Prostate tumors: an overview

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Classification

WHO histological classification of tumors of the prostate (2004):
Epithelial tumours
Glandular neoplasms:
Adenocarcinoma (acinar)
- Atrophic
- Pseudohyperplastic
- Foamy
- Colloid
- Signet ring
- Oncocytic
- Lymphoepithelioma-like
Carcinoma with spindle cell differentiation
(carcinosarcoma, sarcomatoid carcinoma)
Prostatic intraepithelial neoplasia (PIN)
Prostatic intraepithelial neoplasia, grade III (PIN III)
Ductal adenocarcinoma
- Cribriform
- Papillary
- Solid
Urothelial tumours:
Urothelial carcinoma
Squamous tumours:
Adenosquamous carcinoma
Squamous cell carcinoma
Basal cell tumours:
Basal cell adenoma
Basal cell carcinoma
Neuroendocrine tumours
Endocrine differentiation within adenocarcinoma
Carcinoid tumour
Small cell carcinoma
Paraganglioma
Neuroblastoma
Prostatic stromal tumours
Stromal tumour of uncertain malignant potential
Stromal sarcoma
Mesenchymal tumours
Leiomyosarcoma
Rhabdomyosarcoma
Chondrosarcoma
Angiosarcoma
Malignant fibrous histiocytoma
Malignant peripheral nerve sheath tumour
Haemangioma
Chondroma
Leiomyoma
Granular cell tumour
Haemangiopericytoma
Solitary fibrous tumour
Hematolymphoid tumours
Lymphoma
Leukaemia
Miscellaneous tumours
Cystadenoma
Nephroblastoma (Wilm's tumour)
Rhabdoid tumour
Germ cell tumours
Yolk sac tumour
Seminoma
Embryonal carcinoma & teratoma
Choriocarcinoma
Clear cell adenocarcinoma
Melanoma
Metastatic tumours

TNM classification: The TNM system is the most widely used for the stratification of prostatic carcinoma and is the standard system only for prostate adenocarcinomas. The current revision of the TNM system is shown in the Table (ICD-O C61):

Classification of Prostatic Carcinoma: TNM classification:
T-Primary Tumor
TX Primary tumour cannot be assessed
T0 No evidence of primary tumor
T1 Clinically inapparent tumour, neither palpable nor visible by imaging
T1a Tumour incidental histological finding in 5% or less of tissue resected
T1b Tumour incidental histological finding in more than 5% of tissue resected
T1c Tumour identified by needle biopsy (e.g., because of elevated PSA)
T2 Tumour confined within prostate
T2a Tumour involves one-half of one lobe of less
T2b Tumour involves more than one-half of one lobe, but not both lobes
T2c Tumour involves both lobes
T3 Tumour extends through the prostatic capsule
T3a Extracapsular extension (unilateral or bilateral) including microscopic bladder neck involvement
T3b Tumour invades seminal vesicle(s)
T4 Tumour is fixed or invades adjacent structures other than seminal vesicles: external sphincter, rectum, levator muscles and/or pelvic wall
N-Regional Lymph Nodes
N0 No regional lymph node metastasis
N1 Regional lymph node metastasis
M-Distant Metastasis
M0 No distant metastasis
M1 Distant metastasis
M1a Non-regional lymph node(s)
M1b Bone(s)
M1c Other site(s)

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<td>Stage I T1, T2a N0 M0</td>
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In the US, it is estimated that 217730 men will be diagnosed with and 32050 men will die of cancer of the prostate in 2010 (NCI). In Europe, the number of new cases was estimated at 338732 men in 2008 with 70821 deaths (21.1%). The age-adjusted incidence rate was 156.0 per 100000 men per year in the US, 110.5 per 100000 men per year in the Europe and 178.7 men per year in the France.

The worldwide mortality was estimated to 258000 deaths in 2008. Mortality rates are generally high in predominantly black populations, very low in Asia and intermediate in Europe and Oceania.

Prostate cancer rates increase with age. From 2004-2008, the median age at diagnosis for cancer of the prostate was 67 years of age in the US and the median age at death for cancer of the prostate was about 80 years.

Incidence can be influenced by several risk factors including genetic susceptibility, environmental exposure (cigarette smoking, alcohol consumption, infectious agents, dietary fat, endogenous hormones, ...) and difference in health care and cancer registration. Large clues on risk factors for prostate cancer are still not found.

**Clinics**

Patients with prostate cancer can present with a variety of symptoms but are most often asymptomatic. The presenting symptoms can be broadly divided into:
- Bladder outflow obstruction symptoms: poor stream, hesitancy, sensation of incomplete emptying, frequency, urgency, urinary incontinence due to chronic urinary retention, recurrent urinary sepsis... The symptoms of bladder outlet obstruction are commonly related to concomitant nodular hyperplasia but may also result from the prostate cancer, especially if it is locally advanced.
- Symptoms related to local extension of the tumor: Pain, haematuria, rectal obstruction or bleeding, haemospermia, symptoms of renal failure...
- Symptoms of metastatic disease: include bone pain and tenderness. Involvement of the spinal column may lead to cord compression. Lymphatic involvement from prostate cancer may also cause a variety of symptoms including lymphoedema (particularly lower limb).

Digital rectal examination (DRE) should be performed and serum PSA measured in patients in whom there is clinical suspicion of prostate cancer or in those who wished to be screened.

Prostate-specific marker

PSA (Prostate-Specific Antigen) represents the best serum marker for prostate carcinoma. It is a 34 kD glycoprotein which is exclusively secreted from epithelial cells of the prostatic ducts. A small portion is absorbed into the blood. PSA is useful in diagnostic, staging and monitoring men who have prostate carcinoma, 4 ng/ml was established as the upper limit of normal. An elevation in the total PSA level can be the result of benign prostate hyperplasia, prostatitis,
prostatic manipulations (DRE, needle biopsy, transrectal ultrasound) and intravesical BCG therapy rather than prostate carcinoma. Despite the reasonable performance of PSA in this setting, serum PSA lacks high sensitivity and specificity for prostate cancer. To improve this point, several approaches have been used including PSA density, PSA velocity and the fraction of free serum PSA.

Pathology

Definitive diagnosis requires biopsies of the prostate, using a needle under transrectal ultrasound-Guidance. The most prostate cancers are adenocarcinomas (95%) which develop from the secretory luminal cells of the prostatic gland, 70% arise in the peripheral zone, 15-20% arise in the central zone and 10-15% arise in the transitional zone. The most prostate cancers are multifocal.

The studies from J. Mc Neal on the zonal topography of the prostate and from D. Gleason on the tumoral differentiation constituted the basis of anatomopathological descriptions of prostate cancers. The most widely internationally accepted grading model is the Gleason score based on the progressive loss of the gland pattern and the increased peritumoural stroma invasion. Histopathological grading and Gleason scores range as follow:

- Histopathological Grading and Gleason score
- GX: Grade cannot be assessed.
- G1: Well differentiated, Gleason 2-4.
- G2: Moderately differentiated; Gleason score 5-6.
- G3-4: Poorly differentiated/undifferentiated; Gleason score 7-10.

Treatment

Guidelines exist to help for treatment decisions. The choice of a specific treatment is based on the TNM staging and the evaluation of the low, intermediate or high-risk of prostate cancer. Therapeutic options for localized prostate cancer:

- Watchful waiting: or active surveillance is the best option for low risk cancers or for patients with a short life expectancy, estimated by age and co-morbidities. This therapeutic decision is also based on patient general health, potential side effects of treatment and patient preference.
- Radical prostatectomy: is the best treatment in localised prostate cancer for patients with at least 10 years life expectancy. This option is classically used in low or intermediate intra-capsular tumors. Otherwise, surgical treatment may include not only prostatectomy but lymphadenectomy.
- External-beam radiation therapy (EBRT): is one of the primary treatment modalities in localised and locally advanced prostate cancer. The introduction of 3D conformal radiotherapy in the early 1990s allows higher doses and safety radiation. The second generation 3D technique, intensity modulated radiotherapy (IMRT) is now required.
- Brachytherapy: interstitial permanent brachytherapy as monotherapy is indicated for patients with low risk cancer. Intermediate risk patient could benefit from the brachytherapy if they have only one of the pre-mentioned risk factors.
- Androgen deprivation: is not recommended in low and intermediate risk disease.
- Cryotherapy, HIFU (High-intensity focused ultrasound) and other focal therapies are not recommended as standard initial treatment.
- Therapeutic options for locally advanced prostate cancer (T3-4N0M0):
  - Management of locally advanced prostate cancer is usually EBRT combined to androgen deprivation therapy. Actually, a study of Messing and al. suggests that immediate androgen deprivation by surgical castration or LHRH agonist therapies decrease the risk of mortality. The benefit of neo-adjuvant hormone therapy remains unclear.
- Radical prostatectomy in this case is indicated only occasionally in a very high selected patients.

Management of metastatic disease

Androgen suppression using surgical castration or LHRH agonist therapies are the first line. Short-course anti-androgen should be used to prevent the disease flare up on starting an LHRH agonist.

Management of castration-refractory prostate cancer (CRPC)

- The patients with castration refractory prostate cancer should be considered for second line hormonal therapies by anti-androgen, corticosteroid, oestrogen or ketoconazol.
- Docetaxel (Taxotere): Chemotherapy with Docetaxel should be envisaged after the failure of all second lines hormonal manipulations and in a symptomatic disease. In a phase III trial, Tamock et al. demonstrated that Docetaxel plus prednimone every 3 weeks improved patients survival by 3 months over Mithoxantrone plus prednisone.
- Cabazitaxel (Jevtana): In june 2010, chemotherapy with Cabazitaxel was approved by the FDA (US Food and Drug Administration) for CRPC previously treated with Docetaxel containing regimen.
- Abiraterone (Zytiga): abiraterone is a new generation androgen inhibitor approved for CRPC previously treated with Docetaxel.
- Sipuleucel-T (Provenge): is an immunotherapy approved by the FDA since April 2010 for asymptomatic or minimally symptomatic prostate cancer resistant to standard hormonotherapy.
- Biphosphonates: mainly zoledronic acid (Zometa) may be offered to patients with skeletal metastasis to prevent osseous complications. Recently, RANK ligand inhibitor, such as denosumab has been developed. Denosumab has the effect to slow...
down metastasis in patients with hormonal therapy and also prevents osseous complications much more than zoledronic acid.
- Radiopharmaceuticals: like samarium and strontium, can be used for the management of painful osseous metastasis.

**Evolution**

The natural history of prostate cancer begins with a local tumour burden and a subsequent metastatic dissemination, preferentially to bones and the inescapable evolution to a castration refractory prostate cancer (CRPC).

**Prognosis**

Predicting prognosis is essential for patient decision making. Prognostic factors are used to select the appropriate treatment option(s) and to predict biochemical recurrence (also called PSA failure) after radical local therapy.

The most established independent prognostic factors are: the anatomical extend or stage of the disease (TNM stage) evaluated by digital rectal examination supplemented when necessary by ultrasound or MRI, the Gleason score, PSA level and post prostatectomy margin status. Clinically localized prostate cancer should be classified as low, intermediate and high risk.
- Low risk is corresponding to T1-T2a, Gleason 6, 10< PSA
- Intermediate risk is T2b, Gleason=7, 10< PSA
- High risk is =T2c, Gleason >7, PSA>20.

Imaging investigations are not recommended for men with low risk disease, in contrast CT or MRI and bone scintigraphy should be considered in high risk disease. Other prognostic normograms can be used to predict disease progression and patient survival.

**Genetics**

Several loci associated with an increased familial prostate cancer have been mapped to 1q24-25 (HPC1), 17p11 (HPC2), and Xq27-28 (HPCX). HPC1 corresponds to RNASEL which encodes an endoribonuclease for ssRNA implicated in the interferon response to viral infection.

**Cytogenetics**

Numerous single-nucleotide polymorphisms (SNPs) were reported that are associated with cancer risk. 8q24 is the most frequently gained chromosomal region in prostate tumors. Gain in 8q24 region has been associated with aggressive tumors, hormone independence, and poor prognosis. Although this region is frequently amplified in prostate tumors, it covered few known or predicted genes. The known genes that are closest to 8q24 are FAM84B and MYC.

Other independent risk loci for prostate cancer risk are MSMB (10q11.23); HNF1B (17q12); NUDT10/NUDT11 (Xp11.22); 17q12; KCLK2/3 (19q13.33); JAZF1 (7p15.1); 3p12.1; EHHB1 (2p15); CTBP2 (10q26.13); SLC22A3 (6q25.3); 22q13.

CGH (comparative genomic hybridization) was used to identify gains or losses of chromosomal regions. The most common alterations include chromosome 8 (23%) and chromosome 7 (20%). The most commonly reported are gains of 2p, 3q, 7q, 8q, 9q, 17q, 20q, and Xq. deletions of 2q, 5q, 6q, 8p, 10q, 12p, 13q, 16q, 17p, 17q, 18q, 21q, and 22q, hyperdiploidy, and aneusomy of chromosomes 7 and 17.

Outlier profile analysis was used to identify the TMPRSS2-ERG fusion as the single most prevalent gene fusion in prostate cancer (>60%) on 21q22. TMPRSS2 encodes for the androgen-activated transmembrane protease serine 2 and ERG encodes a member of the ETS transcription factor. Other gene fusions have been found with ETS genes (including ERG, ETV1, ETV4, and ETV5) involving different 5' partners (~10%): TMPRSS2 (t(7;21)(p21;q22)), SLC45A3 (t(1;7)(q32;p21.2)), AC8L3 (t(2;7)(q36.1;p21.2)), HES1 (t(7;22)(p21.2q11.23)), FLJ35294 (t(7;17)(p21.2;q13.1)), FOXP1 (t(3;7)(p13;q21)), EST14 (t(7;14)(p21.1;q21)), C15orf21 (t(7;15)(p21.3;q21)), and HNRPA2B1 (t(7;7)(p21.2;q15)) for ETV1; TMPRSS2 (t(17;21)(q21.1;q21)), KLK2 (t(17;19)(q21;q13)), CANT1 (inv(17;17)(q22;q22)) and DDX5 (t(17;17)(q24;q21)) for ETV4; TMPRSS2 (t(3;21)(q28;q22)) or SLC45A3 (t(1;3)(q32;q28)) for ETV5. TMPRSS2-ERG fusions may occur as an early event in the development of prostate cancer, but the gene fusion is not sufficient to cause cancer formation by itself. The prognostic significance of the TMPRSS2-ERG fusion and other ETS rearrangements in prostate cancer is still controversial. Integrative genomic profiling was used to identify a narrow deletion on 3p14 is highly associated with TMPRSS2-ERG fusion-positive tumors. PTEN loss is also associated with the presence of the fusion gene TMPRSS2-ERG. For more details on the fusion gene TMPRSS2-ERG, see the section "Deep insight": TMPRSS2:ETS gene fusions in prostate cancer.

**Genes involved and proteins**

Note
Somatic point mutations in prostate cancer are rare relative to other tumor types such as glioblastoma, lung cancer and melanoma; alterations of gene copy number are more frequent.

**AR**

Location
Xq11-12
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Protein
AR gene encodes for the nuclear receptor for the androgens, which belongs to the steroid/thyroid hormone receptor gene superfamily. This nuclear receptor mediates hormone action by binding to hormone in the cytoplasm and translocating to the nucleus where it dimerizes and binds DNA at androgen-responsive gene promoter to modify the transcription of target genes.

Somatic mutations
Almost all prostate cancers express AR. AR expression is maintained during prostate carcinogenesis from primary prostate cancer to castration-refractory prostate cancer (CRPC). AR was found overexpressed in all CRPC and plays a predominant role in the transition from androgen-dependent to independent cancer.
The AR plays a role of a dominant oncogene in castration-resistant prostate cancer. AR gene amplification has been recovered in ~30% of CRPC.
AR is also activated by missense mutations (frequencies of 10-30%) that modify the scope of hormone specificity and/or enhance hormonal response. Activation of AR in androgen-independent disease may also be accomplished by activation of co-regulators that interact with AR to activate gene transcription. Another potential AR activation mechanism is a ligand-independent activation by growth-factors or cytokines. Several structurally different AR variants with divergent biologic activity recently identified could emerge as a primary cause of resistance to androgens. All of those variants required full-length AR through dimerisation to activate endogenous AR target genes and confer castration-resistant growth.

BRCA2
Location
13q12.3
Protein
BRCA2 encodes a large DNA repair protein. As for BRCA1, the primary roles of BRCA2 as a tumor suppressor gene is to maintain genomic stability through DNA repair system and is involved in transcriptional regulation process.

Germinal mutations
Mutations in BRCA2 and BRCA1 are responsible for the largest number of inherited breast cancers and a large proportion of ovarian cancers. While in BRCA1 mutation carriers, the risk increase did not reach significance, BRCA2 mutations were associated with a significantly increased risk of prostate cancer. Its mutation accounts for 30-35% of familial breast cancers. Prostate cancer screening by PSA assay and digital rectal examination annually are recommended at age 40-50 years for male carriers of BRCA mutations.

Somatic mutations
The BRCA2 mutation does not occur in sporadic prostate cancer.

CDKN1B
Location
12p13.1
Protein
The gene CDKN1B encodes the cyclin-dependent kinase inhibitor p27Kip1 that inhibit cell proliferation.

Somatic mutations
Loss or reduced expression of CDKN1B correlates with high grade prostate cancer and reduced disease-free survival. The CDKN1B gene is haploinsufficient, so hemizygous deletion reduces the expression of CDKN1B and affects its normal function through dosage reduction. In mouse, concomitant inactivation of PTEN and CDKN1B accelerates spontaneous neoplastic transformation and incidence of tumors.

EZH2
Location
7q36.1
Protein
The Polycomb group gene EZH2 encodes a histone lysine methyltransferase which methylates histone H3 on lysine-9 and dimethyl lysine-27 leading to transcriptional repression of the affected target gene.

Somatic mutations
The gene EZH2 is frequently up-regulated in advanced prostate cancer, in some cases through gene amplification. Overexpression of EZH2 confers a poor prognosis in localized prostate cancers.

KLK3
Location
19q13.41
Protein
The KLK3 gene encodes a kallikrein of the serine protease family. KLK3, like KLK2, is involved in semen liquefaction in the normal prostate by degrading seminogelin I/II, fibronectin and laminin. Nevertheless, KLK3 function can also stimulate prostate cancer cell growth and promote metastasis by activating and/or degrading a variety of substrates such as PTHRP, IGFBP3, IGFBP4 and TGFbeta. Independently of its serine protease activity, KLK3 is able to stimulate the production of reactive oxygen species in prostate cancer cells, probably by binding to a cell surface receptor.
The KLK3 protein is commonly known as the Prostate-Specific Antigen (PSA) which is the most acceptable and broadly used cancer biomarker. Serum level of PSA in the clinical setting is useful in the diagnosis and monitoring of prostatic carcinoma.

GSTP1
Location
11q13.2
Protein

The GSTP1 gene encodes an enzyme that plays an important role in detoxification by catalyzing the conjugation of many hydrophobic and electrophilic compounds with reduced glutathione. The GSTP1 gene is involved in the DNA methylation of CpG island in prostate cancer and other cancers.

Somatic mutations

Silencing of GSTP1 expression by promoter hypermethylation is an early event in prostate cancer initiation, which is seen in up to 70% of PIN lesions and in 90-95% of prostate carcinomas. Decreased GSTP1 expression may predispose luminal cells to increased oxidative damage which in turn result in accumulation of genetic changes ultimately resulting in cell transformation and cancer.

**MYC**

**Location**
8q24

**Protein**

The proto-oncogene MYC encodes a transcription factor involved in activation of the cell-cycle progression and protein biosynthesis which is frequently amplified in variety of human malignancies.

Somatic mutations

Amplification is observed in approximately 30-40% of primary prostate tumors and 90% of metastatic prostate cancers and is associated with poor patient outcome. The locus at 8q24 was identified as a major susceptibility locus in several large-scale genome-wide association studies of prostate cancer as well as other epithelial cancers. Experiments with mouse models of prostate cancer provided strong causal connection between the overexpression of MYC and the development of prostate cancer.

**NKX3.1**

**Location**
8p21.2

**Protein**

The NKX3.1 gene encodes a prostate-specific homeobox transcription factor considered as a putative prostate tumor suppressor that is expressed in a largely prostate-specific and androgen-regulated manner.

Germinai mutations

Human germ line mutations in Nkx3.1, which decreases its interaction with DNA, result in genetic predisposition to prostate cancer.

Somatic mutations

Loss of NKX3.1 protein expression is a common finding in 12% prostatic intraepithelial neoplasia (PIN) and in 85% human prostate carcinomas. Nkx3.1 has been shown to be a critical regulator of prostate epithelial differentiation and stem cell function in mouse models. NKX3.1 represents a haploinsufficient tumor suppressor gene that acts as a "gatekeeper" gene for prostate cancer initiation.

**p53**

**Location**
17p13

**Protein**

The tumor suppressor gene PTEN encodes a lipid phosphatase that negatively regulates intracellular levels of phosphatidylinositol-3,4,5-trisphosphate in cells to activate PI3K/Akt signaling pathway.

Somatic mutations

17% of primary prostate tumors show a loss of PTEN protein and there is significant correlation between PTEN loss and Gleason Score and clinical stage. PTEN loss is associated with the presence of the fusion gene TMPRSS2-ERG.

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