved through an incisional penile biopsy and immunohistochemical staining, underscoring the importance of histopathological confirmation when evaluating persistent penile lesions. In our patient, NKX3.1 immunohistochemistry demonstrated nuclear positivity, confirming prostate origin. NKX3.1 is a highly sensitive and specific marker for prostatic tissue, shown to be superior to serum PSA, especially in poorly differentiated metastatic adenocarcinoma [9]. The elevated serum PSA (489 ng/mL) supported the diagnosis and reflected the burden of metastatic disease. A limitation of this case is that contrast-enhanced computed tomography (CT) and prostate-specific membrane antigen positron emission tomography/computed tomography (PSMA PET/CT) were not performed, as advanced imaging was unlikely to alter the initial management.

Most reported cases of penile metastases from prostate adenocarcinoma occur after an established diagnosis of prostate cancer, often during disease progression or biochemical relapse [2,4,10]. In contrast, the present case is notable as the cutaneous penile metastases were the presenting manifestation, preceding the diagnosis of prostate cancer. Similar presentations have been described, with authors reporting diagnostic delays prior to histological confirmation [4,7,8].

Although Androgen Deprivation Therapy (ADT) remains the backbone of treatment for metastatic prostate cancer, it is now combined with hormone therapies, taxane-based chemotherapy, or radiotherapy [11,12]. Given the advanced, hormone-sensitive nature of the disease [13], all management options were discussed with the patient. Considering his residence in a remote rural area and the anticipated challenges with long-term follow-up, a bilateral orchidectomy was performed as definitive androgen deprivation therapy. Surgical castration remains an effective, rapid,



and cost-efficient approach for reducing circulating testosterone and slowing disease progression [14–16].

This case illustrates the challenges associated with the delayed diagnosis due to atypical presentation and emphasizes the need for clinicians to maintain a broad differential diagnosis for persistent penile lesions in older men. Furthermore, if it were assumed that the lesions were a primary penile pathology based on their appearance, a crucial diagnosis might have been overlooked in a patient presenting with similar skin nodules but no other symptoms.

The significant psychological impact of penile metastases and bilateral orchidectomy, including effects on sexual function and body image, further emphasizes the importance of counseling and psychosocial support. Multidisciplinary care remains vital to ensure comprehensive treatment, symptom control, and psychological support [17,18].

Conclusion

Cutaneous penile metastases from prostate adenocarcinoma are a rare phenomenon, but a clinically significant manifestation of late-stage disease. Clinicians should maintain a high index of suspicion when evaluating persistent penile lesions, especially in patients with other systemic symptoms, and pursue prompt histopathological evaluation to avoid delayed diagnosis of an underlying malignancy.

Acknowledgments

None

Author Contributions

Van Jaarsveld Navan: Writing – original draft; writing – review and editing; validation

Chimusoro Litha: Writing – original draft; writing – review and editing; validation

Ilett Chiara: Writing – original draft; writing

– review and editing; validation

Jeff John: Supervision; writing – original draft; writing – review and editing; validation

Ethics Approval and Consent to Participate

Our institution does not require ethical approval for individual case reports.

Consent for Publication

Written informed consent was obtained from the patient for anonymized information and the accompanying images to be published in this article.

Funding

The authors received no financial support for the research, authorship, or publication of this article.

Competing Interests

The authors declare no potential conflicts of interest concerning the research, authorship, or publication of this manuscript.

Availability of data and materials

Not applicable.

References

1. Bray F, Laversanne M, Sung H, Ferlay J, Siegel RL, Soerjomataram I, et al. Global cancer statistics 2022: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin.* 2024;74(3):229-63.
2. Cherian J, Rajan S, Thwaini A, Elmasry Y, Shah T, Puri R. Secondary penile tumours revisited. *Int Semin Surg Oncol.* 2006;3:33.
3. Martz N, Benziane-Ouaritini N, Gautier M, Brenot-Rossi I, Montagne L, Salem N, et al. Brachytherapy for oligometastatic prostate cancer to the penis. *J Contemp Brachytherapy.* 2021;13(5):593-7.
4. Chaux A, Amin M, Cubilla AL, Young RH. Metastatic



- tumors to the penis: a report of 17 cases and review of the literature. *Int J Surg Pathol*. 2011;19(5):597-606.
5. Jin JK, Dayyani F, Gallick GE. Steps in prostate cancer progression that lead to bone metastasis. *Int J Cancer*. 2011;128(11):2545-61.
6. Soma S, Reddy PC, Bhat R, Prabhu S. Penile metastases from prostate adenocarcinoma: a rare presentation. *J Clin Diagn Res*. 2015;9(9):PD03-PD04.
7. Mahmoud B, Allouche M. Penile metastasis from prostate cancer: a rare case. *Asian J Res Rep Urol*. 2021;4(4):117-21.
8. Brito Y, Wilson BW, Bacchus KI, Mwaniki J, Jorge J, Tiesenga F. Cutaneous metastases in progressive prostate adenocarcinoma: a case report and literature review. *Cureus*. 2023;15(9):e45864.
9. Gurel B, Ali TZ, Montgomery EA, Begum S, Hicks J, Goggins M, et al. NKX3.1 as a marker of prostatic origin in metastatic tumors. *Am J Surg Pathol*. 2010;34(8):1097-105.
10. Nova-Camacho LM, Collins K, Trpkov K, Acosta AM, Sangoi AR, Akgul M, et al. Metastatic solid tumours to the penis: a clinicopathologic evaluation of 109 cases from an international collaboration. *Histopathology*. 2023;83(1):31-39.
11. John J, Mngqi N, Aldera AP. Metastatic prostate carcinoma presenting as a gluteal soft tissue mass. *Ther Adv Urol*. 2022;14:17562872221096384.
12. Mottet N, van den Bergh RC, Briers E, Van den Broeck T, Cumberbatch MG, De Santis M, et al. EAU-EANM-ESTRO-ESUR-SIOG guidelines on prostate cancer-2020 update. Part 1: screening, diagnosis, and local treatment with curative intent. *Eur Urol*. 2021;79(2):243-62.
13. Sweeney CJ, Chen YH, Carducci M, Liu G, Jarrard DF, Eisenberger M, et al. Chemohormonal therapy in metastatic hormone-sensitive prostate cancer. *N Engl J Med*. 2015;373(8):737-46.
14. Sun Y, Wang BE, Leong KG, Yue P, Li L, Jhunhunwala S, et al. Androgen deprivation causes epithelial-mesenchymal transition in the prostate: implications for androgen-deprivation therapy. *Cancer Res*. 2012;72(2):527-36.
15. Borno HT, Lichtensztajn DY, Gomez SL, Palmer NR, Ryan CJ. Differential use of medical versus surgical androgen deprivation therapy for patients with metastatic prostate cancer. *Cancer*. 2019;125(3):453-62.
16. Garje R, Chennamadhavuni A, Mott SL, Chambers IM, Gellhaus P, Zakharia Y, et al. Utilization and outcomes of surgical castration in comparison to medical castration in metastatic prostate cancer. *Clin Genitourin Cancer*. 2020;18(2):e157-e166.
17. Saad F, Gleason DM, Murray R, Tchekmedyian S, Venner P, Lacombe L, et al. Long-term efficacy of zoledronic acid for the prevention of skeletal complications in patients with metastatic hormone-refractory prostate cancer. *J Natl Cancer Inst*. 2004;96(11):879-82.
18. Roth AJ, Weinberger MI, Nelson CJ. Prostate cancer: psychosocial implications and management. *Future Oncol*. 2008;4(4):561-8.