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Case Report

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Ovarian Carcinoma: A Rare Case of Post-Kidney Transplant Malignancy – A Case Report

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Abstract: Background: Post-transplant malignancies are a recognized complication of long-term immunosuppression in renal transplant recipients, with incidences significantly higher than in the general population [1,2]. Ovarian carcinosarcoma remains a particularly rare and aggressive post-transplant tumour [3]. Case Presentation: A 54-year-old female renal transplant recipient presented with progressive abdominal distension and weight loss 3 years posttransplant. Imaging revealed massive ascites and a large complex ovarian mass. Histopathology confirmed carcinosarcoma of the ovary. Despite initiation of chemotherapy, the patient died three weeks later. Conclusion: This case highlights the importance of heightened surveillance for rare malignancies in immunosuppressed transplant recipients. Ovarian carcinosarcoma is a rare but intervention diagnosis, necessitating fatal prompt and potential immunosuppression adjustment.

Keywords: Carcinosarcoma, ovary, renal transplant, post-transplant malignancy, immunosuppression.

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INTRODUCTION

Post-transplant malignancies are a growing cause of late morbidity and mortality in renal transplant recipients, with a 2–15-fold higher incidence compared to the general population [1,2]. Among these, ovarian carcinosarcoma—a malignant mixed Müllerian tumour—remains exceedingly rare, accounting for only 1–2% of all ovarian neoplasms [3,4]. These tumours contain both epithelial and sarcomatous malignant components [5] and are known for their aggressive clinical course and poor prognosis [6].

This report presents a rare case of ovarian carcinosarcoma occurring in a renal transplant recipient, underlining the need for increased vigilance and early diagnosis in such patients.

CASE PRESENTATION

A 54-year-old woman who had undergone kidney transplantation three years earlier due to hypertensive nephropathy presented with a 6-month history of progressive abdominal distension, weight loss, and bilateral pitting oedema of the lower limbs. She was compliant with her immunosuppressive regimen (tacrolimus, mycophenolate mofetil, and prednisolone).

Clinical examination revealed massive ascites. No organomegaly or abdominal masses could be appreciated due to fluid accumulation. Laboratory investigations showed markedly elevated serum CA-125 (13,069.2 U/mL; reference: 0–35). Abdominopelvic CT scan with IV contrast revealed a large ($12.8 \times 8.5 \times 11.4$ cm) complex left adnexal mass with both solid and cystic components, as well as massive ascites.

Ultrasound-guided biopsy of the mass revealed features of ovarian carcinosarcoma. Despite prompt

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initiation of chemotherapy, the patient died three weeks after diagnosis.

DISCUSSION

The risk of malignancy in renal transplant recipients is significantly elevated due to chronic immunosuppression, which reduces immune surveillance and facilitates oncogenic viral infections [2,7]. The most common post-transplant malignancies are skin cancers and lymphoproliferative disorders, but solid organ malignancies—though less common—carry a worse prognosis [8].

Ovarian carcinosarcoma is rare, accounting for only 1–2% of all ovarian neoplasms [3,4]. Its occurrence in transplant recipients is even rarer. The tumour's biphasic nature, comprising both carcinomatous and sarcomatous elements, contributes to its aggressiveness [5]. Immunosuppressive agents such as calcineurin inhibitors and antimetabolites are known to impair cellular immunity and may contribute to tumour growth [6].

This patient's poor outcome was likely due to advanced disease at presentation, high tumour burden, and limited time for therapeutic intervention. While tumour debulking and chemotherapy are the mainstay of treatment [6], immunosuppressive dose adjustment or switching to mTOR inhibitors might be considered as part of a comprehensive cancer control strategy in transplant recipients [10].

Regular cancer screening in transplant patients, particularly those who present with vague systemic symptoms or paraneoplastic features, is essential for early detection and improved outcomes [11].

CONCLUSION

Ovarian carcinosarcoma is an extremely rare but aggressive tumour in renal transplant recipients. Clinicians should maintain a high index of suspicion for malignancy in transplant patients with unexplained systemic symptoms. Early diagnosis and multidisciplinary care, including possible adjustments in immunosuppressive therapy, are key to improving survival.

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Ethical Approval: Approved by the University of Maiduguri Teaching Hospital Ethics Committee.

Consent: Informed consent for publication was obtained from the patient's next of kin.

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