Prostate-specific kallikrein-related peptidases and their relation to prostate cancer biology and detection

Established relevance and emerging roles

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Summary

Kallikreins are a family of serine proteases with a range of tissue-specific and essential proteolytic functions. Among the best studied are the prostate tissue-specific KLK2 and KLK3 genes and their secreted protease products, human kallikrein 2, hK2, and prostate-specific antigen (PSA). Members of the so-called classic kallikreins, these highly active trypsin-like serine proteases play established roles in human reproduction. Both hK2 and PSA expression is regulated by the androgen receptor which has a fundamental role in prostate tissue development and progression of disease. This feature, combined with the ability to sensitively detect different forms of these proteins in blood and biopsies, result in a crucially important biomarker for the presence and recurrence of cancer. Emerging evidence has begun to suggest a role for these kallikreins in critical vascular events. This review discusses the established and developing biological roles of hK2 and PSA, as well as the historical and advanced use of their detection to accurately and non-invasively detect and guide treatment of prostatic disease.

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Introduction

Tissue kallikrein and kallikrein-related peptidases (KLKs) comprise a family of 15 homologous secreted trypsin- or chymotrypsin-like serine proteases. These enzymes play important roles in a variety of complex processes including reproductive function, inflammation, skin homeostasis, blood clotting, fibrinolysis, and possibly cancer. Kallikreins are expressed in numerous tissues, are commonly co-expressed (1), and are localised predominantly in the cytoplasm of glandular epithelia. The multiplicity of these enzymes and their diverse roles has attracted considerable interest for use as diagnostic markers and for therapeutic targets across many pathologies. Despite their importance, many of the functions of this family of proteases remain unknown. The presence and forms of the kallikreins found in glandular excretions, such as breast milk, seminal fluid, and other fluids, such as blood and urine, have long been investigated in order to better understand the vital functions of KLK-gene products and to detect disease.

KLK2 and KLK3 have the most organ restricted expression profile of all KLKs; specifically, they are abundantly expressed in the luminal epithelium of the prostate. KLK2-KLK3 constitute two of the few highly expressed genes and secreted proteins produced by this gland. While low expression may be detected in other organs, the levels found in extra-prostatic tissue are many orders of magnitude lower, and the biological significance of KLK2-KLK3 in these tissues remains uncertain. The abundant production in the prostate helps to explain why the KLK3-gene product is commonly called prostate-specific antigen or PSA. By convention, we will refer to KLK2 as “human Kallikrein 2” (or hK2).

KLK2 and KLK3 are also unique in that they are regulated by androgens, with expression levels reflecting the functional status and activity of the nuclear androgen receptor (AR) and its response to supply of testosterone or other androgens (2). The unique combination of tissue-specificity and androgen-driven expression profile of KLK2 and KLK3 provide a straightforward and biologically relevant read-out of the activity of the prostate epithelium. This is particularly useful in the context of detecting and monitoring prostate cancer. This review will focus on these two most well studied and prostate-specific kallikreins, PSA and hK2. We will introduce and discuss their roles in prostate and prostate cancer biology, in detecting prostate cancer, the most commonly diagnosed malignancy in men (3) and their potential contribution to vascular disorders.

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Kallikrein genetics

Located in series at chromosomal region 19q13.3-q13.4, the kallikreins comprise the largest cluster of peptidases in the human genome (4) (Figure 1A). Each of the tissue kallikrein genes shares similar encoding features. This includes five exons of similar size separated by four introns of varying sizes (5) (Figure 1B). KLK1–3 are considered the "classic kallikreins" due to phylogenetic analysis showing a distinct evolutionary history from the other human kallikreins. It is likely that a duplication of KLK1 created the progenitor of KLK2 in the early evolution of eutherian mammals, with subsequent primate-specific duplication of the predecessor that gave rise to KLK3 and KLK2 (6, 7). The progenitor of KLK2 is a non-functional pseudogene in many mammals including rodents, but not in canines in which expression is regulated by AR and abundant in the canine prostate (6). Distinct from KLK4–15, KLK1–3 also share a unique surface loop called the "kallikrein loop". The evolutionary relationship is also reflected in the amino acid sequences; the identity in amino acid sequence with KLK1 is much higher for KLK2 or KLK3 (62–67%) than for KLK4–15 (27–39%). KLK2 and KLK3 manifest 80% identity in amino acid sequence, making them the most closely homologous KLK-gene products.

KLK2,3 activity

The KLK family of proteases has important functions in a range of tissues. Therefore, regulation of the enzymes’ activities may be essential for the maintenance of health, and dysregulation of such activity can be used as a marker for disease (Table 1, and following sections). This perhaps explains why the proteases are produced as non-catalytic preproenzymes requiring multiple post-translational modifications to result in a catalytically active form (Figure 1B). To obtain the catalytic PSA or hK2, the proteolytic cleavage of the signal sequence by a signal peptidase precedes the proteolytic release of the short N-terminal activation peptide by trypsin-like peptidase(s) converting the non-catalytic zymogen to an enzymatically active 237-amino acid single-chain form of PSA or hK2 (8). The trypsin-like tertiary structure differs between each kallikrein, resulting in variation in specificity and cleaving/activating motifs of physiologic substrates (Figure 1C).

Catalytically active PSA is secreted into the seminal fluid (9). It is there, along with secreted and possibly auto-activated hK2 [10], that the main gel forming proteins (SEMG1 and SEMG2) contribute by the seminal vesicles of the ejaculatory mix can be proteolytically degraded by KLK3 (possibly also KLK2) to liquefy the seminal gel (9, 11, 12). This critical action results in the freeing of spermatozoa for the necessary translocation to the womb to fertilise the egg.

Cleavage of semenogelin and fibronectin in the ejaculate suggest that these serine proteases could degrade extracellular matrix proteins elsewhere (11). Accordingly, investigations have focused on a potential link between KLK2–3 function and the invasion and growth of prostate cancer. While the explicit role(s) of the prostate-tissue kallikreins in prostate cancer development and progression remains unclear, there is considerable interest in determining if such links exist. For example, the activation of urokinase by hK2 provides a scenario in which this enzyme initiates a proteolytic cascade to degrade the tissues surrounding a tumour, contributing to expansion (13). In addition to degrading the surrounding tissue, it has been reported that hK2 and PSA activity can directly promote cancer cell growth. Studies have shown that the kallikreins are able to cleave insulin-like growth factor proteins, yielding a growth factor for prostate cancer (14–16) and may regulate the activity of PTHrP in vitro (17).

Described in detail in the following section, catalytically active PSA interacts with native α2-macroglobulin (A2M) to form stable covalent complexes (18), while A2M has been shown to bind cell-surface glucose-regulated protein (GRP78). This activates canonical proliferative and survival pathways such as Akt, MEK and ERK signalling cascades. This work suggests a critical role for PSA in driving aggressive disease as it induces a feed-forward loop for the production of more of the PSA-activated α2-macroglobulin complex (19). Recently, galectin-3 in the seminal fluid has been identified as a substrate for PSA (20). This may explain the presence of cleaved galectin-3 at sites of prostate cancer, with potentially important consequences for tumour development and angiogenesis by this biologically important lectin (21).

An important role for hK1 (the product of the KLK1 gene) involves the regulation of blood pressure homeostasis and inflammation. hK1 acts on kinogen to release the vasoactive peptide lys-bradykinin (22). At present, there exists no evidence that the catalytic action of PSA or hK2 could contribute a critical role in the kallikrein/kinin system. While hK2 has been shown to have some activity, it is strictly limited by its 1,000-fold lower kinin cleavage relative to hK1 (23).

Emerging work has evaluated the role of PSA activity on endothelial cell proliferation and migration, suggesting an anti-angiogenic effect. In vitro experiments have shown that human endothelial umbilical cord (HUVEC) proliferation and invasion was inhibited by addition of PSA to culture media. In vivo, using a metastatic melanoma model, high concentrations of systemically administered PSA reduced the number of metastases detected (24, 25). Indeed, in vitro studies have shown that inhibition of PSA activity (by small molecules [26], antibodies [27] and peptides [28]) negates the putative anti-angiogenic effects of PSA. With continually refined models of kallikrein expression in cancer prone and healthy mice, it is expected that greater clarity of the putative effects of the PSA and hK2 on the vasculature will be gained in the near future.

Prostate cancer and PSA and hK2 as biomarkers

Prostate cancer is highly dependent on AR signalling for both cancer cell proliferation and survival. There is no definitive explanation for how the generally slow-growing adenocarcinoma forms. However, the dependence of the disease on AR can be utilised for...
Historically, the only early detection strategy for prostate cancer was digital rectal exam (DRE). However, as a screening tool DRE is far from perfect most notably for finding many tumours that are no longer curable by the time they are palpable (29). Hence, there was a need for a better test that could detect asymptomatic disease at an earlier stage. This call was met with the discovery and implementation of the PSA blood test in the late 1980s (30, 31). The serum PSA test, which is the most commonly used cancer biomarker assay, refers more specifically to the measuring of the “total” immune-reactive concentration of PSA in serum or plasma. Normally, the architecture of the prostate gland maintains the highly abundant levels of PSA (10–100 μg/l) tightly confined to the glandular ducts and the secretory epithelium. In this healthy state, PSA occurs in the blood circulation at approximately one-millionth of the intra-ductal levels, with median blood levels in healthy adult males below age 50 close to 0.6 ng/ml (32, 33). Disease processes such as malignant transformation of the prostate epithelium may result in degradation of the basement membrane, loss of basal cell layer and glandular architecture, but experimental evidence is lacking as to whether this leads to increased release of PSA into the blood circulation where the levels may elevate up to Tissue kallikrein 

<table>
<thead>
<tr>
<th>Tissue kallikrein</th>
<th>Alternative names</th>
<th>Protease specificity</th>
<th>Tissue expression</th>
<th>Disease biomarker</th>
<th>Prostate-specific KLKs vascular substrates</th>
</tr>
</thead>
<tbody>
<tr>
<td>KLK1</td>
<td>Pancreatic/renal Kallikrein (hPRK), hK1</td>
<td>Trypsin/Chymotrypsin</td>
<td>A, K</td>
<td>P', B', K'</td>
<td></td>
</tr>
<tr>
<td>KLK2</td>
<td>Human (glandular Kallikrein 2 (hK2; hGK1)</td>
<td>Trypsin</td>
<td>P</td>
<td>P', B'</td>
<td>Fibronectin</td>
</tr>
<tr>
<td>KLK3</td>
<td>Prostate Specific Antigen (PSA; hK3, APS)</td>
<td>Chymotrypsin</td>
<td>B, P</td>
<td>P', B'</td>
<td>Plasminogen, Fibronectin</td>
</tr>
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<td>KLK4</td>
<td>Kallikrein-like 1 (KLK-L1), PRSS17, Enamel Matrix Serine Protease 1 (EMSP1), Androgen-Regulated Message 1 (ARM1), hK4, Prostate Serine-like Antigen</td>
<td>Trypsin</td>
<td>CNS, G, H, P, T</td>
<td>P', O', B'</td>
<td></td>
</tr>
<tr>
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<td>Kallikrein-like 2 (KLK-L2), Human Stratum Corneum Tryptic Enzyme (hSCTE), hK5</td>
<td>Trypsin</td>
<td>B, D, K, S</td>
<td>P', O', B', T', K'</td>
<td></td>
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<tr>
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<td>Serine Protease 9, Serine Protease 18, Zyme, Neurosin, Protease M, Myelen cephalon-specific Protease, hK6</td>
<td>Trypsin</td>
<td>CNS</td>
<td>B', O', K'</td>
<td></td>
</tr>
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<td>Chymotrypsin</td>
<td>A, C, D, K, L, O</td>
<td>O', K'</td>
<td></td>
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<tr>
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<td>Trypsin</td>
<td>B, CNS, D, K, S</td>
<td>O'</td>
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<tr>
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<td>O'</td>
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<td>B, K, O, P</td>
<td>P', O', B', T', K'</td>
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<td>O', P', K'</td>
<td></td>
</tr>
<tr>
<td>KLK12</td>
<td>Kallikrein-like 5 (KLK-L5), hK12</td>
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<td>I, P, S, X</td>
<td>B'</td>
<td></td>
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<td>K, S</td>
<td>B'</td>
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<tr>
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<td></td>
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<tr>
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<td>Prostinojen, ACO protease, hK15</td>
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<td>P, C, T</td>
<td>P', O', B', K'</td>
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Table 1: Kallikrein tissue distribution and function. This table summarises the alternative names of the 15 KLK gene products. These enzymes display either trypsin or chymotrypsin behaviour (or both). The tissue specific profile of the kallikrein family members is noted: P, prostate; C, colon; T, thyroid; A, arteries; K, kidney; B, Breast; CNS, central nervous system; G, adrenal; H, enamel of teeth; S, salivary glands; D, skin; O, ovary; L, liver; F, adipose tissue; I, intestine; X, bone. The dysregulation of these enzymes are established and potential biomarkers in a diverse number of malignancies, as noted by apostrophe (e.g. T', testicular cancer). Additionally, the prostate-specific kallikreins have speculated roles in vascular events by acting on such substrates as fibronectin and plasminogen.
Due to the catalytic activity of PSA and the 10,000-fold molar excess of proteinase inhibitors present in extra-cellular fluids, different forms of PSA are detected in blood (34). When released into blood, enzymatically-active PSA forms irreversible and covalently linked complexes with extra-cellular protease inhibitors such as α1-antichymotrypsin (ACT or SERPINA3), α2-macroglobulin (A2M), pregnancy-zone protein (PZP), α1-antitrypsin (SERPINA1) (35), or the protein C inhibitor (SERPINA5) (36). PSA bound to ACT constitutes the predominant fraction of the immunoreactive forms of PSA in blood (34, 37, 38).

In addition to these irreversibly linked complexes of the activated PSA, there are non-catalytic PSA-forms (19, 38). These proteins are not bound by any of the inhibitors and commonly referred to as "free PSA" and are present at proportions varying from less than 5% up to greater than 45% of the "total" immunoreactive PSA present in blood. The free PSA is proteolytically processed to an inactive form and is therefore not able to form complexes with inhibitors noted above, nor is it thought to have a role in cleaving vascular substrates. The different inactive free PSA-forms have been thoroughly characterised and comprise proprotein subtypes with varying extensions of the N-terminal propeptide (e.g. [-2] proPSA) (39), single-chain form(s) with truncated N-terminal (40), and non-catalytic single-chain protein, nicked PSA, which contains internal peptide bond cleavage(s) at Lys145 or Lys146, resulting in the loss of an antigenic epitope critical for high-affinity antibody binding to the non-catalytic single-chain free PSA-forms referred to as intact PSA (iPSA) (41). Tests have been developed that are able to distinguish many of the forms (42-44) and "total PSA" refers to the sum of the different forms of detectable PSA (19). With respect to hK2, levels in blood are commonly 10–2 of those determined for total PSA. Here, the majority of the non-catalytic hK2-forms in blood appear to be unbound to any of the inhibitors discussed above (45).

The measurement of the several fractions of PSA has been proposed with the aim of improving the specificity and sensitivity of total PSA. Multiple studies, including several meta-analyses, have shown that the determination of the percentage of free PSA (%fPSA) is useful to improve the detection of prostate cancer (46). More recently, several research teams have been successful in in-

**Figure 1: Kallikrein locus and enzymes.** The kallikrein family comprises the longest cluster of peptidases in the human genome. A) The 15 KLK genes, numbered, are found in rough series, located at 19q.13.3–19q.13.4. The KLK2 and KLK3 genes are prostate-tissue specific trypsin-like serine proteases, highlighted with dashed-line. B) The kallikreins have a conserved five exon, four intron, profile. Protein products are produced in a pre(i)-pro(ii)-enzyme(iii) form, requiring subsequent peptidase-activation in order to become catalytically active. C) The three-dimensional structure of trypsin is shown on the left. The structure of the two enzyme products from KLK2 and KLK3 are hK2 and PSA, respectively, as shown in the middle and at right. Here, the active serine site is visualised with the residue’s ball-and-stick model.

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creasing accuracy by also including the subforms iPSA or [-2] pro-PSA, and thereby creating detections systems called “indexes” or “panels”. The Prostate Health Index (phi) is a mathematical combination with [-2] proPSA according to the formula [-2] proPSA/ fPSA) x √tPSA (47). The panel of four kallikrein markers includes intact PSA and hK2 (48, 49). The underlying hypothesis of this approach is that the relative amounts of the different prostate-kallikrein subtypes are indicative of underlying disease biology, with implications for patient selection and management. While promising, a rigorous comparison, or combination, of these systems has yet not been completed.

**PSA screening, validation and concerns**

The introduction of the PSA-test marked a significant global shift towards earlier detection and improved management of prostate cancer. Over the past two decades, the age-adjusted death rate from prostate cancer has decreased by 40% in the US, owing to the combination of early detection with improvements in treatment (3).

The utility of PSA detection as a screening tool has been evaluated in trials in Europe and the US. The world’s largest randomised controlled PSA screening trial, with nearly 180,000 participants in eight countries was initiated in 1995 as the European Randomized Study of Screening for Prostate Cancer (ERSPC) (50). In January 1995, the Göteborg Randomized Population-based Prostate Cancer Screening Trial (51) began in Sweden, inviting 10,000 men for biennial PSA screening. This trial was integrated with the ERSPC in 1996. These two high-quality trials have demonstrated significant reductions in prostate-cancer mortality by screening men 55–69 years every two-four years and men 50–64 years every two years. The results show a 21% prostate-cancer mortality reduction in favour of screening after 11 years of follow-up and 44% after 14 years (51, 52).

The largest US trial, the Prostate, Lung, Colorectal and Ovarian (PLCO) Cancer Screening Trial, was also initiated in the early 1990s. However, in contrast to Europe, where PSA testing was not recommended and very few men had taken a PSA-test, opportunistic PSA testing had already become widespread in the US by this time (53). About half of the men in the control arm of the PLCO trial had a PSA test during the first years of the trial and 30% of men in the screening arm were pre-screened (53-55). This so-
called contamination diluted the PLCO's capacity to distinguishing any significant differences in prostate-cancer mortality between the groups. Accordingly, at 13 years of follow-up, there was no statistically significant difference in prostate-cancer mortality risk between the study groups (risk ratio 1.09; 95% confidence interval, 0.87–1.36) (56).

The evidence of reduced prostate cancer mortality at the population level may not translate to survival advantages at the individual level, generating considerable controversy. The benefits of PSA screening include early detection and enhanced cure, prevention of metastatic disease (57) and reduced prostate cancer mortality. However, these benefits must be weighed against potential harm to patient management in a generally slow growing disease. Impairments include anxiety/distress, the potential for false-positives, unnecessary prostate biopsies, over-diagnosis, infectious complications from biopsies and side-effects from treatment such as impotence and urinary incontinence with subsequent impact on quality of life (58). This imbalance between benefit and harm, is one of the reasons why the United States Preventive Services Task Force recommended against PSA screening in 2012 (59).

Although the relative cost-benefit ratio for the PSA testing continues to be examined, incorporating long-term follow-up data of studies such as the ERSPC, it is obvious that advanced screening strategies are needed to reduce disadvantages. Improvements to sensitivity and specificity can include reduction in false-positives and over-diagnosis. The way forward involves risk stratification approaches, where PSA-testing is used together with other clinical risk factors and markers, to predict individual risk – and takes into account the individual’s preferences (60). Many men demand the PSA-test and most guideline groups and professional urological organisations agree on and emphasise shared decision-making, stipulating that “PSA screening must be preceded by a discussion regarding risks and benefits” (60, 61) and that well-informed men suitable for screening should have access to PSA testing upon request (62).

**Advanced screening with KLK2 and KLK3**

Population representation, storage conditions of samples, completeness of follow-up and reproducibility are some of many important factors that need to be considered with respect to interpreting outcomes from retrospective biomarker studies. One of the few cohorts that meet these stringent criteria, and has a very low rate of PSA testing, is the Malmö Preventive Medicine cohort (MPP). The MPP is a large, representative, population-based study of cardiovascular risk factors that took place in Malmö, Sweden between 1974 and 1992 (63). The study included 22,444 men (participation rate 71.2%), of whom 21,277 were ≤50 years of age at the baseline blood draw. Through the end of 1999, 498 of the 21,277 participants who were aged less than 50 years at study participation had been diagnosed with prostate cancer.

Strikingly, studies based on the MPP data demonstrate that a single measurement of either PSA, or hK2, measured at or before age 50 predicts advanced prostate cancer diagnosed up to 30 years in advance (64–66). Interestingly, total PSA (the sum of PSA complexed with ACT and the free, uncomplexed form) is a stronger predictor in younger men, while percent free PSA and hK2 add important predictive value in older men and much closer to diagnosis (67). In summary, these studies show that the use of early PSA to stratify risk would allow a large group of low-risk men to be screened less often but increase frequency of testing on a more limited number of high-risk men.

**Beyond serum ELISAs: Emerging technologies to utilise PSA activity and subforms**

**Kallikrein inhibitors**

A deeper understanding of the molecular properties of PSA has engendered new approaches to target this protein for prostate cancer diagnostics and therapy. A seminal example is how the discovery of PSA biochemistry provoked several fruitful endeavors to identify active site directed ligands. Indeed, upon the discovery that PSA is a serine protease, many groups drew from the prior medicinal chemistry literature on this enzyme family to design and synthesise avid and selective peptide-based substrate mimetics. As expected, incrementally more potent binding (and inhibition) was achieved by grafting electrophiles (e.g. aldehydes or boronic acids) onto the peptide leads to engage PSAs catalytic serine nucleophile (68, 69). In addition, small molecules bearing an electrophilic chemotype known to react with activated serine nucleophiles (in this case, a β-lactam) have been shown to irreversibly inhibit PSA activity (26). Several of these compounds suppress angiogenesis in preclinical models, further underscoring the potential value of curbing PSA proteolytic activity. The inhibition of other kallikrein-activities has already been translated to the clinic (Aprotinin; to reduce peri-operative bleeding in cardiac surgery [70]). Pending first in man studies, the ultimate clinical utility of this strategy for PSA can be assessed.

PSA possesses a unique substrate specificity among serine proteases (to hydrolyse amide bonds on peptides bearing a glutamine residue at the P1 position [69]). This feature has inspired some creative chemistry to develop pro-drugs for the selective activation of therapies at the prostate pre-neoplastic or tumour microenvironment. In such a strategy, cytotoxic chemotherapies are rendered inactive by conjugation to a short peptide sequence (HSSKLQ). This sequence is specifically cleaved by PSA in situ with an efficiency vastly exceeding that of other abundant serine proteases (71–73). This then enables site-specific activation and delivery of the chemotherapeutic agent. Among these drug candidates, preclinical trials have shown that systemic administration of a HSSKLQ-doxorubicin adduct incurs no obvious toxicity in mice at doses several fold higher than the MTD for doxorubicin. This pro-drug system demonstrated tumour regression in a prostate cancer model that endogenously produces and secretes catalytically active PSA. Again, the promise of these preclinical findings argues strongly for a human trial to assess the therapeutic potential of this strategy.
Imaging free PSA

The complex posttranslational regulation of PSA protein has also born a clever strategy to image PSA expression at a prostate tumour. Much like the thinking that motivated the pro-drug technology, Ulmert et al. reasoned that tumour imaging could be achieved by selectively targeting the pool of PSA uncomplexed to serpins ("free" PSA) known to transiently exist in pericellular space (74). As PSA in blood is overwhelmingly bound to serpins, this strategy circumvents the inherent challenges in imagining a tumour biomarker with an abundant circulating isoform (in principle, the circulating isoform could impair tumour visualisation by binding a majority of the systemically administered imaging tool). Consistent with this hypothesis, a radiolabelled monoclonal antibody to an epitope on PSA that is masked by bound serpins ($^{89}$Zr-labelled 5A10) produced high contrast images of PSA-secreting prostate cancer tumours in animal models with positron emission tomography (PET). This radiotracer also detected small orthotopic tumours in the bone, suggesting it could be a powerful tool for prostate cancer staging at diagnosis. Of note, some of the aforementioned substrate mimetics have been successfully conjugated to the radionuclide $^{89}$Tc, and the promise of $^{89}$Zr-5A10 PET suggests these radiolabelled ligands may demarcate prostate tumours as well (75).

Ironically, current gaps in our knowledge of PSA biology have also inspired new technologies to expand the utility of this biomarker. Specifically, there is a strong need to understand the paradoxical observation that declines in serum PSA do not lead to a survival benefit from androgen deprivation therapy in castration-resistant prostate cancer (CRPC). It is thought that a more refined monitoring of AR activity, as discussed below in tumours or single cells, rather than an average serum value, could shed light on this phenomenon. The rationale for this hypothesis is based on the widely accepted observation that the concentration of intratumoural PSA vastly exceeds that present in serum, and there is no known role for AR in the rate limiting steps of PSA secretion or leakage into serum. In this regard, one can reasonably speculate that relative changes in secreted PSA levels need not necessarily reflect corresponding changes in AR activity, and by extension, a technology that measures a pool of PSA more closely linked to AR transcriptional activity could provide a higher fidelity readout of AR signalling.

This concept was formally introduced to the literature by comparing $^{89}$Zr-5A10 PET to serum PSA levels post MDV3100 therapy in preclinical models of CRPC. Downregulation of $^{89}$Zr-5A10 localisation to a prostate cancer tumour post MDV3100 treatment was observed without a corresponding change in serum PSA levels from the respective animals over an acute treatment interval. Although far short of validating this concept, these findings nevertheless support the promise in exploring PSA measurements orthogonal to the serum ELISA.

A second example of using PSA measurements to determine AR activity has utilised a novel microfluidics platform to capture and analyse CTCs on a per cell basis. Miyamoto et al. have reported that PSA expression (and that of other markers) can be effectively quantified in CTCs by staining with fluorescently labelled monoclonal antibodies and subsequent immunofluorescent analysis to determine AR status (76) (Figure 2A). This enables single-cell molecular characterisation within and across patients. Significant (and expected) changes in the molecular profile of PSA expression were detected from the CTCs assayed from patients undergoing therapy with Lupron" (an androgen deprivation therapy) or abiraterone acetate (an androgen biosynthesis inhibitor).

In this proof-of-concept report, no pairwise comparison was made between the predictive value of the molecular changes in CTCs versus serum PSA levels of patients. However, the results clearly showed that the CTCs from patient with previously untreated disease show an "AR-on" phenotype, which turned "AR-off" after initial treatment, which also eliminated the CTCs from circulation. Upon progression to CRPC and reemergence of CTCs, the molecular profiling reveals a mixture of phenotypes (Figure 2B). This surprising finding stands in contrast to current popular opinion of the role AR plays in CRPC. It also adds to the discourse regarding inter- or intra-patient tumour heterogeneity at this stage of the disease and may yield molecularly distinguished populations for enhanced patient management.

The relative ease with which blood samples can be acquired from large groups of patients following therapy for such evaluation foreshadows a potentially landmark study for the field. To this debate, serum PSA measurements unfortunately offer little clarity, as it by nature represents an averaged sum of production from lesions in the body. In this regard, it is also important to highlight how developing new technologies for even established serum biomarkers like PSA might enhance our understanding of the fundamental biology of clinical disease.

Conclusion

PSA and hK2 are serine proteases in the kallikrein family that are highly and specifically expressed in the prostate. Their detection in the general blood circulation has become an established biomarker for the detection of prostate cancer. Increasing interest is being directed to their role in regulation of angiogenic and vascular events. While debate continues on the weight that the PSA level should have in directing options for treatment, advancements in the accuracy and information for individualised characterisation of disease continue to be made. New technologies have recently demonstrated the visualisation of PSA pools in vivo with high-resolution, quantitative and non-invasive techniques. Expansion of the preclinical successes of $^{89}$Zr-5A10 to man may ultimately result in the same paradigmatic shift in patient management as occurred with the advent of the modern PSA screening era.

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Conflicts of interest
Dr. Lilja holds patents for free PSA, intact PSA, and hK2 assays. Dr. Lilja is named as co-inventor on a patent application for a statistical method to predict the result of prostate biopsy. None of the other authors declares any conflicts of interest.

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