Editorial

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The Metamorphosed Girdle-Carcinoma Prostate

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Introduction

Carcinoma prostate is a frequently discerned malignancy of prostate gland emerging from prostatic secretory epithelium. Tumefaction is devoid of speci c or sensitive clinical and radiological features which may assist appropriate disease discernment. Neoplasm is frequently detected by non-targeted ne or core needle tissue samples obtained secondary to evaluation of elevated serum prostate speci c antigen (PSA) levels [1, 2]. Additionally designated as prostate cancer or prostatic adenocarcinoma, carcinoma prostate exhibits distinctive subtypes as acinar adenocarcinoma, ductal adenocarcinoma, atrophic adenocarcinoma, pseudo-hyperplastic adenocarcinoma, microcystic adenocarcinoma, foamy gland adenocarcinoma, pleomorphic giant cell adenocarcinoma or sarcomatoid adenocarcinoma [1, 2].

Majority of carcinoma prostate are multifocal and predominantly (~80%) con ned to posterior or posterolateral peripheral zone. Few

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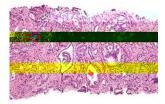


Figure 1: Carcinoma prostate depicting neoplastic glands layered by transformed columnar epithelium admixed with glands imbued with intraluminal eosinophilic

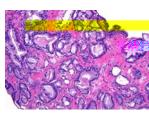


Figure 2: Carcinoma prostate delineating glands lined with neoplastic and

TNM classi cation of carcinoma prostate is designated as **Primary** tumour.

• T1: Tumour discerned upon TURP, ne or core needle biopsy.

• T2: Tumour con ned to organ and categorized as ~T2a: Tumour discerned with Digital Rectal Examination (DRE) or transrectal ultrasound and appears unilateral, ~T2b: Tumour discerned with Digital Rectal Examination (DRE) or trans-rectal ultrasound and extends to contralateral segment, ~T2c: Tumour discerned with Digital Rectal Examination (DRE) or trans-rectal ultrasound and appears bilateral

• T3 designated as ~T3a: Tumour with focal or multifocal extra-prostatic extension or microscopic invasion of dense musculature of urinary bladder neck with absence of adjacent non-neoplastic glands, ~T3b: Tumour invasion into musculature of seminal vesicle.

• T4: Tumour invasion into external urethral sphincter, rectum, bladder, levator muscles or pelvic wall.

Regional Lymph Nodes

- NX: Regional lymph nodes cannot be assessed
- N0: Regional lymph node metastasis absent
- N1: Regional lymph node metastasis present

Regional lymph nodes are exempli ed by peri-prostatic, pelvic, hypogastric, obturator, internal iliac, external iliac or sacral lymph nodes.

Distant Metastasis

M0: Distant metastasis absent.

• M1 is designated as ~M1a: Distant metastasis into nonregional lymph nodes as aortic, common iliac, super cial inguinal, deep inguinal or retroperitoneal lymph nodes, ~M1b: Distant metastasis into bone, ~M1c: Distant metastasis into diverse viscera [2, 5].

Gleason grading of carcinoma prostate is contingent to tumour architecture and represents a morphological continuum of wellformed glandular pattern to complex glandular proliferation or lack of glandular di erentiation. Gleason grades are designated as:

Grade X: Gleason score cannot be determined.

• Grade 1(Gleason score 6): Individual, discrete, well con gured glands and serum PSA <10 ng/ml or between 10 ng/ml and 20 ng/ml.

• Grade 2(Gleason score 3+4=7): Predominant well con gured glands with a minimal component of inadequately de ned, fused or cribriform glands and serum PSA <20ng/ml.

• Grade 3 (Gleason score 4+3=7 or 8): Predominant inadequately con gured, fused or cribriform glands with miniscule component of well con gured glands and serum PSA <20ng/ml.

• Grade 4 (Gleason score 8): Singularly composed of inadequately con gured, fused or cribriform glands or predominantly well con gured glands with minor component devoid of glands or a tumefaction predominantly devoid of glands with miniature component of well con gured glands and serum PSA 20 ng/ml.

• Grade 5 (Gleason score 9 or 10): Absence of glandular formation or glands associated with necrosis along with or devoid of inadequately con gured, fused or cribriform glands and an elevated serum PSA level [2, 5].

Clinically, grade 1 denominates a morphological 'low grade', grade 2 exempli es an 'intermediate grade' and grade 3 or elevated grade enunciates 'high grade' neoplasms [2, 5].

Staging of carcinoma prostate is denominated as:

• Stage I: Tumour is preliminary, non-palpable, and gradually progressive and incriminates unilateral aspect of prostate. Serum PSA levels are minimal. Tumour cells simulate uninvolved prostatic epithelium.

• Stage II: Tumour con ned to prostate. Serum PSA levels are minimal to moderate. Progression and dissemination of miniature tumefaction is exempli ed.

• Stage IIA: Tumour is non-palpable and unilateral. Serum PSA levels are moderately elevated. Neoplastic cells are well di erentiated. Enlarged neoplasms composed of well di erentiated prostatic epithelial cells con ned to prostate are categorized as stage IIA neoplasms.

• Stage IIB: Tumour is con ned to prostate and appears palpable upon direct rectal examination (DRE). Serum PSA levels are moderately elevated. Tumour cells are moderately di erentiated.

• Stage IIC: Tumefaction is con ned to prostate and appears palpable upon DRE. Serum PSA levels are moderately elevated. Neoplastic cells are moderately or poorly di erentiated.

• Stage III: Serum PSA levels are signi cantly elevated. Tumour is high grade and progressive. e stage is constituted of locally advanced prostatic carcinoma with signi cant tumour progression and metastasis.

• Stage IIIA: Tumefaction extends beyond extrinsic layer of prostate into circumscribing so tissues or seminal vesicles. Serum PSA levels are signi cantly elevated.

• Stage IIIB: Tumefaction extends beyond prostate gland with extension into adjacent viscera as urinary bladder or rectum.

• Stage IIIC: Tumour cells appear poorly di erentiated.

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• Stage IV: Tumefaction extends beyond prostate ~ stage IVA: Tumefaction depicts regional lymph node metastasis, ~stage IVB: Tumefaction disseminates into distant lymph nodes, diverse viscera or bones [2, 5].

Recurrent carcinoma prostate is denominated by neoplastic reappearance following therapy. Tumour may re-emerge within prostatic zone or diverse areas. Extent of tthin