Sarcomatoid Carcinoma: A Rare and unusual histological variant of prostatic carcinoma

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Submitted: 24-09-2014

Revised: 26-01-2015

Published: 01-05-2015

ABSTRACT

Carcinosarcoma of Prostate also known as Sarcomatoid carcinoma is a rare biphasic malignancy with a malignant epithelial component (carcinomatous) and a malignant mesenchymal component. We report a case of this rare and unusual histological variant of prostatic carcinoma.

Key Words: Sarcomatoid, Prostate specific antigen, Prostate carcinoma

INTRODUCTION

Malignant tumors of the prostate that display biphasic patterns are very rare. They include sarcomatoid carcinomas (SC) and carcinosarcomas (CS). It is an admixture of high grade epithelial and sarcomatoid components, although it is still unclear that these cases represent a co-existent epithelial and mesenchymal elements (carcinosarcoma) or an evolution of an underlying adenocarcinoma into a lesion with associated sarcomatoid features and occasional heterologous elements (sarcomatoid carcinoma).¹ However, WHO classification of urinary tract tumor does not distinguish between Sarcomatoid carcinoma (SC) and Carcinosarcoma (CS) and mentions all these lesions under the term "Sarcomatoid carcinoma". In contrast to adenocarcinoma Prostate, it is an aggressive malignancy, while the treatment is non-standardized. The patient may present with irritative symptoms (frequency, nocturia, urgency) and/or obstructive symptoms (poor stream, incomplete emptying or retention). Hematuria, perineal/ rectal pain and burning on ejaculation are uncommon symptoms.² Hereby, we report a case of a 65 year old gentleman with Sarcomatoid carcinoma Prostate and discuss the treatment protocol of this uncommon tumor.

Case report

Address for Correspondence:

A 65 year old gentleman presented to our department with complaints of increased frequency of urine for last 6 months and back ache for the last two months. Digital rectal examination revealed grade II prostatomegaly, with left lobe of Prostate disproportionately enlarged and hard in consistency. There was no palpable lymphadenopathy, and the rest of his physical examination was unremarkable. His serum PSA was 495 ng/ml. A tru-cut biopsy of Prostate showed malignant epithelial cells forming glands (Adenocarcinoma) scattered in a stroma composed of bizarre mesenchymal cells (Sarcoma) with infiltration into adipose tissue consistent with diagnosis of Carcinosarcoma Prostate (Gleason score 4+5) (Figures 1 and 2).

In view of biopsy findings patient underwent further evaluation and found to have bone metastases. Patient was started on anti-androgen therapy and bilateral subcapsular Orchiectomy was done. Palliative External beam Radiotherapy was recommended to painful bone metastases along with Inj Bisphosphonates monthly.

DISCUSSION

Carcinosarcoma of Prostate also known as Sarcomatoid carcinoma is a rare biphasic malignancy with a malignant epithelial component (carcinomatous) and a malignant mesenchymal component (sarcomatous) with the presence or absence of heterologous elements.^{3,4} The histogenesis of prostate carcinosarcoma is not completely understood. Proposed mechanisms include transformation of

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Access this article online Website: http://nepjol.info/index.php/AJMS DOI: 10.3126/ajms.v6i5.12336

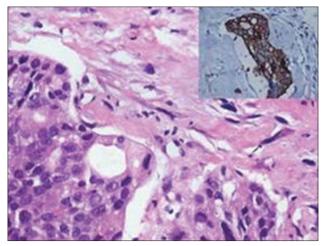


Figure 1: High grade adenocarcinoma admixed with malignant spindle cells (Hematoxylin Eosin stain X 400). Inset: Carcinomatous element is strongly positive for cytokeratin (cytokeratin immunohistochemistry x 100).

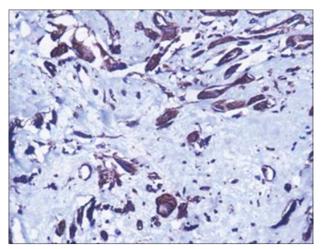


Figure 2: Malignant Spindle cells showing vimentin positivity (immunohistochemistry for vimentin x 400).

epithelial elements into a sarcomatous component and divergent differentiation of epithelial stem cells into both malignant components. WHO classification use the term "Sarcomatoid carcinoma" to denote all of these lesions.⁵ The carcinomatous component is usually acinar but may be ductal adenocarcinoma.⁶ Squamous or adenosquamous carcinoma⁷ or mixed urothelial squamous components⁸ and is usually of high grade.⁹ The type of sarcoma may vary, including elements such as osteosarcoma, chondrosarcoma, rhabdomyosarcoma, leiomyosarcoma and angiosarcoma. In about half of all cases, the initial diagnosis was a usual acinar adenocarcinoma, followed by hormonal and/or radiation therapy, with a subsequent diagnosis of sarcomatoid carcinoma. By immunohistochemistry, epithelial elements react with cytokeratins, PSA, PSAP whereas sarcomatoid elements react with vimentin or specific markers corresponding to the mesenchymal differentiation.¹⁰ The sarcomatoid component is considered to be Gleason grade 5, with the glandular element, if present, assigned a grade according to the usual rules. PSA values are usually within normal limits, possibly due to "dedifferentiation" of tumor cells.⁸

The prognosis of prostate carcinosarcoma is poor, with death invariably occurring within 2 years regardless of treatment modality.

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Authors Contribution:

AG & BC - operated the case, Pu P-collection of photographs; AG- compiled the case report; NW & PP - pathological content of case report.

Source of Support: Nil, Conflict of Interest: None declared.